## Genetic analysis of DMSP metabolism in the marine Roseobacter clade

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Mark Kirkwood

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### **Declaration**

I declare that the work in this thesis has not been previously submitted for a degree at the University of East Anglia or any other university, and all research has been carried out by myself, unless otherwise stated.

Signed

Mark Kirkwood

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## Abstract

Genetic, biochemical, bioinformatic and molecular approaches were used to analyse microbial catabolism of dimethylsulfoniopropionate (DMSP), an abundant anti-stress compound made by marine phytoplankton.

Members of the Roseobacter clade of marine  $\alpha$ -proteobacteria may catabolise DMSP by two different routes; demethylation to form methylmercaptopropionate (MMPA), and cleavage by DMSP-lyases, yielding volatile dimethylsulfide (DMS) plus acrylate.

The DMSP-lyase, DddP, was purified from *Roseovarius nubinhibens* ISM and characterised *in vitro*. Nuclear magnetic resonance spectroscopy and gas chromatography confirmed *bona fide* DMSP lyase activity and mutation of predicted active-site residues abolished DMS production.

DddP was also detected in the fungal coral pathogen *Aspergillus sydowii*, likely acquired from bacteria by inter-Domain horizontal-gene-transfer.

A new DMSP-lyase, DddW, was identified in another Roseobacter species, *Ruegeria pomeroyi* DSS-3, initially by microarray-based demonstrations that transcription of *dddW* was induced in cells grown with DMSP. An adjacent gene encoded the cognate transcriptional regulator. *Escherichia coli* cells that over-expressed DddW cleaved DMSP into DMS plus acrylate. Thus, *Ruegeria pomeroyi* has three DMSP-lyases, with DddP and DddQ being known already; mutational analyses showed that all three contributed to its DMSP-dependent DMS (Ddd<sup>+</sup>) phenotype.

Moran's laboratory had shown that the DMSP demethylase was encoded by *R. pomeroyi dmdA*. I unveiled intimate links between the demethylation and the cleavage pathway(s). A key player is *acuI*, which is co-transcribed with *dmdA*, both genes being induced by DMSP and, more markedly, the DMSP-catabolite, acrylate. Furthermore, AcuI<sup>-</sup> mutants failed to grow on acrylate as sole carbon source and were more sensitive to its toxic effects. AcuI<sup>-</sup> mutants failed to grow on DMSP so, surprisingly, *Ruegeria* likely uses lyase pathway(s) to grow on this compound. A potential regulatory gene, transcribed divergently from *dmdA*, was also identified.

The microarray also, wholly unexpectedly, revealed a suite of *cox* genes involved in carbon monoxide oxidation that was up-regulated in response to DMS.

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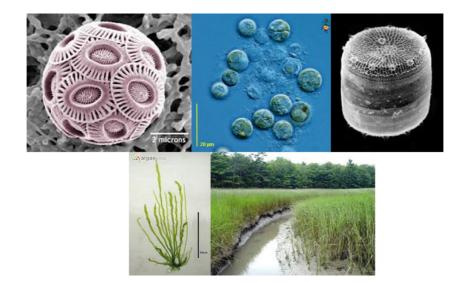
# **Chapter 1**

## **General Introduction**

#### 1.1 Dimethylsulfoniopropionate

Dimethylsulfoniopropionate (DMSP) is a zwitterionic sulfonium compound that is synthesised globally in huge amounts, as much as ~ $10^9$  tons annually (Kettle *et al.*, 1999; Stefels, 2000; Stefels *et al.*, 2007). This biosynthetic ability occurs sporadically throughout a wide range of organisms, including species of the Phyla Haptophyta (e.g. the coccolithophore *Emiliania huxleyi* and *Phaeocystis* sp.), Dinoflagellata (e.g. *Symbiodinium* sp. and *Crypthecodinium cohnii*) and Heterokontophyta (e.g. the diatoms *Melosira numnuloides*, *Thalassiosira pseudonana* and *Fragilariopsis cylindrus*) in the Kingdom Chromoalveolata, as well as widely dispersed members of the Plantae Kingdom, from single-celled Prasinophycae (e.g. *Tetraselmis* sp.) to seaweed macroalgae such as *Ulva intestinalis* (green alga), *Polysiphonia* (red alga) and the angiosperms *Spartina* (salt marsh cord grass) and the beach sunflower *Wollastonia biflora* in the *Compositae* (Stefels, 2000; Otte *et al.*, 2004; Broadbent *et al.*, 2002).

Figure 1.1 Organisms that synthesise DMSP



Clockwise from top left: *Emiliania huxleyi*, *Symbiodinium sp.*, *Thalassiosira pseudonana*, *Spartina alterniflora* and *Ulva intestinalis*.

Soluble DMSP is present at concentrations of less than ~1-2 nM in the open oceans but can reach several  $\mu$ M in phytoplankton blooms (Van Duyl *et al.*, 1998). For example, diatom-dominated sea-ice communities have DMSP concentrations 2-3 orders of magnitude greater

than the average ocean concentration (Kirst *et al.*, 1991; Trevena and Jones, 2006). The intracellular concentrations of DMSP that may accumulate in such organisms are astonishing, ranging from 0.1 to as high as 1 M (Stefels, 2000; Yoch, 2002). The DMSP may comprise 1-16% of the total cellular carbon in some phytoplankton, accounting for the fate of ~11% of the total carbon that is fixed, and between 26% and 44% of the sulfur demand in blooms of these organisms (Matrai and Keller, 1994; Archer *et al.*, 2001). Diatoms alone contribute to ~45% of the total oceanic primary production, and this represents ~25% of global carbon fixation (Nelson *et al.*, 1995). Organisms that do not synthesise DMSP may still exhibit high concentrations of the compound within them. Thus, many cnidarians contain high levels of DMSP, due to symbiotic dinoflagellates, known as zooxanthellae, which associate with these invertebrates in corals (Van Alstyne *et al.*, 2006, 2009; Raina *et al.*, 2009).

#### **1.2 DMSP synthetic pathways**

DMSP biosynthetic pathways are not identical throughout the above taxonomic types, and the sporadic distribution of DMSP production is indicative of several independent evolutionary events. However, all of the known pathways begin with *L*-methionine. There are three general mechanisms for DMSP synthesis, differing between the higher plants, algae and diatoms (Hanson *et al.*, 1994a; James *et al.*, 1995; Kitaguchi *et al.*, 1999).

#### 1.2.1 DMSP synthesis in angiosperms

In the angiosperm, *Wollastonia biflora*, *L*-methionine is first methylated to form *S*-methylmethionine (SMM), followed by successive decarboxylation and deamination to DMSPaldehyde, and finally, an oxidation step to form DMSP (Hanson *et al.*, 1994a). In the saltmarsh cordgrass, *Spartina alterniflora*, the reaction is thought to proceed in a similar way, but with an additional, putative DMSP amine intermediate between SMM and DMSP aldehyde as shown in figure 1.2 (Kocsis *et al.*, 1998; Kocsis and Hanson, 2000).

#### 1.2.2 DMSP synthesis in algae and phytoplankton

Despite the abundance of DMSP in plankton communities, no gene, expressed sequence tag or enzyme responsible for its synthesis had been identified. Recently, however, progress has been made in investigating the molecular basis of DMSP synthesis in the algal diatoms *Fragilariopsis cylindrus* CCMP1102 and *Thalassiosira pseudonana* (Lyon *et al.*, 2011; J D Todd, personal communication).

Diatoms had been shown to have a very different method of making DMSP, via the so-called transamination pathway (Figure 1.2). There is evidence that this pathway also operates in coccolithophores and the green alga *Ulva* (Gage *et al.*, 1997; Summers *et al.*, 1998). The first step is catalysed by a methionine aminotransferase, transferring the amino group to 2-oxoglutarate, yielding glutamate plus 4-methylthio-2-oxobutyrate (MTOB), which is then reduced to *D*-4-methylthio-2-hydroxybutyrate (MTHB) by MTOB reductase. An MTHB *S*-methyltransferase then catalyses the *S*-adenosyl-*L*-methionine (SAM)-dependent *S*-methylation of MTHB, yielding *D*-4-dimethylsulfonio-2-hydroxybutyrate (DMSHB). Finally, DMSHB is decarboxylated to form DMSP by a DMSHB decarboxylase.

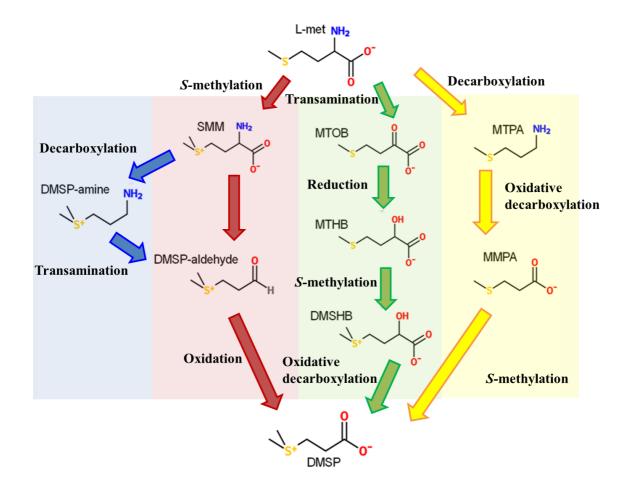
Importantly, the activity of the *S*-methyltransferase only occurs at significant levels within DMSP-producing organisms and is predicted to be specific for DMSP biosynthesis. In the sea-ice diatom *Fragilariopsis cylindrus*, a proteomics approach identified a number of proteins that were increased in their expression in the presence of DMSP, whose intracellular concentration itself increased following raised salinity; under such conditions, one third of the proteins were related to amino acid pathways, and members from all four of the enzyme classes involved in the algal DMSP transaminase biosynthesis pathway were elevated (Lyon *et al.*, 2011). Additionally, a number of specific enzymes have now been ratified, with the genes for both the MTOB reductase and MTHB *S*-methyltransferase from diatoms having been cloned and expressed successfully in the  $\gamma$ -proteobacterium, *Escherichia coli* (J D Todd, personal communication).

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#### **1.2.3 DMSP synthesis in dinoflagellates**

The biosynthesis of DMSP in dinoflagellates differs quite significantly from those described in the angiosperms and algae (Figure 1.2). The dinophyceae, *Crypthecodinium cohnii*, uses *L*methionine as a starting point, as in the other organisms, but this is decarboxylated to form methanethiolpropanamine and then converted to methylmercaptopropionate (MMPA) through oxidative decarboxylation. A final *S*-methylation step yields the DMSP product (Kitaguchi *et al.*, 1999).

# Figure 1.2 The DMSP biosynthesis pathway in angiosperms, phytoplankton and dinoflagellates



Chemical reactions are labelled, with the predicted enzymes illustrated. Panels indicate systems identified in different organisms.

Blue panel: *Spartina alterniflora*; Pink panel: *Wollastonia biflora*; Green panel: coccolithophores (*Emiliania huxleyi*), diatoms (*Thalassiosira pseudonana* and *Fragilariopsis cylindrus*) and green algae (*Ulva*); Yellow panel: dinoflagellates (*Crypthecodinium cohnii*).

*L*-met, methionine; SMM, *S*-methylmethionine; DMSP, dimethylsulfoniopropionate; MTOB, 4methyl-2-oxobutyrate; MTHB, 4-methyl-2-hydroxybutyrate; DMSHB, 4-dimethylsulfonio-2hydroxybutyrate; MTPA, methanethiolpropanamine; MMPA, methylmercaptopropionate; DMSP, dimethylsulfoniopropionate.

Adapted from Trossat et al., 1998; Hanson et al., 1994a; Gage et al., 1997; Kocsis et al., 1998.

#### 1.2.4 Cellular localisation of DMSP synthetic enzymes

DMSP synthesis in plants may occur in the chloroplast, with DMSP and the enzyme reportedly being concentrated in this organelle in *W. biflora* (Trossat *et al.*, 1998). Tentatively, the localisation of the relevant enzymes in the diatoms matches this, since the methionine aminotransferase, MTHB *S*-methyltransferase and DMSHB decarboxylase polypeptides of *F. cylindrus* and *T. pseudonana* contain N-terminal leader sequences that are predicted to target them to the chloroplast (Gruber *et al.*, 2007; Lyon *et al.*, 2011; J D Todd, personal communication).

#### 1.3 Physiological uses of DMSP

DMSP has been implicated in a number of important physiological processes, and is primarily thought to act as compatible solute that protects the cells that accumulate it from osmotic stress in the saline marine environment (Otte and Morris, 1994; Edwards, *et al.*, 1988). However, direct evidence for this is scarce and not wholly consistent. For example, phytoplankton species such as *Ulothrix* spp. (Chlorophyta) and *Enteromorpha bulbosa* (Now *Ulva*, Chlorophyta) exhibit a clear, positive DMSP-accumulation response to salinity (Karsten *et al.*, 1992), as do diatoms, but *Spartina* spp. display no such response (Otte *et al.*, 2004). Studies using a heterologous host, *E. coli*, have shown that DMSP bestows halotolerance under artificial, laboratory conditions (Cosquer *et al.*, 1999). Interestingly,  $\beta$ alanine betaine, a structural analogue of DMSP, is used by members of the Plumbaginaceae as a salt stress response molecule (Hanson *et al.*, 1994b). Likewise, DMSP resembles glycine betaine, a widely utilised osmo-protectant in terrestrial organisms. It has been suggested that DMSP is favoured in marine environments due to the abundance of sulfur and scarcity of nitrogen in the seas; on land, the reverse is true, favouring the use of the amino acid osmoticum (Kiene *et al.* 2000; Galinski *et al.*, 1995).

Other biological functions for DMSP have also been proposed. One such use is as an antistress compound, for protection from UV light and/or amelioration of oxidative damage (Otte *et al.*, 2004; Sunda *et al.*, 2002). Known oxidative stressors such as carbon and iron limitation in *E. huxleyi* and *Thalassiosira pseudonana* also increase production of DMSP (Sunda *et al.*,

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2002). It has also been suggested that DMSP acts as an herbivore deterrent via the release of one its catabolic products, acrylate (Van Alstyne *et al.*, 2003a). This might explain why species that readily synthesise DMSP are less likely to be targeted by zooplankton predators than those that accumulate lower concentrations (Wolfe, 1997). DMSP may also act as a biological sulfur sink in the form of reduced sulfur (Kiene *et al.*, 1999; see below), as an overflow mechanism during nitrogen limitation (Stefels, 2000), or as a cryoprotectant (Karsten *et al.*, 1996).

Of course, these proposed functions are not mutually exclusive. These may be resolved (at last!) if and when a mutant that is defective in DMSP production becomes available for at least one organism.

#### **1.4 DMSP utilisation**

The release of DMSP from the producing plankton cells may occur through viral lysis, senescence or predation (Malmstrom *et al.*, 2004a). Several marine bacteria can import DMSP from the environment, including members of the cyanobacteria (*Synechococcus* and *Prochlorococcus*) and many heterotrophic species (Malmstrom *et al.*, 2004b; Vila-Costa *et al.*, 2006, Sun *et al.*, 2011). These include the SAR11 clade of the  $\alpha$ -proteobacteria, which are prodigiously abundant in the oceans (see later) and which favour the use of reduced forms of sulfur, even ahead of such abundant S sources as sulfate which are bio-energetically more expensive to assimilate (Tripp *et al.*, 2008).

In addition, a few of the algae and phytoplankton that synthesise DMSP may catabolise it. Indeed, one of the first DMSP catabolic reactions to be described was in the red seaweed *Polysiphonia lanosa*, and involves the cleavage of DMSP into acrylate plus the volatile, dimethyl sulfide (DMS) (Cantoni and Anderson, 1956; Steinke *et al.*, 1998). However, the cultures were never reliably confirmed to be axenic, and although DMSP biotransformation has since been reported in some species of single-celled phytoplankton, such as the haptophyte *E. huxleyi*, this criticism also applies to many of these studies on Eukaryotic algae (Franklin *et al.*, 2010; S Newton-Payne, personal communication). What is indisputable,

though, is that many marine bacteria, and a few fungi, can catabolise DMSP, with acrylate and DMS being the products in some examples of this phenomenon.

In total, it has been estimated that of the *ca*. one billion tons of DMSP catabolised annually, around one third is subjected to this form of cleavage; the remainder being degraded by demethylation, which does not liberate any DMS – see below. Much of the resultant DMS appears to be retained in these microbial communities, but roughly 10% escapes into the atmosphere. Here, it may be spontaneously oxidised into sulfates, sulfur dioxide, methanesulfonic acid, and other products that act as cloud condensation nuclei (Hatakeyama *et al.*, 1982). This involves major movement of S from sea to air thence back to land, via precipitation, making DMSP a significant player in the global sulfur cycle (Andreae, 1990). The increase in albedo created by DMS oxidation products has been predicted to affect the climate via a positive feedback loop, termed the CLAW hypothesis, shown in Figure 1.3 (Charlson *et al.*, 1987). However, recent counterclaims have suggested DMS has little to no effect as cloud condensation nuclei, other than at a local level in regions of high primary productivity, such as coral reefs (Quinn and Bates, 2011). These are the richest source of DMSP, and hence of DMS, because of the dinoflagellate *Symbiodinium* and the many bacteria in these ecosystems (Jones and Trevena, 2005; Raina *et al.*, 2009; 2010).

Atmospheric or dissolved DMS can also act as a potent chemo-attractant for marine animals, including shearwaters, seals, penguins and crustaceans (Steinke *et al.*, 2002; 2006; Nevitt, 2008; 2011; Cunningham *et al.*, 2008).This was even found to act through more than one trophic level. For example, bacterial production of DMS from the DMSP released by phytoplankton encourages grazing by crustaceans and acts as a spatial foraging cue for procellariiform birds, such as petrels (Nevitt and Bonadonna, 2005).

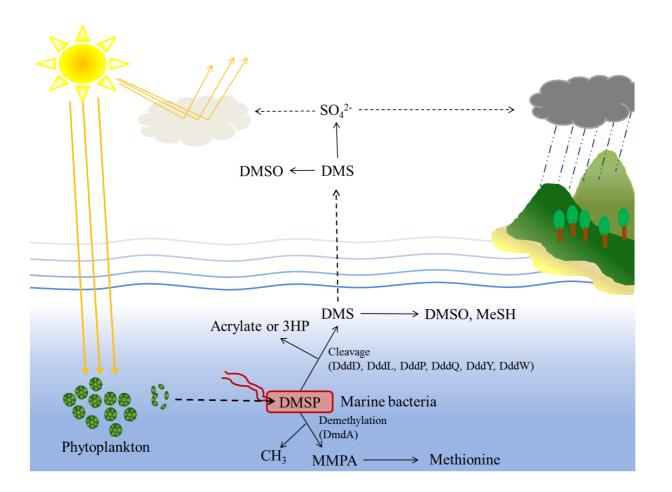


Figure 1.3 Formation and catabolic fate of DMSP

Once released by phytoplanktonic producers, through grazing, senescence or viral lysis, dimethylsulfoniopropionate (DMSP) is subjected to bacterial degradation by two general pathways. The demethylation pathway leads to the production of methylmercaptopropionate (MMPA), whereas the cleavage pathway yields dimethyl sulfide (DMS) and either acrylate or 3-hydroxypropionate (3HP). The DMS can be further transformed by bacteria into dimethyl sulfoxide (DMSO) and other products. Some of the DMS escapes to the atmosphere, where it is oxidised, forming cloud condensation nuclei, perhaps leading to increased albedo. The transfer of these compounds to the Earth's surface, via rain or snow, is an important step in the global sulfur (S) cycle, creating a flow of S from sea, to air thence to land. MeSH, methanethiol. Adapted from Curson *et al.*, 2011a.

#### 1.5 α-proteobacteria, Rhodobacterales, Roseobacters and *Ruegeria* pomeroyi DSS-3

There is one group of organisms that is consistently the dominant player in DMSP utilisation. These are members of the sub-phylum of  $\alpha$ -proteobacteria, and particularly one or two taxa within this group. The  $\alpha$ -proteobacteria form one of the largest and most diverse groups within the Eubacteria, with diverse metabolic profiles, and with examples of phototrophs, chemolithotrophs, chemoorganotrophs and aerobic photoheterotrophs (Venter *et al.*, 2004; Kersters *et al.*, 2006). Among this sub-phylum, two groups, the SAR11 clade and the Roseobacters are especially abundant in the oceans and play a pivotal role in global turnover of DMSP.

#### **1.5.1** General introduction to the Roseobacters

DMSP is a major component of the lifestyle for a group collectively as the Roseobacters, as exemplified initially by physiological, and then by genetic studies. The first description of members of the Roseobacter clade occurred with the culturing from Japanese coastal waters of a high density population of pink, aerobic, anoxygenic phototrophs (AAoP) that contained bacteriochlorophyll- $\alpha$  (Shiba et al., 1979). The genus Roseobacter and the species Roseobacter denitrificans (previously Erythrobacter sp.) and R. litoralis, were created for AAoP that contained bacteriochlorophyll- $\alpha$  and which were isolated from marine algae (Shiba et al., 1991). Later, examinations of estuarine and coastal seawater samples using 16S rRNA PCR showed that these species, along with other Roseobacters such as Sulfitobacter EE-36 and members of the  $\alpha$ -proteobacteria, were a numerically dominant group in marine bacterioplankton communities (González and Moran, 1997). Although the term Roseobacter is not an officially recognised taxonomic term, it is widely used as informal shorthand for a group of some 17 taxonomically coherent genera, within the Rhodobacteraceae family (Buchan et al., 2005; Tang et al., 2010). Apart from their taxonomic relatedness, members of the Roseobacters share one other pivotal feature – they are all marine-living, unlike other genera in the Rhodobacteraceae, such as Paracoccus. Indeed, the Roseobacters are now thought of as one of the ten main taxa found in coastal and open waters, comprising up to 30% of the total number of cells in these environments. They can represent ~3% of all cells in

open ocean waters and also occur in hypersaline lagoons, coastal waters, marine sediments, and sea ice (Labrenz *et al.*, 1998; Eilers *et al.*, 2000; Brinkmeyer *et al.*, 2003; Buchan *et al.*, 2005; Biers *et al.*, 2009). During algal bloom events in coastal waters large amounts of DMSP are released into the milieu and Roseobacters are the dominant taxon in these environments (González *et al.*, 2000).

Over 40 members of the Roseobacter lineage have had their genomes sequenced, yet only recently has a robust phylogenetic analysis occurred (Newton *et al.*, 2010; Tang *et al.*, 2010). Different Roseobacter genera display at least 89% identity in their 16S rRNA sequences, and share a similar G + C content (*ca.* 50-60%). However, there is considerable variation in their genomes, with genome sizes that range from 3.5 Mbp (Mega base pairs) to 5.3 Mbp - corresponding to 3500 to 5500 genes (Buchan *et al.*, 2005; Tang *et al.*, 2010). In addition to a conserved set of core genes retained through vertical evolution, there exists a significant subset of accessory genes, which have undergone extensive lateral gene transfer during the evolution of the Roseobacters (Tang *et al.*, 2010; see chapters 4, 5, 6). Included in this portfolio of accessory genes are those involved in DMSP catabolism (see below).

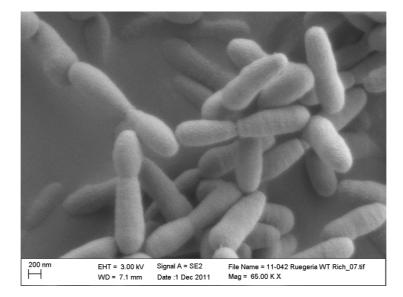
Some characteristic physiologies of the Roseobacters include:

- the ability to carry out organic and inorganic sulfur metabolism, contributing significantly to the biogeochemical cycling of this element (Moran *et al.*, 2003; see below);
- the oxidation of carbon monoxide, a potentially significant response to the photolysis of dissolved organic matter (Zuo and Jones, 1995; King, 2003);
- symbiotic relationships with eukaryotic marine organisms such as the dinoflagellate, *Pfiesteria* (Alavi *et al.*, 2001), and including pathogenic effects on scleractinian corals (Cooney *et al.*, 2002).

#### 1.5.2 Ruegeria pomeroyi DSS-3

The model Roseobacter, *Ruegeria pomeroyi* DSS-3 (originally named *Silicibacter pomeroyi* but subsequently re-classified [Yi *et al.*, 2007]), was isolated from the Coastal Atlantic, off the shores of Georgia. It forms short, rod-shaped cells with a complex, polar flagellum that confers motility (Figure 1.4). It was clear, soon after its isolation, that this strain can catabolise DMSP by both of the general pathways that had been partially described earlier, but which had been seen in different organisms. *Ruegeria pomeroyi* is rather a glutton for DMSP, as it can catabolise this compound by two different mechanisms, known as demethylation and cleavage (González *et al.*, 1999; 2003; see below).

#### Figure 1.4 Microscopic analysis of Ruegeria pomeroyi DSS-3



## Scanning electron micrograph of *R. pomeroyi* DSS-3 cells grown in rich ½ YTSS medium. Scale bar given. (R T Green, personal communication).

*R. pomeroyi* DSS-3 is easily cultured compared to many of its relatives (González and Moran, 1997), and was the first Roseobacter to be genome-sequenced (Moran *et al.*, 2004). It has 4,283 genes in a total genome that is 4.1 Mbp in size, and includes a 491 kilo base pair (kbp) megaplasmid (www.roseobase.org). The original sequencing study identified a number of genes related to its metabolic profile, as discussed in following chapters.

#### 1.5.3 The SAR11 Clade

The second of the two groups of  $\alpha$ -proteobacteria that are known to metabolise DMSP is the SAR11 clade, or "Pelagibacteraceae", belonging to the Order Rickettsiales. These were first identified by culture-independent approaches, using environmental 16S rRNA sequences obtained from the <u>Sargasso Sea (Giovannoni *et al.*, 1990)</u>. A member of the SAR11 clade was cultured, with some difficulty, in free-living form (the doubling time of a cell is *ca.* 29 hours). This species, *Candidatus* Pelagibacter ubique, has a rod shape with one of the smallest lengths of any self-replicating cell, at 0.4-0.9 µm and a diameter of only 0.1-0.2 µm. It also has the smallest genome of any free living organism at 1,308,759 bp, and yet these bacteria are massive in terms of their global population and distribution. There are *ca.* 10<sup>28</sup> cells of Pelagibacter in the seas, comprising around 50% of cells in the temperate oceans during summer (Morris *et al.* 2002; Giovannoni *et al.*, 2005; Thrash *et al.*, 2011). It is a scavenger, whose ability to recycle dissolved organic carbon means that it plays a major role in the Earth's carbon cycle. Pelagibacter spp. are only known to harbour the genes for DMSP demethylation at present (see below).

Both of these groups of organisms are important to global DMSP utilisation. However, whilst the SAR11 clade only demethylates DMSP, many Roseobacters harbour the enzymes necessary for both forms of DMSP degradation. The pathway that occurs in both Roseobacters and the SAR11 clade, DMSP demethylation, is a good starting point to examine these systems in more detail.

#### **1.6 DMSP demethylation**

The predominant pathway of DMSP catabolism, which accounts for approximately 70% of the globally consumed DMSP, occurs through the "demethylase pathway" (Figure 1.5; Kiene, 1996; Kiene *et al.*, 1999; 2000). The discovery of DMSP demethylation began with the observation by Mopper and Taylor (1986) that methane thiol (MeSH) and methyl mercaptopropionate (MMPA) were the dominant thiols present in anoxic marine intertidal

Tracer experiments on the catabolic fate of <sup>35</sup>S labelled DMSP, showed that 15-40% of DMSP in natural populations of bacterioplankton from subtropical and temperate marine waters was incorporated into macromolecules, via conversion of MeSH to methionine, and it was inferred that this was via the demethylation route (Kiene and Service, 1991).

Although there had been several suggestions for the pathway(s) of DMSP demethylation, it is only recently that a combination of biochemistry and genetics has generated what appears to be the definitive pathway, at least in one strain, namely the model Roseobacter, *Ruegeria pomeroyi* DSS-3.

#### 1.6.1 DmdA

The first gene found to have a role in the demethylation pathway was *dmdA*, encoding a DMSP demethylase enzyme, DmdA (Howard *et al.*, 2006). A *Ruegeria pomeroyi* DSS-3 Tn5 transposon mutagenesis library was screened for mutants that were unable to produce MeSH, but could still release DMS. The DNA from one such mutant was sequenced out from the transposon site and the gene identified as *SPO1913*. Complementation assays confirmed that the gene product was responsible for the DMSP to MeSH phenotype, and cell extract assays narrowed this down to the first step in the reaction, that of DMSP demethylation to MMPA.

Bioinformatic analysis of DmdA placed it within a family of glycine cleavage proteins (EC 2.1.2.10), exemplified by the GcvT of *E. coli* (Stauffer *et al.*, 1993; Moran *et al.*, 2004). DmdA also shares domains with dimethylglycine oxidase and sarcosine oxidase, but exhibits greatest homology with the glycine cleavage system (Schuller *et al.*, 2012). There is a ~25% identity between homologues of DmdA and other members of the GcvT family (e.g. *E. coli*) and phylogenetically, the DMSP demethylases form a distinct out-group (Reisch *et al.*, 2008).

A common characteristic among these enzymes is the requirement for a tetrahydrofolate (THF) co-factor. In the case of DmdA, this involves the transfer of the *S*-methyl group from DMSP to THF, to form 5-methyl THF and MMPA, as noted in *Candidatus* Pelagibacter

ubique and *Ruegeria pomeroyi* (Reisch *et al.*, 2008). Interestingly, this co-factor differs from other enzymes in the GcvT family, whose reactions involve the production of 5,10-methylene-THF (Schübert *et al.*, 2003). The use of THF as a cofactor is discussed in more detail in chapter 6.

#### **1.6.2 DMSP demethiolation**

It had been thought that two mechanisms may exist for the subsequent catabolism of the MMPA produced from DMSP demethylation. The first is via a second demethylation, which converts MMPA to 3-mercaptopropionate (MPA), as noted in anoxic marine sediment (Kiene and Taylor, 1988; Visscher *et al.*, 1992; Visscher and Taylor, 1994). However, there is limited evidence for the physiological significance of this pathway in aerobic marine surface waters.

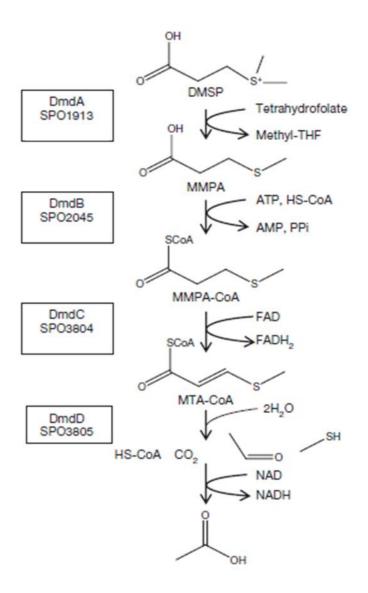
The second proposed, but non-ratified, pathway is known as demethiolation and, crucially, allows the siphoning of DMSP into central metabolism, and as a precursor to amino acids via conversion of MeSH (Kiene *et al.* 1999, 2000). The pathway involves a fatty acid  $\beta$ -oxidation-like reaction, in order to produce MeSH, acetaldehyde, CO<sub>2</sub> and CoA (Figure 1.5; Reisch *et al.*, 2011a; 2011b; see chapters 5 and 6). These products can then be easily shuttled into systems such as amino acid synthesis (methionine) and central metabolism (the citric acid cycle). These successive steps are carried out by the DmdB, DmdC and DmdD enzymes, recently characterised in *Ruegeria pomeroyi* DSS-3 (Reisch *et al.*, 2008; 2011a; 2011b).

#### 1.6.3 DmdB/DmdC/DmdD

DmdB (the product of *R. pomeroyi* gene, *SPO2045*) is an ATP dependent enzyme, originally labelled as a fatty acid CoA ligase, which forms an MMPA-CoA thioester from MMPA. The FAD-dependent enzyme DmdC (*SPO3804* in *R. pomeroyi*) converts this to methylthioacryloyl-CoA (MTA-CoA). Finally, the crotonase-like DmdD (encoded by

The transcript levels responsible for the three enzymes were seen to increase in response to DMSP or MMPA, and mutations in *dmdA*, *dmdB*, *dmdC* or *dmdD* of *R. pomeroyi* abolished DMSP-dependent production of MeSH (Howard *et al.*, 2006; Reisch *et al.*, 2011a; 2011b). Interestingly, the *dmdC* mutant was able to grow well on DMSP as a sole C source, and a *dmdD* mutant was inhibited for growth on the same substrate. Thus, it is likely that products of DMSP cleavage are sufficient for growth of *R. pomeroyi* in the absence of MMPA, and it seems that the MTA-CoA intermediate may be toxic, given its accumulation in the *dmdD* mutant. Excitingly, this also implies that MMPA may be used for growth in a pathway independent of DMSP demethylation, and this will be explored further in chapters 5 and 7.

## Figure 1.5 DMSP demethiolation pathway in *Ruegeria pomeroyi* DSS-3



Enzymes given with relevant "SPO" gene number. DMSP is first demethylated to MMPA by DmdA, using tetrahydrofolate as a cofactor. An MMPA-CoA thioester is then formed, with the concomitant consumption of ATP and production of AMP. The MMPA-CoA is then dehydrogenated to an enoyl-CoA intermediate, MTA-CoA. This is then hydrated, releasing MeSH, CoA, CO<sub>2</sub> and acetaldehyde, which is then oxidized to acetic acid by acetaldehyde dehydrogenase. From Reisch *et al.*, 2011b.

#### 1.6.4 Distribution of the *dmd* genes

DmdA is almost exclusively found in the SAR11 bacteria and in the Roseobacters, with a high level of conservation at the amino acid level among most of these homologues, with weaker matches in the  $\gamma$ -proteobacterium, HTCC2080 and a newly discovered member of the SAR116 clade (Howard *et al.*, 2006; 2008; Reisch *et al.*, 2008, Varaljay *et al.*, 2010).

The distribution of the secondary *dmd* genes is curious, with *dmdB* and *dmdC* being present in all 36 *dmdA*-possessing organisms in the genomic database, but also in several species that lack DmdA, including many  $\beta$ - and  $\gamma$ -proteobacteria; representatives of these bacteria produced MeSH from MMPA (Reisch *et al.*, 2011b), implying that there are alternative sources of MMPA that do not originate from DMSP catabolism.

The DmdD enzyme appears to be significantly less abundant than the other enzymes. A BLASTp interrogation of the Global Ocean Sampling (GOS) database, which is a metagenomic database of environmental DNA samples taken from the world's oceans (Rusch *et al.*, 2007), only retrieved 16 weak homologues. DmdD may be replaced by non-orthologous iso-functional enzymes in other organisms, given the activity of this pathway in *R. lacuscaerulensis*, which possesses DmdB and DmdC, but not DmdD (Reisch *et al.*, 2011b).

#### **1.7 The DMSP lyases**

Although the DMSP demethylation pathway may account for the majority of the degradation of DMSP in the natural marine environments, and the *dmdA* gene is very widespread in marine metagenomic data bases, this pathway does not release the environmentally influential gas, DMS. For many years, it was thought that this volatile was formed by a cleavage of the DMSP by an enzyme known generically as DMSP lyase, which would form acrylate as the other, C3 catabolite.

However, it is only thanks to recent genetic work that has extended earlier physiological and biochemical work that we have a clear idea of the mechanisms and the pathways that are

involved. As seen below, these newer studies have ratified previous thinking in some cases, but in others, there have been some unexpected outcomes. To a large extent, this is because there has been a wholly unexpected degree of diversity, with different microbes (some of which were wholly unanticipated) using a range of different enzymes and generating a collection of different intermediate catabolites.

The overall phenotype that describes the breakdown of DMSP with the concomitant liberation of DMS has been termed  $Ddd^+$  (<u>DMSP-dependent DMS</u>). Thanks to work in the UEA lab, no fewer than six different "primary" Ddd enzymes have been shown to mediate the cleavage of DMSP, and many other polypeptides and the corresponding genes have been shown to be involved in such ancillary functions as the transport of the DMSP substrate into the cells, the regulation of the expression of the *ddd* genes, in the catalysis of subsequent, downstream catabolic steps and even in the protection of the cell against damage inflicted by one of the intermediate catabolites.

The isolation and characterisation of these six different *ddd* genes, namely *dddD*, *dddL*, *dddP*, *dddQ*, *dddW* and *dddY* followed a standard approach, as follows. First, either novel strains of DMSP-catabolising bacteria were isolated, or known strains with this ability were obtained from other labs or culture collections. Next, genomic libraries of such strains were made in a wide host-range cosmid vector. These cosmids in this library were then introduced into an appropriate bacterial host strain that does not catabolise DMSP and screened for transconjugants that either cleaved DMSP to release DMS (Ddd<sup>+</sup> phenotype) and/or which grew on DMSP as sole carbon source. These genes would then be localised by sub-cloning and/or mutagenesis. The availability of genomic sequences of bacteria that are known to catabolise DMSP was exploited to search for the presence or absence of the various *ddd* genes. Once identified, the selected microbes were examined for those that contain homologues of the *ddd* genes already identified as above and confirmed that these can cleave DMSP even if they had hitherto not been suspected of having this ability. This cycle was then repeated for those cases where it was known that a given strain had a Ddd<sup>+</sup> phenotype, but its genome lacked any of the previously identified *ddd* genes.

The features of these six primary genes and their products are now presented, with the exception of DddW, which is a result of the work described in chapter 4.

#### 1.7.1 DddD

The first *ddd* gene was discovered in the  $\gamma$ -proteobacterium, *Marinomonas* sp. MWYL1, isolated from the rhizosphere of the salt marsh grass *Spartina anglica*, on the basis of its ability to grow well on DMSP as sole carbon (C) source (Todd *et al.*, 2007). A fosmid library of this strain was introduced into *E. coli* and screened for a Ddd<sup>+</sup> phenotype using gas chromatography. This identified two transcriptional units that were responsible for DMS release in *Marinomonas* (see below; Figure 1.6). It was shown that both *dddD* and the divergently transcribed *dddTBCR* were required for a Ddd<sup>+</sup> phenotype in *Marinomonas* itself, but that, *dddD* alone conferred a Ddd<sup>+</sup> phenotype to *E. coli* if the cloned gene was expressed from an active ectopic promoter (Todd *et al.*, 2007).

The DddD polypeptide belongs to the Class III CoA-transferases and shares homology with CaiB of *E. coli*. CaiB is involved in an anaerobic respiratory pathway, transferring CoA to the amino acid carnitine, which is used as an electron acceptor (Eichler *et al.*, 1994; Elssner *et al.*, 2001). CaiB is a homodimer whose individual polypeptides are approximately half the size of the DddD polypeptide. Interestingly, DddD comprises two repeated CaiB-like domains with a short linker region (Rangarajan *et al.*, 2005; Todd *et al.*, 2007) suggesting that DddD acts as a form of "intra-molecular dimer". This would allow it to transfer CoA to DMSP from an acyl-CoA donor before the release of DMS, and would result in an acyl-CoA intermediate prior to the DMS releasing step (Todd *et al.*, 2007). Unfortunately, no *in vitro* study of DddD has confirmed this mechanism.

#### 1.7.1.1 Distribution of DddD

Following the discovery of DddD in *Marinomonas* sp. strain MWYL1, further Ddd<sup>+</sup> strains were identified by growth on DMSP as a sole C source. Thus, Curson *et al.* (2010) isolated two species of  $\gamma$ -proteobacteria from the gut of the Atlantic herring, *Clupea harengus* which possessed the *dddD* gene. These species, *Pseudomonas* strain J465 and *Psychrobacter* strain J466, were found to possess homologues of closely linked *ddd* genes that resembled those in *Marinomonas* sp. strain MWYL1 (Figure 1.6 and see below).

Using bioinformatics as a form of gene mining, close homologues of DddD were found in silico using BLAST analysis. In addition to matches to other species of Marinomonas (>90% identity at peptide level) and another  $\gamma$ -proteobacterium, Oceanimonas doudoroffii, there were also homologues of DddD in certain members of the  $\alpha$ -proteobacteria, namely the Roseobacters, Sagittula stellata E37, Hoeflea phototrophica DFL-43, Dinoroseobacter shibae DFL 12, Rhodobacterales sp. KLH11 and Ruegeria pomeroyi DSS-3 (Johnston et al., 2008; Todd et al., 2010; Curson et al., 2011a). Interestingly, DddD also appeared in other "terrestrial" strains. These included the  $\alpha$ -proteobacterium Rhizobium NGR234, which is a particularly unusual strain, in that it nodulates a wide range of hosts, including Parasponium, which is a non-legume plant (Trinick, 1973). Another homologue was found in the  $\beta$ proteobacterium, Burkholderia cepacia AMMD, which is present in the rhizosphere of many angiosperms (Ramette et al., 2005). The two latter examples were shown not to be able to grow on DMSP as a sole C source, but were confirmed to exhibit Ddd<sup>+</sup> phenotypes when provided with DMSP and an alternative C source. The DddD-like polypeptides from D. shibae, Rhodobacterales sp. KLH11 and R. pomeroyi DSS-3 form a distinct out-group from the other Roseobacter DddD enzymes, being >72% identical to each other, yet only sharing 40% identity with the original Marinomonas peptide. Additionally, mutations into the gene responsible for this "dddD" gene in R. pomeroyi (SPO1703) had no discernible effect on DMSP dependent DMS production and quantitative real time PCR (qRT-PCR) analysis of dddD illustrated low (if any) expression of its messenger RNA (Todd et al., 2011).

The presence of DddD in these organisms almost certainly arose via Horizontal Gene Transfer (HGT) between the lineages (Todd *et al.*, 2007). Indeed, an inter-Domain HGT event may even have occurred between bacteria and the coccolithophore, *Emiliania huxleyi*, which has a deduced polypeptide with ~30% identity to *Marinomonas* DddD (Curson *et al.*, 2011a).

#### 1.7.1.2 DddD in Halomonas HTNK1

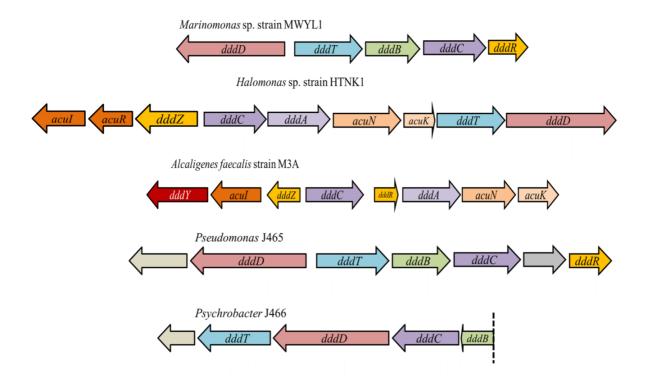
Although many of the DddD-containing bacteria, (notably the  $\gamma$ -proteobacteria) grew well on DMSP, they did not grow on acrylate as sole C source. Initially, this was surprising because the generally accepted view was that DMSP lyase generated acrylate as the initial, C3 catabolite. It was therefore surprising that none of the DddD-containing organisms (above) could grow on acrylate as a sole C source. Therefore, a search was made to find a strain that could grow on both acrylate and DMSP. This led to the isolation from the surface of the DMSP-producing macroalga *Ulva lactuca* (sea lettuce), of the  $\gamma$ -proteobacterium, *Halomonas* strain HTNK1 (Todd *et al.*, 2010) which, like *Marinomonas*, is in the *Oceanospiralleles*.

The gene cluster responsible for the growth phenotype in *Halomonas* was then discovered using similar methods to *Marinomonas*, as above. It contains *dddD*, *dddT* and *dddC* homologues (see below), plus several other genes that had not been seen in the *ddd* gene clusters described above (Figure 1.6). Therefore, *Halomonas* HTNK1 was chosen to undertake a biochemical study of the fate of DMSP (Todd *et al.*, 2010).

# **1.7.1.3** The Ddd and Acu enzymes – a mechanism for DMSP or acrylate dependent 3HP production

As noted, DddD mediates the initial transfer of CoA onto DMSP, and when *Halomonas* DddD was expressed in *E. coli*, it was found that cell-free extracts of the engineered strain converted DMSP to 3-hydroxypropionate (3HP) plus DMS. Consistent with this, *Halomonas* itself could grow on 3HP, the subsequent catabolic steps being mediated by the DddA and DddC enzymes which, respectively, catalyse the conversion of 3HP to malonate semialdehyde (MalSA) and the latter to acetyl-CoA (Todd *et al.*, 2007; 2010; Curson *et al.* 2008; Figure 1.7). Note that *Marinomonas* MWYL1 does not have a *dddA* gene but that the *ddd* cluster includes *dddB*, which is conversely absent from that of *Halomonas*. Both genes encode an alcohol dehydrogenase enzyme, but belong to different families; DddA and DddB are thought to bind flavin and Fe, respectively. However, it is thought that they achieve similar physiological goals.

# Figure 1.6 The *dddD/TBCR*, *dddZ*, *acuR*, *acuI* and *dddY* genes in strains of *Marinomonas*, *Halomonas*, *Alcaligenes faecalis*, *Pseudomonas* and *Psychrobacter*



Gene names given inside arrows. Not to scale. Dashed line indicates break point in cosmid library.

dddD: DMSP lyase; dddY: DMSP lyase;

*dddT*: Betaine carnitine choline transport (BCCT) family, DMSP transporter;

*dddB*, *dddA*: Flavin containing alcohol dehydrogenase; *dddC*: methyl-malonate semialdehyde dehydrogenase;

*dddR*: LysR-like regulator; *dddZ*: DddR-like regulator;

*acuI*: acryloyl-CoA reductase (see chapter 5); *acuR*: TetR-like regulator;

*acuN*: CaiB-like gene product, crotonobetainyl-CoA / carnitine CoA transferase; *acuK*: CaiD-like gene product, crotonobetainyl-CoA hydratase;

*ddd*: <u>D</u>MSP <u>dependent D</u>MS production; *acu*: <u>Ac</u>rylate <u>u</u>tilisation.

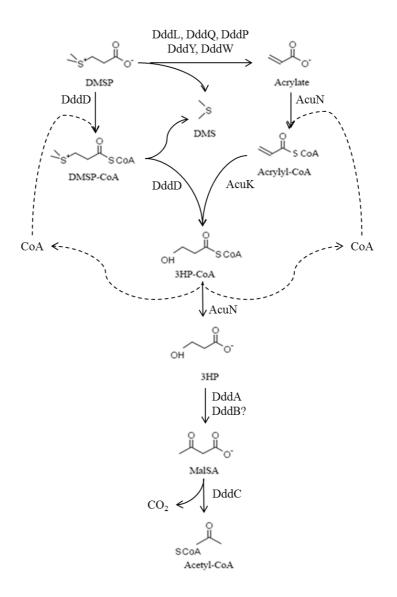
Adapted from Todd et al., 2007; Curson et al., 2011b; 2011c.

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Although this explained the pathway for DMSP catabolism, it did not account for the growth of *Halomonas* on acrylate. For that ability, two other genes in the *ddd* cluster are needed. These *acuN* and *acuK* (Acu = <u>ac</u>rylate <u>u</u>tilisation) genes respectively encode gene products that resemble the CaiB and CaiD enzymes of *E. coli*. CaiB forms a homodimeric crotonobetainyl-CoA / carnitine-CoA transferase, while CaiD is a crotonobetainyl-CoA hydratase (Elssner *et al.*, 2001). It is hypothesised that 3HP is produced from DMSP or acrylate via one or more CoA intermediates, through the actions of DddD or AcuN/K, respectively. Thus, DddD produces DMSP-CoA, and AcuNK produce acryloyl-CoA, 3HP-CoA and 3HP (Figure 1.7). Importantly, mutations in AcuN or AcuK abolish growth on acrylate, as found in *Alcaligenes faecalis* M3A (see below; Curson *et al.*, 2011c).

To recap, DddD shares homology with the CaiB enzyme and this is reflected in the similar activities of the two systems. DddD may in fact act as several enzymes at once, carrying out the initial addition of CoA to DMSP, followed by a hydration, with concomitant DMS release. An exquisite suggestion is that the 3HP-CoA formed by the actions of AcuNK enzymes may act as the acyl-CoA donor in the first step of the reaction, creating a self-perpetuating system (Figure 1.7; Todd *et al.*, 2010). Alternatively, an as yet unknown CoA donor may be utilised.

### Figure 1.7 Predicted mechanism of DMSP and acrylatedependent 3HP production



The suggested convergence of the DddD, DMSP lyase and the AcuN/K pathways at a shared intermediate, 3HP-CoA, before conversion to 3HP is shown. The 3HP-CoA may act as the acyl CoA donor for the initial addition of CoA to DMSP or acrylate by DddD or AcuN, respectively, as shown by dashed arrows. From 3HP, there is a conversion to malonate semi-aldehyde (MalSA), catalysed by very different types of alcohol dehydrogenases – DddA in the case of (e.g.) *Halomonas* and DddB in (e.g.) *Marinomonas*. The MalSA is then converted to acetyl CoA and CO<sub>2</sub> via DddC. In organisms that do not contain AcuNK enzymes, such as *Ruegeria pomeroyi*, a different catabolic pathway for acrylate is predicted (see chapters 5 and 7). Adapted from Elssner *et al.*, 2001; Todd *et al.*, 2007; 2010.

Thus, the ability to grow on DMSP and on acrylate is not due to the conversion of the former to the latter, but to the independent catabolism of each, converging at 3HP.

Interestingly, some *Archaea*, such as *Metallosphaera sedula* contain a pathway known as the 3HP/4-hydroxybutyrate pathway. In this pathway, CO<sub>2</sub> and acetyl CoA are combined to produce malonyl-CoA, with this being reduced to MalSA by a NADPH-dependent malonate semialdehyde dehydrogenase. It is then converted through via 3HP, 3HP CoA and acryloyl-CoA (Berg *et al.*, 2007; Stines-Chaumeil *et al.*, 2006; Hügler *et al.*, 2003). This is the reverse reaction that is predicted to occur in *Halomonas* HTNK1 (Figure 1.7; Todd *et al.*, 2010).

#### 1.7.1.4 DddD and DMSP Transporters

It was noted that the *dddD*-containing gene clusters of several  $\gamma$ -proteobacteria included one or more genes that were strongly predicted to encode polypeptides that would import the DMSP substrate. Strikingly, these were of two very different types, namely BCCT-type and ABC-type, both of which had been shown in other bacteria to import molecules that closely resemble DMSP.

Thus, the *dddD*-containing clusters of *Halomonas* HTNK1, *Oceanimonas doudoroffii*, *Psychrobacter* J465 and *Pseudomonas* J466 include *dddT*, whose product is a strongly predicted betaine importer of the BCCT (Betaine, Carnitine Choline Transporter) family. These are defined by the Transporter Classification Database (2.A.15) as proteins that transport molecules with quaternary ammonium groups (Saier *et al.*, 2009). They vary in their mode of transport, but nearly all of them depend on Na<sup>+</sup> or H<sup>+</sup> for maintenance of a motive force. Many also contain osmoregulatory or osmosensory properties (Saier *et al.*, 2009). BCCT proteins transport DMSP, in addition to its canonical substrates, likely due to the structural similarity of these compounds (Todd *et al.* 2010; Sun *et al.*, 2011). DddT expressed in an *E. coli* mutant defective in betaine transport conferred the ability to import DMSP. The flexibility of DddT, as with other BCCTs, was also illustrated by the recovery of growth when provided with glycine betaine (Todd *et al.*, 2010; Sun *et al.*, 2011).

In contrast to the examples above, the *dddD* genes of *Burkholderia ambifaria* AMMD, and the α-proteobacteria, *Rhizobium* sp. NGR234, *Rhodobacterales* KLH11 and *Hoeflea phototrophica* DFL-43, are all clustered with genes that encode ATP-binding cassette (ABC) transporters, belonging to sub-families involved in proline/glycine betaine transport (COG2113) and spermidine/putrescine binding transport systems (COG0687). The ABC transporters are massively widespread, and comprise a periplasmic binding substrate binding protein, a membrane-bound transporter and an energy-providing ATP-ase domain (Eitinger *et al., 2011). E. coli* and other enteric bacteria were also able to import DMSP and glycine betaine in response to increased salinity, via a ProU-mediated system (Cosquer *et al., 1999).* The ProU transport system, originally discovered in *Salmonella typhimurium*, imports betaines such as proline in response to osmotic stress (Cairney *et al., 1985; Haardt et al., 1995).* 

#### 1.7.1.5 The DddR, DddZ and DddH regulators

In addition to the genes that encode the enzymes for DMSP and acrylate transport and for the import of DMSP, some regulatory *ddd* genes have been identified.

Firstly, it was noted that a mutation in *dddR* of *Marinomonas* MWYL1 abolished the Ddd<sup>+</sup> phenotype. Furthermore, a *dddD-lacZ* transcriptional fusion was not expressed in the *dddR*<sup>-</sup> mutant strain, whereas in the wild-type background, it was transcribed at high levels, but only in DMSP-grown cells (Johnston *et al.*, 2008). Consistent with this, the DddR protein resembles a LysR-type transcriptional activator, a huge family of regulators that activate expression of their target genes and which, usually, are negatively auto-regulatory (Maddocks and Oyston, 2008). The DddR protein in *Marinomonas* is unusual in that it is positively auto-regulatory, the opposite of most LysR-type regulators (Curson *et al.*, 2011a). This may induce a positive feedback loop that results in rapid consumption of DMSP upon activation.

The *dddR* gene is also found in other species, including *Pseudomonas* J465, which has a highly similar *ddd* gene arrangement to that of *Marinomonas* and also to others, but these are less similar in sequence (Figure 1.6). A cloned DddR-like gene product in *Burkholderia cepacia* AMMD could activate expression of the *Marinomonas dddD* when expressed

heterologously (Todd *et al.*, 2007). A less similar homologue of DddR also exists in *Rhizobium* NGR234, with an identity of only 25% at the peptide level. Indeed, it was termed DddZ and predicted to be involved in the regulation of the divergently transcribed *dddA*-*dddC* genes in that strain (Johnston *et al.*, 2008). The *dddZ* gene became of more interest when a homologue of DddZ was found in *Halomonas* HTNK1, and the gene encoding it was found to be divergently transcribed from the *dddCA-acuNK-dddT* operon, along with another gene, *dddH* (Todd *et al.*, 2010). The DddH protein is very different from the other two regulators, belonging to the TetR family of transcriptional regulators. A role for either of these proteins has yet to be demonstrated directly.

The significant level of HGT that has occurred during the evolution of *dddD* has resulted in copies of this gene being present in many different genera of bacteria. The *dddD* gene appears to be closely linked to growth on DMSP, being prevalent in bacteria isolated on the basis of their ability to use DMSP as a sole C source. Many of these organisms are located in areas of high DMSP concentration, such as corals, seaweeds, marine sediments and the gut of plankton-eating fish (Curson *et al.*, 2010; 2011a).

The appearance of a non-functional *dddD* gene in *R. pomeroyi* suggests a different physiological role for this enzyme from DMSP-dependent DMS production. *R. pomeroyi* DSS-3 already has several DMSP lyases (see later; chapters 4 and 5) and perhaps this form of DddD is responsible for a reaction involving a substrate similar, but not identical to, DMSP; for example, the addition of a CoA moiety to a molecule similar to DMSP (Todd *et al.*, 2011).

Interestingly, some bacterial strains known to cleave DMSP and release DMS were found to lack any detectable homologues of dddD, so must have possessed an alternative enzyme (Johnston *et al.*, 2008).

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#### 1.7.2 DddL

As mentioned previously, DMSP can be cleaved to release equimolar amounts of DMS and acrylate. Following the discovery of the "a-typical" lyase, DddD, I shall now discuss those enzymes that are involved in the more common and direct, DMSP cleavage reaction, starting with DddL.

#### 1.7.2.1 Identification of *dddL* in *Sulfitobacter* EE-36

One of the bacterial strains that was known to release DMS from DMSP but whose genome was seen to lack *dddD*, was *Sulfitobacter* EE-36, which is a member of the Roseobacter clade. A cosmid library of this strain was made in the wide host-range cosmid pLAFR3 (Staskawicz et al., 1987) and one cosmid was obtained that conferred a Ddd<sup>+</sup> phenotype to a strain of Rhizobium leguminosarum. This was chosen as the heterologous host for two reasons; like *Sulfitobacter* it is an  $\alpha$ -proteobacterium and, secondly, it has many  $\sigma$ -factors, and so may favour the expression of heterologous genes from other bacteria (Young et al., 2006). A single gene, *dddL*, sub-cloned form the original cosmid, was sufficient to bestow a Ddd<sup>+</sup> phenotype in *E. coli*. Importantly, high pressure liquid chromatography (HPLC) analysis identified acrylate as the C3 compound released from DMSP, in addition to DMS (Curson et al., 2008). This made DddL the first bona fide DMSP lyase, in the generally accepted definition of this enzymatic activity. DddL had previously been described as a DUF (domain of unknown function), but has a C-terminal domain with similarity to cupins (Curson et al., 2008; Todd et al., 2011; see later). However, Sulfitobacter is unable to grow on DMSP as a sole C source, a clear distinction from the previously mentioned Marinomonas strain.

#### 1.7.2.2 Distribution of DddL homologues

By using DddL as a probe *in silico*, homologues (~50-65% amino acid identity) were found in other genome-sequenced strains of *Rhodobactereceae*, namely *Rhodobacter sphaeroides* 2.4.1, *Loktanella vestfoldensis*, *Oceanicola batsensis* HTCC2597, *Stappia aggregata*, and

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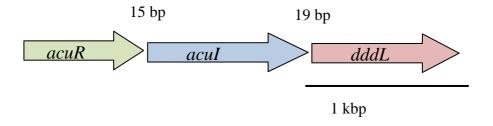
also in *Fulvimarina pelagi* HTCC2506, in the closely related *Aurantimonadaceae* family. It was confirmed that all these strains had a  $Ddd^+$  phenotype (Curson *et al.*, 2008).

*Rhodobacter sphaeroides* is a metabolically diverse, purple, non-sulfur bacterium that can respire through fermentation, anaerobic or aerobic respiration, and can fix molecular  $N_2$  (Ferguson *et al.*, 1987). Also, for decades, it has been a model organism for the study of bacterial photosynthesis (Oh and Kaplan, 2001), but it had never been implicated in DMSP catabolism

#### 1.7.2.3 The *acuR-acuI-dddL* operon

In most of these strains *dddL* was in a one-gene operon, but there was one striking exception, in some strains of *R. sphaeroides*, in which it was the third gene in a three gene operon (Figure 1.8). Two strains (2.4.1. and ATCC17029) of *Rhodobacter sphaeroides* possessed DddL homologues, which were near-identical to each other and 46% identical to DddL of *Sulfitobacter* and both these strains have a Ddd<sup>+</sup> phenotype. In contrast, a third strain of *Rhodobacter sphaeroides* (17025) lacked a *dddL* gene and, reassuringly, did not cleave DMSP (Curson *et al.*, 2008). The *dddL* and neighbouring genes of *R. sphaeroides* 2.4.1 were examined in more detail (Choudhary *et al.*, 2007; Curson *et al.*, 2008; Sullivan *et al.*, 2011).

#### Figure 1.8 Rhodobacter sphaeroides 2.4.1 acuR-acuI-dddL genes



Approximate scale provided. Gene names shown in arrows. Intergenic gaps given in base pairs (bp). Adapted from Sullivan *et al.* (2011).

The promoter-distal gene *dddL*, the central gene, *acuI* and a promoter proximal gene, *acuR* are in a single transcriptional unit (Figure 1.8). The *acuR* gene encodes a TetR-like transcriptional regulator that represses expression of the operon, unless relieved by its co-inducer molecule acrylate. Although growth of the *R. sphaeroides* cells in DMSP also increased the expression of *acuRI-dddL*, this required its conversion to the authentic inducer, acrylate, by the DMSP lyase, DddL. However, neither substrate was sufficient for growth of *R. sphaeroides* as a sole C source (Curson *et al.*, 2008; Sullivan *et al.*, 2011).

From a regulatory point of view, this operon is of particular interest because its mRNA lacks a leader sequence; the 5' end of the transcript coincides with the ATG of *acuR* and therefore lacks a ribosomal binding site. This is highly unusual in bacteria, but may account for the low expression of *acuR* relative to *acuI* and *dddL* as measured by their translational but not transcriptional efficiencies (Sullivan *et al.*, 2011).

Another significant finding from this work was the discovery of the novel gene, *acuI*. AcuI had been (and still is) labelled in the NCBI database as a zinc-dependent alcohol dehydrogenase, in the medium chain dehydrogenase/reductase (MDR) super-family, which has >15,000 members, spread over ~500 peptide families (Persson *et al.*, 2008). This label is a misnomer, as explained further in chapter 5 and chapter 7. Suffice to say here that AcuI has a role in the processing of acrylate in the organisms that harbour it. This was first indicated by Sullivan *et al.* (2011), who observed that AcuI mutants of *R. sphaeroides* 2.4.1 were (a) more sensitive to the toxic effects of acrylate in the growth medium and (b) reduced in their ability to undertake partial catabolism of radio-labelled acrylate.

Other genome-sequenced bacteria were known to make DMS from DMSP, and yet these contained neither *dddD* nor *dddL*. Thus, there were yet other versions of DMSP lyase to be found. One such enzyme was DddP, first identified in a strain of the Roseobacter clade, as follows.

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#### 1.7.3 DddP

DddP was discovered in the  $\alpha$ -proteobacterium Roseobacter, *Roseovarius nubinhibens* ISM, which had been shown to have a Ddd<sup>+</sup> phenotype (Todd *et al.*, 2009) but whose genome lacked *dddL* or *dddD* (Todd *et al.*, 2007; 2009). To identify the gene for the novel lyase in this strain, a cosmid from a newly constructed genomic library was identified that conferred a Ddd<sup>+</sup> phenotype to the heterologous host, *Rhizobium leguminosarum* strain J391, just as was done to identify *dddL* (above). The gene responsible for the phenotype was localised by sub-cloning and sequencing and was termed *dddP*.

#### 1.7.3.1 Notable features of DddP

The DddP enzyme was shown to be a "classical" DMSP lyase, since *E. coli* strains containing *dddP* cloned in an expression plasmid cleaved DMSP into DMS plus acrylate (Todd *et al.*, 2009).Expression of *dddP* is enhanced, *ca*. 4-fold in cells of *Roseovarius nubinhibens* ISM that were grown in DMSP-supplemented medium, but, unlike *dddL* of *Rhodobacter sphaeroides*, its expression was not induced by acrylate (Todd *et al.*, 2009; chapter 2). DddP is a member of the M24B family (COG0006) of peptidases, as shown in the MEROPS database (Rawlings *et al.*, 2010). These enzymes are often metallopeptidases, although work described in chapter 2 will investigate this label (or rather, misnomer!) in further detail.

#### 1.7.3.2 Distribution of DddP

Homologues of DddP were found in several, but not all Roseobacters – including the "model strain" *Ruegeria pomeroyi* DSS-3. Other  $\alpha$ -proteobacteria include *Mesorhizobium* spp. and >55% identity was found to homologues in the genomes of the  $\gamma$ -proteobacteria, *Oceanimonas doudoroffii, Vibrio orientalis* and *Pseudomonas putida*. Thus, DddP appears to be present in both terrestrial and marine organisms. Interestingly, *O. doudoroffii* possesses two very different orthologues of DddP, both of which are active and confer a Ddd<sup>+</sup> phenotype when cloned, individually into *E. coli* (Todd *et al.,* 2009; Curson *et al.,* 2011b). Incidentally, this species also has a functional DddD enzyme.

An exciting discovery was that the *dddP* gene also occurred in some species of fungi and in several cases, these were confirmed as having DMSP lyase activity (Todd *et al.*, 2009). Homologues (~53% identity at peptide level) included members of the Sordariomycetes, *Fusarium graminearum*, *F. culmorum* and *F.oxysporum* f.sp. *lycopersici*. These species are well known as the causative agents of a number of crop diseases, such as head blight of cereals and tomato wilt (Sutton, 1982; Snijders *et al.*, 1990; Cooper *et al.*, 1978). Other homologues (also ~53% identity at peptide level) were found in some Euritiomycetes, such as *Aspergillus oryzae* RIB40 and *A. flavus*. These two *Aspergillus* strains are very similar in their genomic structure, differing in only a few instances, such as the presence of a suite of aflotoxin (*aflR*) production genes in *A. flavus* (Watson *et al.*, 1999). There is a very close similarity between the DddP sequences from *Aspergillus* and *Fusarium* species (~90% identity). The bacterial (i.e. *Roseovarius nubinhibens*) and fungal proteins are also similar in their peptide sequence, with ~54% identity to one another, despite their last common ancestor being ~2.7 billion years ago (Nei *et al.*, 2001). The *Aspergillus* and *Fusarium* copies of the gene lack introns, pointing to a relatively recent acquisition of *dddP* (Nielsen *et al.*, 2004).

In fact, the presence of a fungal 'lyase' was inferred several years earlier from the DMS released by fungi, predominantly Ascomycetes, associated with the rhizosphere of the DMSP synthesising angiosperm, *Spartina alterniflora*, and other salt marsh grasses (Otte, 2004; Bacic *et al.*, 1998). One of these, expressed by *Fusarium lateritium*, was later characterised as a DMSP lyase, and shown to be induced by DMSP (Bacic and Yoch, 1998).

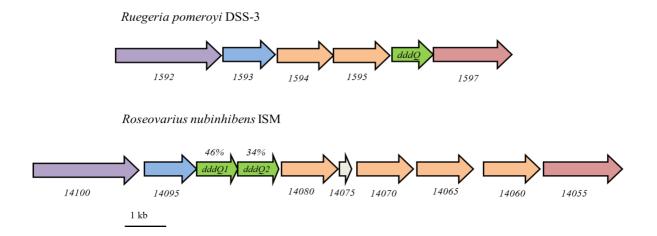
A GOS metagenome database interrogation using DddP from *R. nubinhibens* (Todd *et al.,* 2009) resulted in 92 matches (<  $e^{-80}$ ). The majority (83) of these matches clustered into a single group, which represented proteins in many of the marine  $\alpha$ -proteobacteria, and the fungal representatives clustered together in another subgroup. Interestingly, *dddP* was also found in viral metagenomes isolated from environments as diverse as the Arctic Ocean and a coral reef on the equatorial Pacific Ocean island Kiritimati (Raina *et al.,* 2010).

#### 1.7.4 DddQ

A DddP<sup>-</sup> mutant strain of *Roseovarius nubinhibens* ISM, created for the work mentioned above, still had a Ddd<sup>+</sup> phenotype, albeit at much reduced (~10%) of the wild type level (Todd *et al.*, 2011), suggesting that it had at least one other DMSP lyase. To identify this putative gene, a similar approach to the one used to identify *dddP* was used. Thus, the *Roseovarius nubinhibens* ISM pLAFR3 cosmid library was mobilized into *Rhizobium leguminosarum* strain J391, and the resulting transconjugants were individually assayed for DMS production when provided with DMSP (Todd *et al.*, 2011).

Sequencing of a cloned cosmid that conferred a Ddd<sup>+</sup> phenotype to *Rhizobium*, revealed the presence of two adjacent genes, termed dddQ1 and dddQ2, which were predicted to be in a large, 10 gene transcriptional unit (Figure 1.9). The DddQ1 and dddQ2 gene products were 39% identical to each other and were of interest because, like DddL (above), they had a cupin motif. Indeed, when each of these genes was cloned, alone, in *E. coli*, they conferred the ability to catabolise DMSP, forming DMS and acrylate as the products. The functions of the other genes in the operon are not known but it was noted that there was a similar gene arrangement in *Ruegeria pomeroyi*. However, in this strain there was only one copy of dddQ and only two copies of the enolase superfamily genes, compared to the four in *Roseovarius nubinhibens* (Figure 1.9).

### Figure 1.9 Location of *dddQ* genes in *Roseovarius nubinhibens* ISM and *Ruegeria pomeroyi* DSS-3



Genes in *Roseovarius nubinhibens* ISM and *Ruegeria pomeroyi* DSS-3 are shown as arrows, with their gene tags, but lacking the prefixes 'SPO' and 'ISM\_' respectively. The *R. pomeroyi dddQ* (*SPO1596*) and the *R. nubinhibens dddQ1* (*ISM\_14090*) and *dddQ2* (*ISM\_14085*) genes are coloured green. The %age identities between DddQ and either DddQ1 or DddQ2 are shown above the *R. nubinhibens* genes.

#### Genes for:

- aminomethyl transferase family protein (SPO1592 and ISM\_14100) are purple
- Zn alcohol dehydrogenase family (SPO1593 and ISM\_14095) are blue,
- acetyl ornithine aminotransferase family (SPO1597 and ISM\_14055) are red
- ISM\_14075 of R. nubinhibens (absent from R. pomeroyi) encodes a Cupin-2-like protein and is grey.
- enolase super-family are orange, the two versions in *R. pomeroyi* (SPO1594 and SPO1595) being most similar, respectively, to ISM\_14080 and ISM\_14065 of *R. nubinhibens*.
   Adapted from Todd *et al.*, 2011.

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Given the close phylogenetic relationship between the two species, and the level of synteny between the two genome regions, the dddQ genes are surprisingly dissimilar from one another; the dddQ gene product from *Ruegeria pomeroyi* shares only 46% and 34% identity with the products of dddQ1 and dddQ2 from *Roseovarius nubinhibens*, respectively, and much of this identity relates to the cupin domain present in both homologues.

#### 1.7.4.1 DddL and DddQ Cupin domains

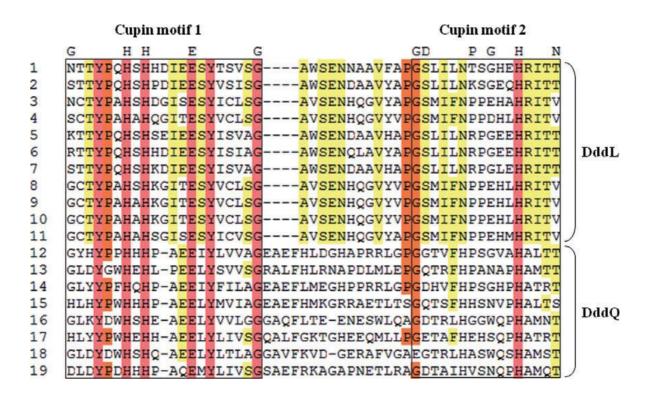
Cupins (Latin "cupa" means "small barrel"), are a super-family of domains with a  $\beta$ -barrel structure (Dunwell, 1998; Dunwell and Gane, 1998). They are functionally diverse, with many examples in all three Domains of life, being initially discovered as germin-like proteins from wheat. Typically, a cupin domain has two conserved motifs, 1 and 2, as shown below:

Motif 1: (X)<sub>5</sub>**H**X**H**(X)<sub>3.4</sub>E(X)<sub>6</sub>**G** 

Motif 2:  $G(X)_5 PXG(X)_2 H(X)_3 N$ 

These motifs are separated by an inter-motif region, between 11 and 100 amino acids in length, depending on the organism and function (Dunwell *et al.*, 2000; 2001; 2004). Cupins are generally metal-binding proteins, with the predicted metal-binding residues being two histidine and one glutamine residues within motif 1, and a single histidine in motif 2 (shown in bold in the above sequences), acting as ligands to bind a variety of metals, including manganese, nickel, zinc, iron and copper (Woo *et al.*, 2000). Different proteins may contain one or more cupin domains, known as mono-, bi- or even multi- cupin proteins. Chapter 4 has further discussion of cupins in other proteins, particularly the novel DMSP lyases.

# Figure 1.10 Alignment of cupin regions of selected DddL and DddQ polypeptides



The region spanning the conserved cupin motifs 1 and 2 (boxed) is shown, with residues conserved in the cupin superfamily (Dunwell *et al.*, 2004) shown above the sequence. Residues marked in red, orange and yellow are, respectively, 100%, >80% and >60% identical in all the sequences. Sequences are from: 1 = Dinoroseobacter shibae DFL 12, DshiDRAFT\_1825; 2 = Fulvimarina pelagi HTCC2506, FP2506\_12684; 3 = Labrenzia alexandrii DFL-11, SADFL11\_5101; 4 = Loktanella vestfoldensis SKA53, SKA53\_01756; 5 = Oceanicola batsensis HTCC2597, OB2597\_08014; 6 = Rhodobacterales HTCC2654, HTCC2654\_RB2654; 7 = Rhodobacter sphaeroides 2.4.1, Rsp\_1433; 8 = Roseobacter GAI101, RGAI101\_3508; 9 = Sulfitobacter sp. EE-36, EE36\_11918; 10 = Sulfitobacter sp. NAS-14.1, NAS141\_17149; 11 = Stappia aggregata IAM 12614, SIAM614\_20126; 12 = Roseovarius nubinhibens ISM, ISM\_14090; 13 = R. nubinhibens ISM, ISM\_14085; 14 = Ruegeria pomeroyi DSS-3, SPO1596;  $15 = Rhodobacterales bacterium HTCC2150, RB2150_06543; 16 = Rhodobacterales bacterium Y4I, RBY4I_2503; <math>17 = Roseobacter$  sp. SK209-2-6, RSK20926\_17292; 18 = Silicibacter lacuscaerulensis ITI\_1157, SL1157\_0332; 19 = Thalassiobium sp. R2A62, TR2A62\_3487.

From Todd et al. (2011).

The DddL and DddQ homologues share many of the conserved residues of cupin domain proteins (Figure 1.10) (Dunwell *et al.*, 2004; Todd *et al.*, 2011). It is not known how the separate proteins evolved, either through convergent or divergent evolution. However, it is clear that the cupin domain is important for cleavage of DMSP.

#### 1.7.4.2 Distribution of DddQ

An NCBI BLASTp found DddQ homologues in a few other Roseobacters, including *Silicibacter lacuscaerulensis*, and also in a SAR116 Clade  $\alpha$ -proteobacterium, HIMB100. Due to the diversity between the groups of these enzymes, a relatively low cut-off value of NCBI bit score >109 was used to interrogate the GOS with DddQ. This retrieved >200 matches and these primarily clustered into 3 main groups (Todd *et al.*, 2011). Given the diversity of DddQ, even ratified ones, it is not clear which of these GOS sequences are *bona fide* DMSP lyases. However, one gene from each group was chosen and synthesised *in vitro*, before being cloned into pBluescript and transformed into *E. coli*. All 3 conferred a Ddd<sup>+</sup> phenotype (Todd *et al.*, 2011).

#### 1.7.5 DddY

Following the discovery of DddQ, the gene for another DMSP lyase, DddY, was identified, but this time, it was one that had been encountered before, from a biochemical approach. DddY was initially found in the  $\beta$ -proteobacterium *Alcaligenes faecalis* M3A, isolated from intertidal sediment containing the DMSP-producing cordgrass, *Spartina* (de Souza and Yoch, 1995a). Prior work had shown that this bacterium could convert DMSP to acrylate, and could also use both of these compounds as carbon sources (Ansede *et al.*, 1999). In a demanding series of experiments, Yoch *et al.* purified the DMSP lyase from *A. faecalis* M3A and even obtained its N-terminal sequence, but they did not extend the work to identify the corresponding gene (de Souza and Yoch, 1995a; 1995b; 1996; Yoch *et al.*,1997). This approach was only undertaken some 14 years later, by Curson *et al.* (2011c), and has led to some intriguing outcomes on what is another, very different system for DMSP catabolism.

#### 1.7.5.1 DddY, a membrane-bound lyase

As before, the isolation of *dddY* involved the construction of a library in a wide host-range cosmid, followed by the screening for function in a heterologous host. In this case, the host was Pseudomonas putida and transconjugants containing the library of cosmids were selected for growth on DMSP as sole C source. Following the sub-cloning from a cosmid that conferred this property, a fragment, and then a single gene, termed dddY, could confer a Ddd<sup>+</sup> phenotype to E. coli, converting DMSP into DMS plus acrylate. Further, a mutation in dddY in A. faecalis M3A itself abolished its ability to make DMS from this substrate (Curson et al., 2011c). Reassuringly, the deduced *dddY* gene product contains an N-terminal sequence which is strongly predicted to include an N-terminal leader that would guide it to the bacterial periplasm. This is in keeping with the earlier suggestion by Ansede et al. (1999) that the lyase in A. faecalis M3A was associated with the cell surface and this was further substantiated by Curson et al., (2011c) who showed by sub-cellular fractionation that the lyase was associated with the periplasm. And, most satisfying of all, when the N-terminal sequence was cleaved in silico at the predicted cleavage site (21 amino acids from the N-terminus) this generated a polypeptide whose deduced sequence matched the one that had been described by de Souza and Yoch (1996) for the mature, functional enzyme. Other than that, there were no sequencebased clues on the enzymatic function of DddY, as demonstrated by the fact that, hitherto, it had been annotated as a Domain of Unknown Function (DUF), with no similarity to any other polypeptide of defined enzymatic activity

#### 1.7.5.2 Distribution of DddY

Following a search of the NCBI database, the DddY polypeptide was not seen in any other species of *Alcaligenes*, but homologues with identities of ~35% were found in certain members of the  $\gamma$ -proteobacteria, *Shewanella* spp. (Curson *et al.*, 2011c). A DddY homologue was also found in the  $\varepsilon$ -proteobacterium *Arcobacter nitrofigilis* DSM7299 (Pati *et al.*, 2010). The sequences of the *Shewanella* and *Arcobacter* strains were ~75% identical to each other and, like that of *A. faecalis*, were strongly predicted to be periplasmic. All of these strains exhibited Ddd<sup>+</sup> phenotypes, meaning that *Arcobacter nitrofigilis* is the first demonstrated  $\varepsilon$ -proteobacterium with this ability (Curson *et al.*, 2011c). Unsurprisingly, *A.* 

nitrofigilis was isolated from the roots of the DMSP synthesising salt marsh plant, Spartina alterniflora (McClung et al., 1983).

Just as they had done with *A. faecalis*, deSouza and Yoch (1996) purified a DMSP lyase from the  $\gamma$ -proteobacterium, *Oceanimonas doudoroffii*, and found that its N-terminus was very similar to that of the *A. faecalis* DddY. However, the near-complete genome of this strain lacks a *dddY* gene (Curson *et al.*, 2011c), although it does contain *dddD* and two versions of *dddP* (see above). The reason for this discrepancy is not known.

There were no detectable DddY homologues in the GOS, or any other, metagenomic database.

#### 1.7.5.3 DddY and acrylate metabolism

Examination of the region close to *dddY* of *A. faecalis* M3A revealed several other genes that corresponded to those in the *ddd* gene clusters of other bacteria, notably *Halomonas* HTNK1 (Figure 1.6). Thus, both these strains contained *dddA*, *dddC*, *acuN* and *acuK*, all of whose corresponding products are ~70% identical in the two species. However, there were two obvious and explicable differences. First, the two strains differed in the nature of the genes that encoded the initial DMSP lyase – *dddY* for *Alcaligenes* and *dddD* for *Halomonas*. Second, the *ddd* cluster of *Halomonas* includes the transporter gene *dddT* but *Alcaligenes* does not. This, of course, is fully consistent with the fact that DddY is located in the periplasm, so there is no need for a dedicated import system for the substrate.

It is striking that both gene clusters include the *acuNK* genes, which are involved in the conversion of acrylate to 3HP (see above). Whereas the AcuNK enzymes of *Halomonas* need only act on exogenously supplied acrylate (DddD does not generate this product from DMSP), in the case of *Alcaligenes* can also be generated via the action of DddY on DMSP. Whatever the source, the resultant 3HP can then be used as the substrate for further catabolism, via DddC and DddA, as described above for *Halomonas*, Interestingly, an intermediate in AcuNK mediated acrylate catabolism is acryloyl-CoA, and this happens to be

the cognate molecule for the AcuI enzyme, encoded by the nearby *acuI* gene in *Halomonas* (see above; chapter 5).

Alcaligenes contains homologues of the previously mentioned dddZ and acuR genes from *Halomonas* HTNK1 (Curson *et al.*, 2011c) and indeed, an *acuI* gene whose product is 60% and 65% identical to the *Rhodobacter sphaeroides* 2.4.1. and *Halomonas* HTNK1 proteins, respectively. AcuR appears to act as a repressor in *A. faecalis* also, with an AcuR<sup>-</sup> mutant exhibiting a ~11 fold increase in DMS production. Growth experiments illustrated that the DddZ protein may be responsible for acrylate catabolism, via regulation of AcuN, AcuK, or perhaps downstream acrylate catabolic enzymes. Transcriptional *lacZ* fusions showed that the *dddY* operon was up-regulated in response to acrylate, and its precursor, DMSP, but no induction of *dddZ* was noted.

Interestingly, *A. faecalis* may also have an alternative method of catabolising DMSP, since DddY<sup>-</sup> strains could still use DMSP as a sole C source, despite producing significantly less DMS from DMSP.

#### 1.8 Current knowledge

These recent genetic studies have provided something of a sea-change in our understanding of bacterial DMSP catabolism, and there are a number of key points to remember, going forward.

- 1. Diversity The genes and enzymes responsible for DMSP degradation are varied and often unique, with at least 6 classified DMSP lyases that display several different characteristics.
- Distribution The DMSP lyases can be found in a wide range of bacteria, particularly the marine Roseobacters. Additionally, the presence of these genes in species of fungi and phytoplankton implies an important role for this pathway throughout the Kingdoms.
- 3. Horizontal Gene Transfer These genes have likely undergone considerable levels of HGT between bacterial species, and even between the Domains.
- 4. Multiple systems in single strains The possession of multiple DMSP degradation systems in certain species (e.g. *Ruegeria pomeroyi* which harbours at least *dddP*, *dddQ*, *dddW* and *dmdA* see chapters 4, 5 and 7) implies a high level of adaptation to the DMSP molecule and has much potential for studies into the portioning of these systems *in vivo*.
- Odd regulation The unusual regulation observed in some of these systems, such as the acrylate mediated induction of *dddL* in *Rhodobacter sphaeroides* 2.4.1. point to more complicated control of these genes than first thought.

#### 1.9 Aims and objectives

In 2008, when this project started, the full extent of the diversity and complexity of the DMSP catabolic pathways was only beginning to emerge. It was clear, though, that the initial genetic studies had formed the platform for a set of new studies, aimed at clarifying and extending what was known in terms of the biochemistry, physiology, genetics, genomics, ecology and even the evolution of these processes.

My project therefore was to gain more information on the mechanism and regulation of DMSP in bacteria, in particular in the Roseobacter clade, for which DMSP catabolism is an important part of their nutritional lifestyle. The experimental programme was to include genetics, bioinformatics, transcriptomics and bacterial physiology.

There exists a significant pool of knowledge concerning the enzymes involved in the degradation of the osmolyte DMSP. The demethylase and lyases have been examined in closer detail, illustrating the remarkable leaps in understanding that have occurred in a relatively short period of time. Despite attempts to investigate the regulation of genes in

response to DMSP in bacterial communities and in particular species, such as *R. pomeroyi* DSS-3, there are still areas that would benefit from further research. Until now, the effects of the DMSP degradation products at a global transcriptomics level have been neglected, discounting the cells response to compounds such as acrylate and DMS, which may provide greater insight into the intricacies of the metabolic regulation that occurs in *Ruegeria pomeroyi* DSS-3, a model organism that utilizes both mechanisms of DMSP degradation and multiple DMSP lyase pathways.

The work in this thesis will address the following:

- To over-express, purify and further characterise, biochemically, the DddP DMSP lyase from the Roseobacter, *Roseovarius nubinhibens* ISM, and to investigate the presence of homologous enzymes in novel fungal species.
- To use transcriptomics to investigate the response of *Ruegeria pomeroyi* DSS-3 to the osmolyte DMSP, and two of its degradation products, acrylate and DMS.
- To identify and ratify novel genes and enzymes involved in DMSP metabolism through bioinformatics and molecular genetic approaches, including the use of transcriptional fusions, RT-PCR and mutations.
- To begin an investigation of the regulatory mechanisms that effect transcriptional responses to DMSP and/or its catabolites.
- To begin to understand the complex interactions between several apparently autonomous metabolic pathways in response to DMSP, and to come to conclusions as to the evolution of such systems in *Ruegeria pomeroyi* DSS-3, and other DMSP utilising marine bacteria.

# **Chapter 2**

## The DMSP lyase, DddP

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#### **2.1 Introduction**

Prior to the studies into the genome-wide response of *Ruegeria pomeroyi* to DMSP, DMS and acrylate, I was involved in some much more specific work regarding one particular DMSP lyase, DddP. As discussed in the introduction, work by Todd *et al.* (2009) had identified a DMSP lyase in *Roseovarius nubinhibens* ISM and shown that it conferred a Ddd<sup>+</sup> phenotype when the corresponding gene, *dddP*, was expressed heterologously in other bacteria. Following on from this discovery I wished to ratify the function of DddP as a *bona fide* DMSP lyase that cleaved DMSP into acrylate plus DMS *in vitro*.

#### **2.2 Results**

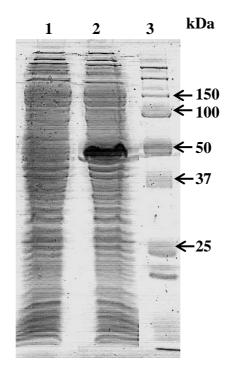
#### 2.2.1 Over-expression of the Roseovarius nuibinhibens ISM DddP protein

To characterise the DddP polypeptide and confirm its function as a DMSP lyase, I set out to purify it to an acceptable level of homogeneity. This was done for the *Roseovarius nubinhibens* version of the enzyme, since this strain had (at the time) become something of a model for the Roseobacter clade in the UEA laboratory, as well as elsewhere. A purification protocol was developed as follows.

A 1392 bp fragment containing the intact *dddP* gene was amplified from *R. nubinhibens*, using primers DddPF and DddPR, and cloned into the pET21a plasmid expression vector (Novagen), to form the recombinant plasmid pBIO1658, which was transformed into *E. coli* strain BL21 pLysS (Miroux and Walker, 1996). The pET21a plasmid contains a *lac1* repressor gene, a viral T7 promoter, a polylinker site and Amp<sup>R</sup>. Thus, when isopropyl  $\beta$ -D-1-thiogalactopyranoside (IPTG) is added to cells containing this plasmid, rapid and significant transcription of the inserted DNA occurs, making it a useful tool for heterologous gene expression. This initial cloning was done by J. D. Todd (see Todd *et al.*, 2009).

To obtain significant amounts of the DddP polypeptide, a 100ml culture of the *E. coli*.: pBIO1658 strain was grown at 37°C in LB containing ampicillin to an OD<sub>600nm</sub> of 0.4-0.6. Then, 10  $\mu$ M IPTG was added to induce expression of the cloned *dddP* gene and the culture was further incubated at 25°C until stationary phase. The cells were was pelleted by centrifugation and re-suspended in TRIS buffer, before being lysed by sonication. This initial extract was run on an SDS-PAGE gel and a ~50 kDa band was observed to be significantly more abundant than any other. This corresponded to the predicted molecular weight of DddP, at 49,972 kDa, and the band was not seen in empty vector (see Figure 2.1).

### Figure 2.1 Cell lysate from *E. coli* cultures expressing empty pET21a vector and vector containing *Roseovarius nubinhibens* ISM DddP, separated by SDS-PAGE



Polypeptides were separated by SDS-PAGE on 12% acrylamide gels and stained with Coomassie Blue. Lanes: 1, crude cell extract from *E. coli* cells containing empty pET21a vector; 2, crude cell extract from *E. coli* cells induced for expression of *R. nubinhibens* DddP (pBIO1658) 3, broad-range protein marker (Bio-Rad).

#### 2.2.2 Purification of DddP from E. coli expressing pBIO1658

The cell lysate was then partially precipitated using  $(NH_4)_2SO_4$ . Increasing the salt concentration, and thus ionic strength, causes proteins to become insoluble and precipitate. An iterative process was initially used to determine at which concentration the  $(NH_4)_2SO_4$  would

pull out as much protein from the lysate as possible, without precipitating DddP. And so, addition of 25% (w/v) (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> precipitated a proportion (*ca.* 40%) of the proteins without removing DddP itself (Figure 2.2, lane 3). The resulting lysate was centrifuged and the supernatant subjected to three separate column chromatograph steps. At each stage of purification, an absorbance trace was used to determine which fractions contained protein, and these were checked using SDS-PAGE for the presence of an appropriate band. DMS assays were also carried out on the pooled fractions to confirm that DddP retained activity.

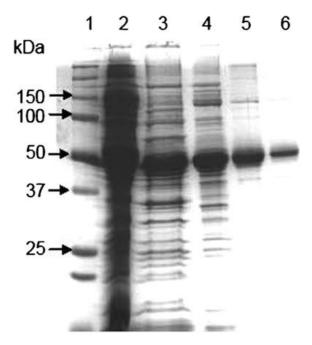
First, the supernatant was run through a phenyl-sepharose high performance column, which utilises the hydrophobic chromatography principle (Hofstee, 1973). The hydrophobic ligand (phenyl-sepharose) reversibly binds the hydrophobic surface residues on proteins in high salt concentrations. As the salt concentration decreases, proteins disassociate and are eluted. Therefore, the column was equilibrated with 20 mM Tris containing 25% (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>. DddP was then eluted by applying a 25% – 0% (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> gradient (flow rate 3 ml min<sup>-1</sup>).

There were three DddP-containing fractions, each of 3 ml (Figure 2.2, lane 4). These were pooled and applied to a DEAE anion exchange column equilibrated with 20 mM Tris, pH8. This separated the proteins on the basis of their charge. DddP has a predicted isoelectric point (pI) of ~5.1 (ExPASy Compute - Bjellqvist *et al.*, 1993), and so at pH8 has a net negative charge, making it ideal for binding to a positively charged solid phase, as used in anion exchange chromatography (Williams and Frasca, 2001). A 0 - 1M NaCl gradient was applied (flow rate 5 ml min<sup>-1</sup>), increasing the ionic strength and causing proteins to elute.

Again, three DddP-containing fractions, each of 2 ml, were retrieved and pooled (Figure 2.2, lane 5), then applied to the final, gel filtration column, to separate the remaining proteins on the basis of molecular weight (Lathe and Ruthven, 1955). To do this, the pooled fractions were concentrated using an Amicon Ultra centrifugal filter (Millipore), loaded onto a Superdex 200 column equilibrated with 50 mM MES buffer, and run through at a flow volume of 1 ml min<sup>-1</sup>. Two DddP-containing fractions, of 5 ml each, were pooled and stored at 4°C for further use, with no detectable loss in activity, even after several weeks.

As shown in Figure 2.2, lane 6, the final step yielded a single visible band on Coomassie Blue with an approximate purity of >95%, and yielded ~ 50 mg DddP L<sup>-1</sup> of initial *E. coli* culture, as estimated from  $A_{280}$  measurements, using the protein extinction coefficient, 76,860 M<sup>-1</sup> cm<sup>-1</sup>, calculated from the tyrosine, tryptophan and cysteine residues in the amino acid sequence (Gill and von Hippel, 1989).

### Figure 2.2 Purified *Roseovarius nubinhibens* ISM DddP protein from *E. coli* cultures, separated by SDS-PAGE



Polypeptides were separated by SDS-PAGE on 12% acrylamide gels and stained with Coomassie Blue. Lanes: 1, broad-range protein marker (Bio-Rad); 2, crude cell extract after sonication; 3, supernatant after treating cell extract with 25% (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>; 4, phenyl sepharose column eluate; 5, DEAE column eluate; 6, gel filtration column eluate. From Kirkwood *et al.* (2011b).

#### 2.2.3 Characterisation of the purified DddP enzyme

#### 2.2.3.1 Effect of pH and temperature on DddP enzymatic activity

Prior to characterisation and enzyme kinetic studies, the effects of pH and temperature on DddP activity were ascertained, as follows. DMS assays using GC were carried out as above, with the enzyme being in MBS buffer over a range of pH values, from 2.0 to 9.0 (see Figure 2.3a).

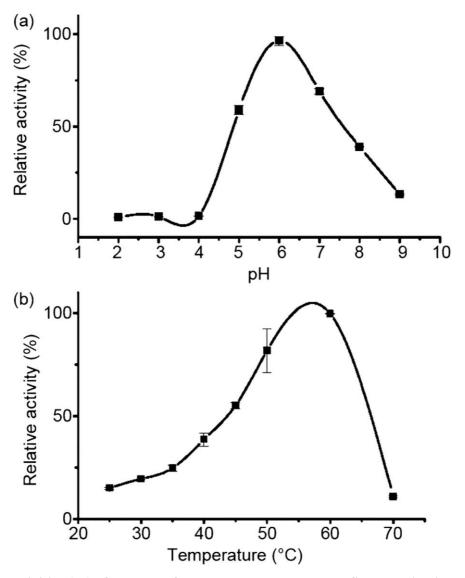


Figure 2.3 Effect of pH and temperature on DddP activity

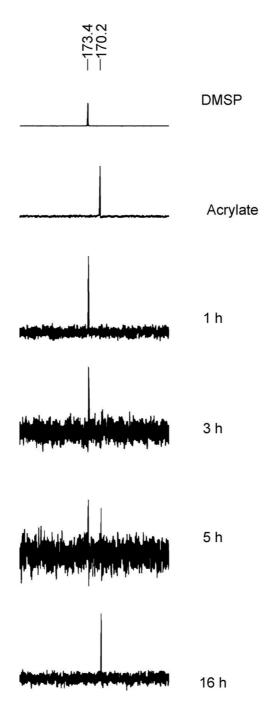
The relative activities (%) of samples of DddP were assayed by DMS production in solutions at different pH (a), and then in 50 mM MES buffer, pH 6.0 at different temperatures (b). Error bars calculated from duplicate experiments.

The highest DMS activity was observed at pH 6.0, which was then used for further studies, including the temperature-responsive activity of the DddP enzyme *in vitro* which showed that activity was lost at temperatures >60°C. All subsequent work was carried out at 30°C, to reflect the *in vivo* conditions (*R. nubinhibens* ISM is routinely cultured at 30°C).

#### 2.2.3.2 Confirmation of DMSP lyase activity

To determine the *in vitro* activity of DddP, samples of purified protein (50  $\mu$ l, 0.9  $\mu$ M) amended with 5 mM labelled [1-<sup>13</sup>C]DMSP were incubated for 16 hours. The <sup>13</sup>C catabolite was identified by nuclear magnetic resonance spectroscopy (NMR) and confirmed that the DMSP had been converted to a newly formed labelled compound, which exhibited an identical peak at the chemical shift of a reference sample of acrylate (Figure 2.4). This work was conducted as described in Todd *et al.* (2010).

# Figure 2.4 NMR illustrating depletion of [1-<sup>13</sup>C]DMSP and appearance of labelled acrylate when in the presence of purified DddP lyase



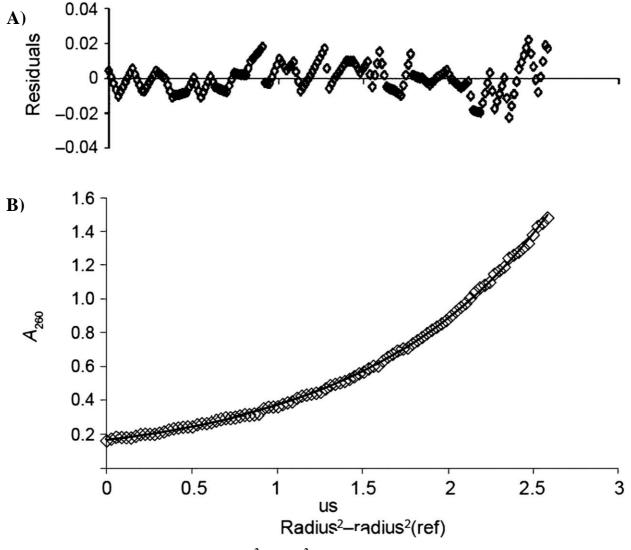
NMR spectra of the products of DddP-mediated cleavage of  $[1-^{13}C]DMSP$ . The <sup>13</sup>C NMR spectra for pure samples of DMSP and acrylate are shown as indicated, with peaks at chemical shifts of 173.4 and 170.2, respectively. The spectra below the reference compounds are from reaction mixes containing 5 mM [1-<sup>13</sup>C]DMSP plus 0.9  $\mu$ M DddP enzyme, sampled after 1, 3, 5 and 16 h incubation, as indicated.

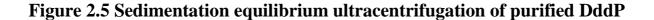
To confirm that DMS was the concomitant product, gas chromatography (GC) was used. A sample of DddP (~10  $\mu$ M) was incubated in a 1 ml screw cap vial in buffer containing 5 mM DMSP, at 30°C. Peaks corresponding to DMS were subsequently observed. This confirms the function of DddP as a conventional lyase, which cleaves DMSP to DMS and acrylate. Approximate estimated rates of production were 0.07 and 0.185 nmol/µg/min for DMS and acrylate, respectively.

#### 2.2.3.3 Association state of DddP

As expected from its deduced  $M_r$  (49, 972 Da), the purified DddP polypeptide was seen as a ~50 kDa band in denaturing gels. To ascertain the association state of functional DddP, this was determined using sedimentation equilibrium analytical ultracentrifugation. This technique works by centrifuging a sample of protein until the opposing forces of diffusion and sedimentation are equal, with different species (of different  $M_r$ ) reaching equilibrium at different times (Cole *et al.*, 2008). This is recorded by a series of scans, and the data can then be applied to known models to ascertain structural composition of the protein. Thus, a sample of DddP (~5  $\mu$ M) was spun at 12,000 rpm at 20°C in a Beckman Optima XL-1 analytical ultracentrifuge. Scans were taken every 4 hours to determine when equilibrium had been reached, at which point 5 more scans were recorded. The data were analysed (Ultrascan; Demeler, 2005) and fitted well to a one-component model, which predicted a  $M_r$  of 95, 300 ± 8000 Da, suggesting that DddP is a homodimer in solution (Figure 2.5).

The addition of 0.45 M NaCl had no obvious effect on the association state of the homodimer, implying that the interaction between the two polypeptides is relatively stable.



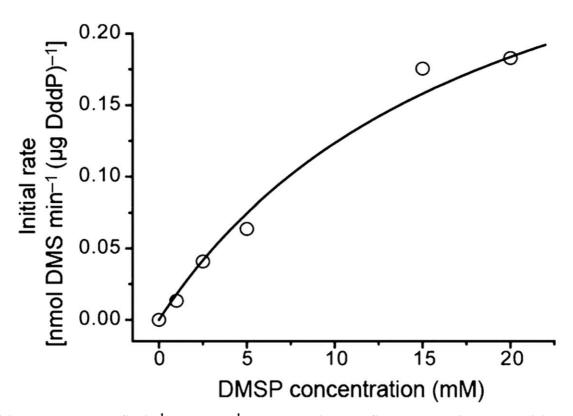


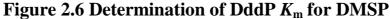
Absorbance at 260 nm against radius<sup>2</sup>-radius<sup>2</sup>(ref) following equilibration of purified DddP (~5  $\mu$ M) at 20°C and 12,000 rpm. A) Residual difference between experimental data and fitted curve. B) A mean molecular weight of 95,300 ± 8000 Da was recorded when data were fitted to a single species (solid line).

The structure of another MEROPS M24B family member, namely creatinase from *Actinobacillus* (Padmanabhan *et al.*, 2002), has been solved and also shows a dimeric enzyme. However, most M24 family members, such as methionine aminopeptidases (M24A – EC: 3.4.11.18) are monomeric.

#### 2.2.3.4 DddP enzyme kinetics

To determine the K<sub>m</sub> and V<sub>max</sub> values of the DddP enzyme preparation, the initial rates of DMS production were measured at intervals using ~0.3  $\mu$ M DddP incubated at 30°C in MES buffer (pH 6.0), between 1 mM and 20 mM DMSP. The data fitted reasonably well to the Michaelis-Menten equation (Figure 2.6) and a  $K_m$  of DddP for DMSP of 13.8 ± 5.5 mM and a V<sub>max</sub> of 0.31 ±0.06 nmol DMS min<sup>-1</sup> ( $\mu$ g DddP)<sup>-1</sup> was extrapolated.





Initial rate (nmol DMS min<sup>-1</sup> (μg DddP)<sup>-1</sup>) plotted against DMSP concentration. DddP (0.3 μM) was in 50 mM MES, pH 6.0. Solid line represents a fit to the Michaelis-Menten equation.

A value of  $K_m$  value of DddP of ~14 mM seemed rather high, but the  $K_m$  of another M24B family member, creatinase is also high, at 13.3 mM for the wild type enzyme in *Pseudomonas putida* (Schumann *et al.*, 1993). Furthermore, the DmdA demethylase enzymes of *Ruegeria pomeroyi* and of *Candidatus* Pelagibacter ubique exhibit high  $K_m$  values, at 5.4 mM and 13.2 mM, respectively (Reisch *et al.*, 2008). To recall, DMSP concentrations in the open oceans can be as little as ~1-2 nM (van Duyl *et al.*, 1998), whereas significant accumulation of

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DMSP occurs intracellularly in these organisms; as high as 70 mM (Reisch *et al.*, 2008). The use of DMSP as an osmoprotectant is therefore likely metabolically linked to its catabolism by the DMSP lyases such as DddP.

## 2.2.3.5 Metal-binding studies

Several, but not all, polypeptides in this M24B family have been shown to bind metal (usually Co or Mn) cofactors (Graham *et al.*, 2005), so I set out to determine if DddP was also metalbinding. To do this, the assays used material that contained metal-chelating compounds. Both 2,2'-bipyridyl (2.5 mM) and ethylene-diaminetetra-acetic acid (EDTA- 25 mM) were added separately to DddP samples and incubated for 15 minutes at room temperature, before adding 5 mM DMSP and carrying out GC DMS assays, as before. No discernible inhibitory effect on DMS production was noted in the presence of either compound. Note that creatinase (another M24B family enzyme, see above) from *Paracoccus*, also retains its activity in the presence of EDTA (Wang *et al.*, 2006). These findings imply that DddP does not require metal cofactors for its activity, although it cannot be wholly discounted that the metal of interest was not sequestered from the enzyme. However, EDTA is an efficient compound in sequestering transition metal ions (such as Co and Mn) from metallopeptidases, and so would be expected to have done so in this experiment (Auld, 1995).

In an attempt to identify putative cofactors (and, in particular, metal ions) more directly, inductively coupled plasma- optical emission spectroscopy (ICP-OES) was used to try to detect metal ions in a sample of DddP. Samples of active 10  $\mu$ M DddP in 2.5% (v/v) nitric acid, were analysed on a Varian Vista pro CCD simultaneous ICP-OES. None of the following metals, Co, Cu, Mn, Ni and Zn was detected above background levels, consistent with the observations when EDTA was added.

## 2.2.3.6 DddP active site residues

The M24 family enzymes (including members of both sub-families such as methionine aminopeptidase [M24A], and creatinase [M24B]), contain a "pita bread" fold that harbours their active sites (Bazan *et al.*, 1994). Five residues in particular have been predicted to form

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the active site and/or coordinate metal-binding in the M24 metallopeptidases, these being D97, D108,E204, E235 H171 (Schiffman *et al.*, 2006). Significantly, all these residues are conserved in DddP, towards its C-terminal end (Figure 2.7).To determine if these residues were important in DddP, site-directed mutagenesis was carried out. The pBIO1658 plasmid, created for the over expression studies (see above), had its cloned *R. nubinhibens* ISM *dddP* gene altered, by introducing 6 different residue substitutions in individual copies of the vector. The D295, D297, D307, H371, E406 and E421 (see Figure 2.7) residues were each individually substituted for an alanine, using a Quikchange XL site-directed mutagenesis kit. Mutagenised plasmids were transformed into *E. coli* BL21 and confirmed by sequencing. Each of these site-directed mutations completely abolished DMS production, as assayed by DMS production. Thus, DddP appears to require similar active site residues as other members of the M24B enzyme family. However, unlike other M24 enzymes, and like creatinase, these residues are important for catalytic activity in a non-metal-dependant manner.

## Figure 2.7 Predicted active site residues, selected for site-directed mutagenesis in DddP of *Roseovarius nubinhibens* ISM

	290	300	310		370	380			
	390								
Nterm-									
EIISFDTDLIGSYGICVDISRSWFQAQKYGCLMHGVGLCDEWPLVAYPDQAVPGSYDY									
400	41	LO 4	20	446					
PLEPGMVLCV <b>B</b> AAVGAVGGNFTIKL <b>B</b> DQV-Cterm									

# A section of the DddP amino acid sequence is shown. Residues highlighted in black, which are predicted to function at the active sites of M24B peptidases (Bazan *et al.*, 1994; Schiffman *et al.*, 2006), were changed to alanine by SDM (Table 8.6).

These findings further expand our understanding, in a biochemical sense, of this relatively recent addition to the DMSP lyase family. However, I wished to also investigate the evolution of this enzyme, particularly in regards to its transferral between the kingdoms, such as the fungi, and the ecological relevance of such events.

## 2.2.4 The presence of a functional *dddP* gene in the opportunistic coral pathogen, *Aspergillus sydowii*

Earlier work by Yoch and colleagues (Bacic *et al.*, 1998; Bacic and Yoch, 1998) had shown that some Ascomycete fungi, notably those that were isolated from decaying remains of the DMSP-producing salt-marsh grass *Spartina*, could make DMS from DMSP. Several Ascomycetes, including *Aspergillus* and *Fusarium spp*. were then shown to contain functional *dddP* genes (Todd *et al.*, 2009).

The presence of these organisms in DMSP-rich environments prompted me to investigate fungi that are associated with another DMSP-rich niche, namely coral reefs. Corals include single-celled, symbiotic photosynthetic dinoflagellates, such as *Symbiodinium*, which have massive levels (as great as 0.5 M) of intracellular DMSP (Matrai *et al.*, 1994; Raina *et al.*, 2010).

## 2.2.4.1 DMS assays of isolates of Aspergillus sydowii confirms a Ddd<sup>+</sup> phenotype

I obtained 20 isolates of *Aspergillus sydowii* from Krystal Riepen (then at Cornell University), sourced from a variety of environments. This species is a putative fungal pathogen of the sea fan *Gorgonia ventalina* (Hernández *et al.*, 2008) and so would be a good candidate for a marine *Aspergillus* species that catabolises DMSP. To test this, mycelial plugs (taken from the growing edge of a mycelium incubated for 48 hours at 28°C on Potato Dextrose Agar) of each strain were placed in liquid Vogel's medium containing 5 mM DMSP, and incubated at 28°C overnight in screw cap vials. After 6 hours, DMS was quantified by GC (Table 2.1). All of the strains exhibited a Ddd<sup>+</sup> phenotype, at varying levels of activity. There was no clear correlation between site of isolation and level of DMS release.

# Table 2.1 Dimethylsulfoniopropionate-dependent dimethyl sulfideproduction in strains of Aspergillus sydowii

Strain (1)	Source (2)	DMS (3)
SOMB	Infected Gorgonia ventalina	0.83±0.09
SABA	Infected G. ventalina	2.01±0.48
DumpD	Infected G. ventalina	0.41±0.03
FK11	Infected G. ventalina	0.5±0.02
15B1	Infected G. ventalina	2.89±0.31
NRRL 242	Environmental	2.17±0.16
NRRL 663	Environmental	4.59±0.20
NRRL 251	Environmental	0.54±0.06
NRRL 247	Environmental	1.67±0.03
KIR 382A	Environmental	2.31±0.28
SRRC 2540	Environmental	2.78±0.44
NRRL 4790	Environmental	0.65±0.28
NRRL 245	Environmental	0.31±0.02
NRRL 249	Environmental	0.26±0.11
NRRL 1732	Environmental	4.31±0.04
NRRL 5913	Environmental	3.22±0.53
NRRL 244	Environmental	0.70±0.02
NRRL 253	Infectious—human	$0.11 \pm 0.02$
297072	Infectious—human	1.20±0.04
SRRC 1112	Unknown	1.10±0.03

Column (1) lists the *A. sydowii* strains used in Rypien *et al.* (2008). Column (2) shows their sources, where known. Column (3) shows levels of DMSP-dependent DMS production, in nmol DMS  $h^{-1}mg^{-1}A$ . *sydowii* mycelial dry weight, with standard errors from two samples. Adapted from Kirkwood *et al.* (2011a).

## 2.2.4.2 PCR amplification of conserved *dddP* from A. sydowii

To examine if these *A. sydowii* contained a *dddP* gene, the SydDddPF and SydDddPR primers (both of which had some redundancy) were designed, based on sequence conservation of the *dddP* genes of *A. oryzae, F. culmorum* and *F. graminearum.* These primers targeted an internal fragment of *dddP*, 238 bp downstream 3' of the ATG start codon and 114 bp upstream of the stop codon.

In all 18 cases, PCR amplification of the genomic DNAs of using *PfuUltra* DNA polymerase (Stratagene) generated fragments of the expected size, *ca.* 1.2 kb. DNA sequencing of these PCR products showed them to be very similar to each other (>97 % identity at the DNA level), ~85% identical to the *dddP* gene of the Ddd<sup>+</sup> *A. oryza*e and ~80% identical to the *dddP* genes of the *Fusarium spp.* (Table 2.2; Todd *et al.*, 2009). Thus, *dddP* was likely present in the last common ancestor of *A. sydowii*. Variations in the level of homology between the *A. sydowii dddP*s did not seem to correlate with the source of each isolate (environmental, human, sea fan), supporting the suggestion (Rypien *et al.*, 2008) that the *A. sydowii* populations are panmictic.

	F. oxy	A. oryzae	NRRL 263	NRRL 4790	Somb	Dump D	KIR 382A	NRRL 52277	Group A strains	NRRL 663	Group B strains	NRRL 5913	Group C strains
F. graminearum	194	195	210	212	210	212	214	216	214	216	215	216	216
F. oxysporum		160	199	200	201	204	200	201	203	200	199	200	200
A. oryzae			149	148	151	154	153	153	155	153	152	153	153
NRRL 263				26	19	24	28	19	26	23	26	25	27
NRRL 4790		,			15	18	20	24	22	16	15	16	16
Somb						3	17	17	15	12	13	14	14
Dump D							20	20	18	15	16	17	17
KIR 382A								8	4	13	10	11	11
NRRL 52277									6	16	16	17	17
Group A strains										13	14	15	15
NRRL 663											3	4	4
Group B strains									0			1	1
NRRL 5913													2

Table 2.2 Conservation of *dddP* regions present in the Ddd<sup>+</sup> fungi

Numbers of nucleotide differences (out of 998, ~88% of a conserved region of *dddP*) in *A. sydowii dddP* and corresponding regions of *dddP* in *A. oryzae, Fusarium graminearum* and *F. culmorum* (Todd *et al.*, 2009) are shown, following comparisons with Megalign. Group A, group B and group C strains each have identical sets of sequences: group A; NRRL 245, NRRL 249; group B; 297072, SRRC 2540, FK11, 15B1, NRRL 251, NRRL 1732; group C; NRRL 242, NRRL 247. The analyses were carried out on 998 bps of unambiguous sequences, which are deposited at GenBank as follows: strain and accession number, respectively; 297072, GQ421799; DumpD, GQ421800; FK11, GQ421801; 15B1, GQ421802; KIR 382A, GQ421803; NRRL242, GQ421804; NRRL245, GQ421805; NRRL247, GQ421806; NRRL249, GQ421807; NRRL251, GQ421808; NRRL263, GQ421809; NRRL663, GQ421810; NRRL1732, GQ421811; NRRL4790, GQ421812; NRRL5913, GQ421813; NRRL52277, GQ421814; SRRC2540, GQ421815; Somb, GQ421816.

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## 2.3 Conclusions

## 2.3.1 DddP is an atypical M24B metallopeptidase

The ability of DddP to cleave a non-peptide bond (C-S) is unusual, but not unique, among the M24B family of peptidases. For example, creatinase of *Pseudomonas putida* generates sarcosine plus urea from water plus creatine, by cleaving a non-peptide C-N bond (Bazan *et al.*, 1994). Indeed, this implicates DddP as the least typical of the peptidases encountered in this group. Many well-characterized M24 enzymes, such as the cobalt-dependent methionine aminopeptidase from *E. coli* have metal cofactors. Yet DddP does not, and so in this regard too it resembles the creatinase of *Paracoccus*, which also lacks a metal-containing active site (Wang *et al.*, 2006). It may be significant that creatinase and DddP, the two known non-metal-dependent M24B enzymes, have structurally related substrates that carry a positive charge at or close to the catalytically cleaved bond. The work described here further extends the types of enzymatic activities that can be accomplished by M24B family members and, importantly, establishes that at least one of them can cleave a C–S bond. The novel structural features of this enzyme compared to other known lyases, such as the cupin-containing DddL, emphasised the diversity of DMSP-cleaving enzymes, and further

encouraged new searches for yet more DMSP lyases that had not yet been discovered (see chapter 4).

## 2.3.2 Evolution of fungal *dddP*

Given the distribution and conservation between the fungal species studied, it can be concluded that *A. sydowii* likely acquired *dddP* through a HGT event, as had been proposed for the other *Aspergillus* spp., and that the distribution of the gene matches the panmictic nature of the *A. sydowii* population (Todd *et al.*, 2009; Rypien *et al.*, 2008). It is debatable if *A. sydowii* and other DddP-containing fungi are mutualistic / commensal symbionts, or opportunistic pathogens (Smith *et al.*, 1996; Geiser *et al.*, 1998; Alker *et al.*, 2001; Hernández *et al.*, 2008). Both marine and terrestrial species of fungi might benefit by using the DMSP synthesised by corals or angiosperms, respectively, and it may have been a co-habitation by these organisms that resulted in the transfer of DddP between species.

# Chapter 3

# Optimisation of *Ruegeria pomeroyi* DSS-3 growth and microarray parameters

## **3.1 Introduction**

One major goal of this study was to discover novel genes in the Roseobacter clade bacterium, *Ruegeria pomeroyi* DSS-3, that respond to a number of inducers, specifically DMSP and the products of its cleavage, acrylate and DMS. Microarrays provide powerful tools to achieve this goal, but a degree of preliminary work was required to try to optimise the procedure. These included the development of suitable growth conditions for *R. pomeroyi* DSS-3, and methods for the isolation of consistent and satisfactory levels of high quality cellular material and RNA.

For this to be done, it was first necessary to establish media and growth conditions that would

- (a) maximise any differences in the expression of those genes whose transcription was affected by growth of the cells in one of the potential co-inducers
- (b) not be distorted by any severe inhibitory effects of these molecules

## 3.1.1 Existing transcriptomics studies involving DMSP

In previous work in the UEA lab's studies on the effects of DMSP- and/or acrylate-mediated induction of known genes involved in the catabolism of these substrates, concentrations of 5 mM and 2.5 mM respectively had been used, in media in which a "conventional" carbon (C) source, usually succinate, was also present. These concentrations may initially seem high. However, acrylate at concentrations of 1.3-6.5 mM has been recorded in the mucus of algal *Phaeocystis* colonies (Noordkamp *et al.*, 2000). It was therefore felt that this would be more informative than the very low levels of DMSP (80  $\mu$ M) used by Bürgmann *et al.* (2007) in their transcriptomic study of the same strain. From our lab's experience it was deduced that such a low concentration would be rapidly consumed in batch cultures prior to the harvesting of the mRNA. Given the very short half-life of mRNA in bacteria, this would likely account for the low-to-absent levels of induction that were seen by Bürgmann *et al.*, even with *ddd* 

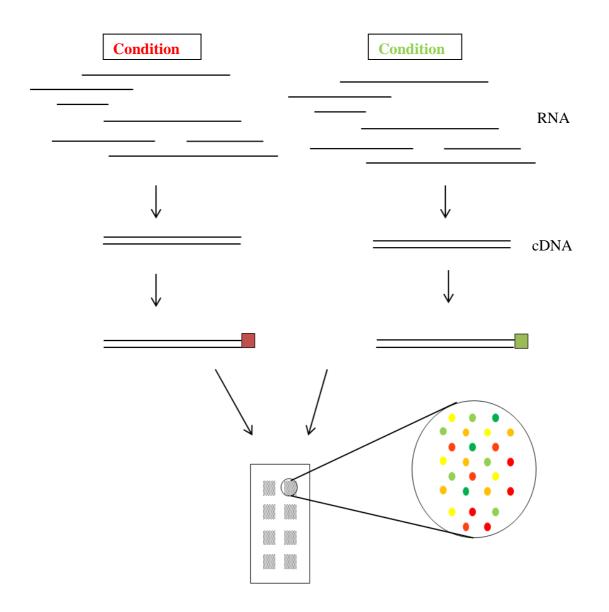
and *dmd* genes that are known to be induced many-fold by one or other of these substrates (see below). Additionally, only the effects of DMSP on gene expression were examined, and not of the two other co-inducers used in this study. As will be shown in chapter 5, the toxic effect of acrylate on growth of wild type *R. pomeroyi* cultures is minimal until well above several mM concentrations, although the myriad of indirect effects of adding a toxic compound may be noted in aspects of the gene expression.

## 3.1.2 An introduction to microarrays

DNA microarrays provide a remarkably powerful method of quantifying the expression of a large number of genes (>10,000), sometimes the entire genome, of an organism in response to an external stimulus and/or genetic background.

The mRNA molecules obtained from the organism grown under two (or more) different conditions are reverse transcribed to cDNA, then labelled, *en masse*, with different markers – usually coloured dyes (see below). This is known as a Type I microarray experiment, and the alternative (Type II) involves the hybridisation of one of the dyes to a reference sample, such as genomic DNA, in all of the arrays (DeRisi *et al.*, 1997).

To quantify these labelled RNAs, a microarray first requires the attachment of thousands of pre-designed DNA oligonucleotide "probes" to a glass slide. The probes comprise short sequences (~60bp), or oligomers, of DNA that contain sections of the target genes. Thus, upon addition of the solution consisting of reverse-transcribed cDNA from the RNA of cells exposed to an experimental condition, these probes become bound. This level of binding infers the level of mRNA that was originally present in the sample for a given probe. The samples, hybridized to fluorophores, give a measure of the signal intensity, or initial RNA concentration, by fluorescence emission, which is detected using a scanner, such as a GeneScan (summarised in Figure 3.1).





Simplified microarray procedure. RNA is isolated from cells grown in two experimental conditions. RNA is reverse-transcribed to cDNA, which is labelled using a coloured fluorophore, such as Cy3 or Cy5. Labelled cDNA is applied to DNA microchip and hybridised. Microchip is scanned, recording the ratio between the two intensities of fluorescence emissions. Red or green spots represent the abundance of one fluorophore, indicating a greater level of cDNA from one sample compared to the other. This illustrates the initial levels of RNA in the sample, and by association, the cells. Spots that appear yellow exhibit roughly equal levels of both fluorophores.

There are two general variants of the microarray procedure. A microarray study may use one, or two fluorophores to label cDNA, and these are named either single-, or dual-channel microarrays, respectively. Thus, a single channel array will provide a measurement of the relative abundance of genes in a given sample, compared to all of the other genes. In a dual-channel array, each fluorophore has a different fluorescence emission spectrum and is hybridised to a different sample, both of which are applied to the slide simultaneously (Shalon *et al.*, 1996). For example, dyes Cy3 and Cy5 have emission wavelengths of 570nm (appears green) and 670 nm (appears red), respectively. A dual-channel array uses the ratio of the intensity of each dye to assess the relative up- or down-regulation of each gene (Tang *et al.*, 2007).

Microarrays are a useful tool for analysing gene expression at a pan-genomic level, and therefore I wished to first optimise growth of *R. pomeroyi* cultures and isolation of good quality RNA for just such an experiment.

## **3.2 Results**

## 3.2.1 Growth characteristics of Ruegeria pomeroyi DSS-3

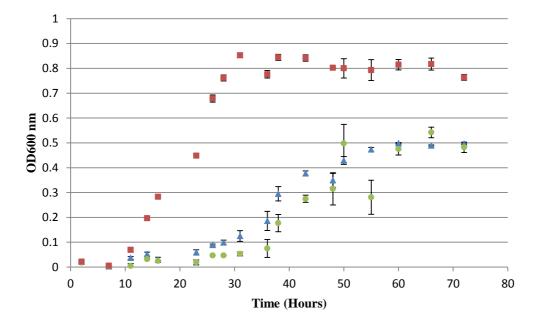
*R. pomeroyi* DSS-3 was originally isolated from seawater samples that had been plated onto seawater medium enriched with 10  $\mu$ M DMSP, although this was not the sole C source (González *et al.*, 1999; 2003). This strain was shown to grow well on complete, ½ strength YTSS (which contains tryptone, in reconstituted seawater) with colonies forming within 2 days at 28°C (González *et al.*, 1996). It also grows readily in a marine basal medium (MBM; Baumann and Baumann, 1981); a defined, minimal growth medium that is amended with a sea salts mix plus vitamins. There is no specific requirement for the vitamins, but enhanced growth has been observed with their addition; biotin, folic acid, pyridoxine, riboflavin, thiamine nicotinic acid, pantothenic acid, cyanocobalamin and *p*-aminobenzoic acid (all of which are B-vitamins) have all been shown to improve growth, (González *et al.*, 1997).

To measure the growth of *R. pomeroyi* DSS-3 on the substrates DMSP and acrylate as sole C sources, batch culture experiments were carried out to produce growth curves. Cells were grown overnight in  $\frac{1}{2}$  YTSS media, then diluted 1:100 into minimal MBM medium amended with 10 mM succinate, or with 5 mM DMSP or with 2.5 mM acrylate as sole C sources. Growth of the cells was determined by spectrophotometric measurement of their optical densities at 600 nm (OD<sub>600</sub>).

As shown in Figure 3.2, all three compounds acted as sole C sources, though, clearly, succinate was most effective, with a shorter lag phase (*ca.* 10 hours compared to *ca.* 20 h for acrylate and DMSP) and a higher final cell density after 60 hours incubation ( $OD_{600}$  of 0.8 for succinate and 0.5 for acrylate or DMSP). All three conditions would supply reasonable amounts of biomass for subsequent studies.

Strikingly, these observations are in contrast to those of González *et al.* (1999), who reported that this strain did not grow on acrylate and only grew on DMSP following iterative inoculations onto a medium containing these compounds. The reasons for this disparity are unclear, but perhaps stem from their higher concentration of acrylate (5 mM compared to 2.5 mM). Unfortunately, we were unable to acquire the strain of *R. pomeroyi* as isolated by González *et al.*, so the possibility that an inadvertent genetic change had occurred in the different versions of this strain cannot be discounted. Oddly, this finding was redacted in 2003 (again by González *et al.*) when it was shown that DSS-3 was indeed found to grow on acrylic acid as a sole C source following iterative inoculations in MBM medium.

# Figure 3.2 Growth of *Ruegeria pomeroyi* DSS-3 in MBM minimal medium with 10 mM succinate, 5 mM DMSP or 2.5 mM acrylate



as sole C sources

Batch cultures of *Ruegeria pomeroyi* strain DSS-3 were incubated at 28°C in MBM minimal media supplemented with vitamin solution and 10mM succinate (red squares), or 5mM DMSP (blue triangles), or 2.5mM acrylate (green circles) as sole C sources.

## 3.2.2 RNA techniques

## **3.2.2.1** Optimisation of *Ruegeria pomeroyi* DSS-3 growth for RNA extraction and microarray analysis

It was decided that the RNA to be used in the microarrays should be harvested from cells in their early exponential stage since, *a priori*, it was felt that any differences in the expression of genes involved in the uptake and catabolism of the different carbon sources would be

greater than in cells that were approaching stationary phase, which was the case in the microarray survey by Bürgmann *et al.*, 2007.

Concerning the timescale of the induction period, it was considered to be important to choose a time that provided sufficient opportunity for the relevant genes to be induced at maximal levels or for the mRNA to decay, for those genes that were repressed by (say) acrylate or DMSP. Cells were grown using succinate as a C source; this had been shown to facilitate good growth and acceptable levels of biomass production for further experiments.

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## 3.2.2.2 RNA extraction

Cultures of *R. pomeroyi* DSS-3 were incubated at 30°C in MBM minimal media supplemented with 10 mM succinate to an  $OD_{600}$  of ~0.4 (*ca.* 16 hours growth). The cultures were then incubated for a further 2 hours in the presence of either 5 mM DMSP, 2.5 mM acrylate, 5 mM DMS or no additional inducer. Following this, a tenth volume of phenol:ethanol (5:95) "stop solution" was added, which inhibits ribonuclease activity, and RNA was extracted. There was some difficulty in achieving adequate levels of RNA, requiring several iterations and the development of a modified SV Total RNA kit (Promega) protocol, using many of the original reagents. A detailed description is given in the Materials and Methods section, but an overview is presented here, as follows.

Essentially, the modified protocol involved increased lysis and additional homogenisation steps; preliminary attempts at RNA isolation without these steps were inconsistent and resulted in low final RNA levels ( $<100 \ \mu g \ ml^{-1}$ ). We first attempted to increase yield by using increased lysozyme, diluted in TE (Tris:EDTA) buffer, in order to more effectively break down the cells following pellet retrieval (100 mg/ml compared to 50 mg/ml). This improved yield but was still not sufficient for further experiments. Both freeze:thaw (using liquid nitrogen) and homogenisation (using glass beads and a ribolyser) steps were tested to improve cellular disruption and RNA retrieval. At first, each was used individually, but it was

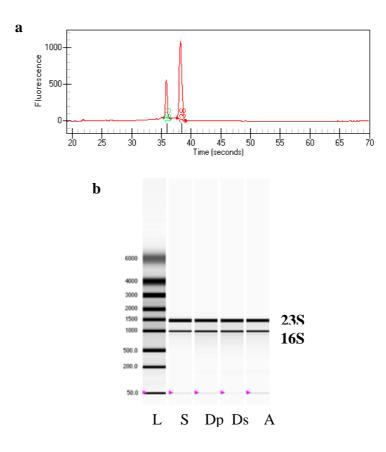
subsequently discovered that using both steps following lysozyme treatment improved RNA yield and quality significantly. Thus, 3 additional techniques were used to isolate RNA from *R. pomeroyi* at an acceptable quality and quantity. This yielded a final RNA concentration of *ca*. 0.8 - 1.5  $\mu$ g ml<sup>-1</sup> sample, which could be used for further procedures (microarray analysis - materials and methods).

## 3.2.2.3 Assessment of RNA quality

To measure the suitability of the RNA for further analyses, the quality and stability of the 16S rRNA in a 1  $\mu$ l sample, was determined. An Experion automated electrophoresis platform was used, the output being in the form of a "virtual" agarose gel. Samples underwent electrophoresis within a microfluidic chip, using a gel-stain solution. Micro-channels in the chip separated the RNA fragments based on their size and charge and these were detected via a bound fluorescent dye and photodiode (Experion Automated Electrophoresis System – user manual, BioRad). The data were converted into an electropherogram (plotting fluorescence vs. time) and then as a "virtual" agarose gel, shown in Figure 3.3a, and 3.3b, respectively.

The protocol used here yielded high quality RNA, as judged by the amounts and integrity of the rRNA molecules. A microarray analysis could then be undertaken.

# Figure 3.3 "Virtual" agarose gel of *Ruegeria pomeroyi* DSS-3 RNA samples



Samples of *Ruegeria pomeroyi* DSS-3 RNA were analysed using an Experion automated electrophoresis platform (BioRad), with 1 µl of RNA applied to each well. Data were initially converted into an electropherogram, plotting fluorescence vs. time (2a), where the peaks are identified (e.g. 16S or 23S rRNA) and integrated, to calculate a concentration of RNA. Clearly defined peaks illustrate a low level of rRNA degradation. Data can also be shown in the form of a "virtual" agarose gel (2b). L, RNA ladder, sizes shown in base pairs; S, succinate; Dp, DMSP; Ds, DMS; A, acrylate. Distinct bands corresponding to the 16S and 23S rRNA molecules are shown. The absence of smearing between the bands indicates lack of RNA degradation. Pink triangles indicate the position of the lowest ladder marker.

## 3.2.3.1 Slide design and labelling

The DNA microchip used in this study was initially designed by me *in silico* using the online tool, eArray (Agilent), by assigning short oligomeric sequences to a pre-determined grid, providing a complete, three-fold coverage of the entire *Ruegeria pomeroyi* DSS-3 genome, both the chromosome and megaplasmid (see materials and methods).

The samples of *R. pomeroyi* mRNA populations isolated above were fluorescently labelled, using a DIG-Easy Hyb kit (Roche). The RNAs from the unamended (succinate only) culture were labelled with Cy3 (red emission) and with Cy5 (green emission) for those samples from cultures containing acrylate, or DMSP or DMS. As noted in the materials and methods, a control array was conducted in which probes were hybridised to genomic DNA, in order to confirm the correct hybridisation to all of the spots on the array slide. A Type I experiment was chosen because I wished to more directly observe the changes in the RNA profile between the different growth conditions.

Dual-channel arrays are more useful in a study such as this, which requires the direct comparison of growth in certain conditions as a marker of gene regulation. A single channel study would require double the number of experiments to ascertain the same expression pattern, and would require additional normalisation, usually with the simultaneous assessment of pre-determined "housekeeping" genes (Stoyanova *et al.*, 2004).

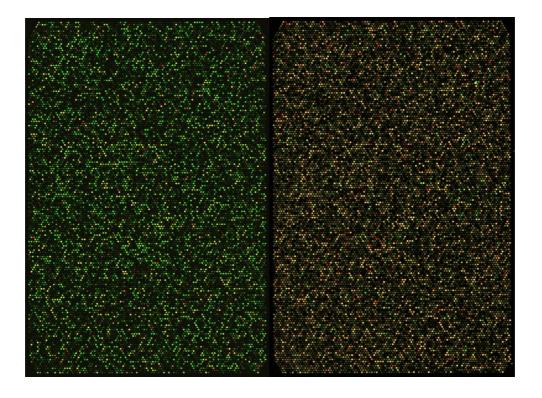
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### 3.2.3.2 Slide scanning

Once the microarray slide was hybridised and ready to scan, a number of efforts were made to improve the accuracy of any data retrieved. Firstly, samples were applied in biological duplicate, and each of the two arrays contained three distinct probes for each gene in the genome or megaplasmid, thus accounting for probe bias or incorrect binding. To reduce variation between the samples, the saturation levels and detector gain of the scanner for each fluorophore were manually calibrated using the software, GenePix (Agilent), following a preliminary scan of the slide. This process records the fluorescence of each dye, highlighting any bias and if the detector is allowing too much, or too little, light in, with the former causing saturation, represented by a white pixel on a scan image. Considering the method by which the program calculates fluorescence, by recording the intensity and wavelength of each individual pixel inside a given spot, or "feature", then white pixels would skew any subsequent results. The GenePix bioinformatics package uses a pre-made template file, which assigns a gene number to each of the features and allows identification of each spot following scanning. For this to be accurate, it is useful to carry out a manual examination of the template overlay and confirm that features are indeed bounded correctly.

Once a satisfactory alignment of template and slide was achieved, GenePix recorded the raw pixel values and fluorescence ratios for each spot on the array, and thus the expression of each gene in the *Ruegeria pomeroyi* DSS-3, in triplicate (shown in Figure 3.4). The raw data were normalised by the LOWESS (LOcally WEighted Scatterplot Smoothing algorithm) method and Marray, to calibrate inter- and intra-array variation such as dye intensity bias (Cleveland, 1979; Yang *et al.*, 2001; Dudoit and Yang, 2002; B Pearson, personal communication). LOWESS normalisation assumes that dye bias is dependent on spot intensity, and fits a smoothing curve to the dataset.

## Figure 3.4 Microarray output of RNA isolated from *Ruegeria* pomeroyi DSS-3 grown in +acrylate conditions

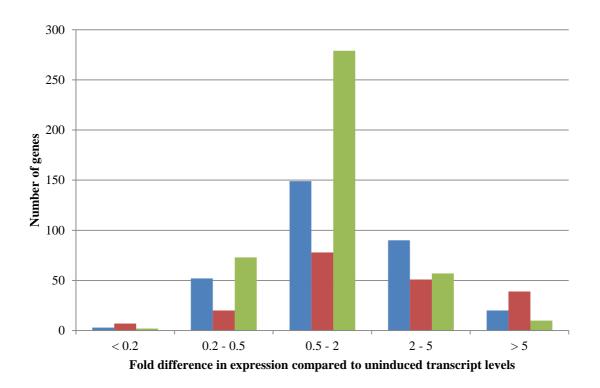


Raw data showing the output from a GeneScan scan of a microarray slide hybridised with RNA isolated from a culture of *Ruegeria pomeroyi* DSS-3 exposed to 2.5 mM acrylate. Each panel represents a separate microarray and thus, individual biological replicates. Each panel represents a three-fold coverage of the *R. pomeroyi* genome. Green spots indicate proportionally more RNA labelled from succinate conditions (Cy5), red spots indicate proportionally more RNA labelled from acrylate conditions (Cy3), and yellow spots indicate approximately equal levels of RNA from each sample. Blank spots indicate either a failed hybridisation, or more likely a control spot allocated by the template file, to allow for scanner calibration. This figure highlights the inter- and intra- array variation that may occur between samples, necessitating the need for post-array normalisation.

## 3.2.3.3 Summary of gene regulation in DMSP, DMS or acrylate growth conditions

In total, 31 genes were marked as having "no data" in any of the arrays following ratio calculation and normalisation, and 350 had at least one set of results absent. This data loss is most likely due to mis-binding of a number of the probes, or dyes, so that dye intensity for one spot was missing, causing the normalisation program to ignore the entire result (Yang *et al.*, 2001). The data retrieved were grouped according to the mean -fold change in expression (based on the ratio of the intensity of one dye over the other), either positively or negatively. An overview of this data is provided in Figure 3.5, or in its complete form in appendix A3. As expected, few genes were significantly (with both array results exhibiting P values of <0.05) up- or down-regulated in the presence of the compounds used.

# Figure 3.5 Summary of -fold differences in gene expression in *Ruegeria pomeroyi* DSS-3 exposed to DMSP, DMS or acrylate



Numbers of genes that are up- or down-regulated relative to the control (succinate-alone) cells. Blue bars, DMSP; red bars, acrylate; green bars, DMS. Data shown includes only genes with a mean fold change as indicated, where both repeats exhibited P values <0.05.

## 3.2.3.4 Limitations of a microarray study

Significant variability between replicates may exist due to the limitations of the microarray technology (Dudoit and Yang, 2002). This may be introduced through factors, such as:

probe-binding bias;

imbalance in the cDNA levels between samples;

variation in the binding of the fluorophore;

uneven spot detection;

over exposure.

To try to minimise these factors, the levels of RNA are accurately measured prior to reverse transcription, along with the quality of the RNA. Triplicate probes were designed for each gene in order to avoid a binding bias of one of the replicates, and a dual channel array was chosen to give a better representation of relative expression between samples. Finally, LOWESS and Marray normalisation were used to alleviate the variation caused by spot intensity or scanning errors, by merging the two colour data and applying a smoothing adjustment that attempts to remove incorrect variation.

The disadvantage of the dual-channel microarray is the possibility of one sample being of higher quality than the other, leading to better hybridisation and a bias of results. There is also the risk of gene-specific dye bias, in both forms of array. This is difficult to normalize, due to variation in the binding of different samples to the dyes, rather than their cognate oligonucleotides (Margaritis *et al.*, 2009). Whilst the absolute levels of gene expression cannot accurately be determined using a dual-channel microarray analysis, they give a clear quantification of the ratio of gene expression between two conditions, which was what was sought here.

Difficulties may also arise from the *in silico* analyses. If either of the green or red data is missing for a particular spot, no data will be recorded for that probe, and information may be lost as a result. The normalisation may also lead to the smoothing of otherwise legitimate variation, which may mask some gene expression unnecessarily. However, the possible loss in fidelity is more than made up for by the reduction in noise that this procedure achieves. In addition to this variability, the baseline expression of a gene of interest may mislead a researcher looking only for relative -fold change in expression. Therefore, a gene that is constitutively highly expressed and involved in (for example) a DMSP catabolic pathway, will change little (if at all) in response to the substrate.

A significant disadvantage to a microarray study is the skewed measurement of the expression of the genome in response to a given condition. Because a microarray records the activities of the cells 'transcriptome', that is, the level of mRNAs (indeed, all RNA molecules), this is not necessarily directly comparable to the proteome. The proteome describes the complete suite of proteins that a cell may express. The relative abundance of the various mRNAs is not always directly proportional to the presence of the translated polypeptide. This may be due to factors such as mRNA or protein stability in the cell, the processing of RNA or its translated protein product, or the mRNA's binding affinity for the ribosome, determined by the composition of the translation initiation sequence (Brown, 2002).

## **3.3 Conclusions**

Microarrays may also have some "biological" limitations, since, for example, they may detect genes whose expression is subject to general stress rather than direct catabolic responses to a particular molecule, and, of course, in the real world organisms are rarely faced with a straightforward "either-or" pair of dramatically different environments that differ in a single parameter. They nevertheless have the potential to give a global view of gene expression that was unthinkable not too long ago.

Certainly, some of the work described in this thesis was initiated solely from the microarray data. Notable discoveries include the gene encoding a wholly new DMSP lyase, and the expression of a suite of genes involved in a rather arcane catabolic pathway with no known link to sulfur, let alone DMS(P). It has to be admitted, though, that this particular set of experiments did not yield as much useful information as was hoped. This was due, largely, to the substantial errors in the different outputs for an individual gene, meaning that several genes whose levels of expression were genuinely affected by one or more of the agents used here were missed and, conversely, perhaps, some genes that were scored as being regulated by these agents were, in reality unaffected.

With these being the first microarrays attempted in our laboratory, it is likely that given increased financial resources, time, replicates and experience, a more accurate picture of the transcriptome of this bacterium would have emerged. However, this pan-genomic inspection of the transcriptional response of *Ruegeria pomeroyi* to DMSP and to related molecules did provide some very useful data.

# Chapter 4

## Discovery of the Novel Lyase, DddW

## **4.1 Introduction**

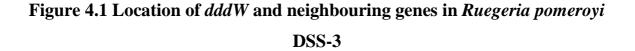
The previously described primary *ddd* genes (*dddD*, *dddL*, *dddP*, *dddQ* and *dddY*) had been identified by screening genomic libraries for any cosmids that conferred a Ddd<sup>+</sup> phenotype to other bacteria. In contrast, the work described in this chapter used a very different approach, in which a candidate *ddd* gene was identified, *de novo*, by its expression pattern, as revealed by the microarray analyses (see chapter 3).

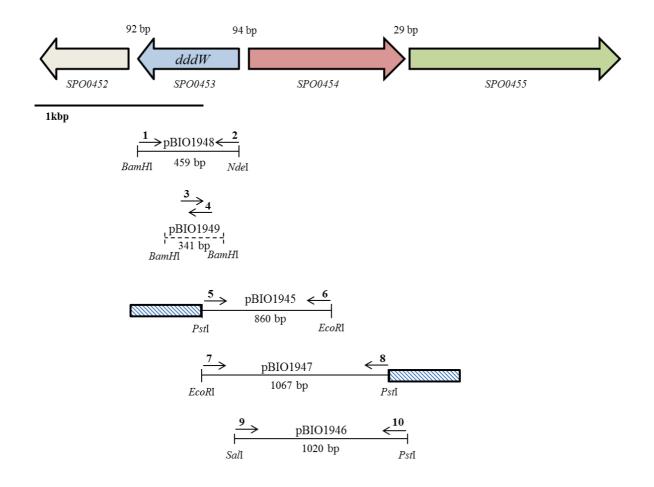
## 4.2 Results

## 4.2.1 Enhanced expression of the SPO0453 transcript in response to DMSP

The microarray analyses showed that the *Ruegeria pomeroyi* DSS-3 gene whose expression was enhanced greatly (average 41.1-fold), following growth of the cells in the presence of DMSP compared to the control medium, was *SPO0453*. Acrylate or DMS had no detectable effect on its expression compared to the control. This gene was of particular interest, not only because of its very pronounced, and specific, induction by DMSP, but also because of the nature of its gene product – the SPO0453 polypeptide is predicted to have a cupin domain, as do two other known DMSP lyases, DddL and DddQ.

The *SPO0453* gene is transcribed divergently from *SPO0454* and is upstream of *SPO0452*, whose 5' start is 92 bps downstream from *SPO0453* (Figure 4.1).To confirm the conclusions drawn from the microarray data, the effects of DMSP on the expression of *SPO0453* were examined, using two other, independent techniques.





Coloured arrows show locations of *SPO0453* and neighbouring genes. The *SPO0452* gene product is a predicted tellurite resistance protein, the *SPO0453* gene product is the novel DMSP lyase, DddW, the *SPO0454* gene product is a predicted LysR type transcriptional activator and *SPO0455* encodes a predicted lysyl tRNAsynthetase. Intergenic gaps are in base pairs (bp).

The 459 bp fragment (amplified using primers Wpet1 and Wpet2 – arrows 2 and 1, respectively) containing intact *dddW* was cloned into the pET21a expression vector to form pBIO1948.

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The 341 bp fragment (amplified using primers Wmut1 and Wmut2 - arrows 4 and 3, respectively) was cloned into pBIO1879 (a pK19*mob* derivative with spec<sup>R</sup>) to form pBIO1949. This was used to disrupt genomic *dddW* through homologous recombination.

The 860 bp fragment (amplified using primers Wprom1 and Wprom2 – arrows 6 and 5, respectively) was cloned into the *lacZ* reporter plasmid, pBIO1878, to create pBIO1945, a *dddW-lacZ* reporter fusion.

The 1067 bp fragment (amplified using primers 454prom1 and 454prom2 – arrows 7 and 8, respectively) was cloned into pBIO1878 to create pBIO1947, a *SPO0454-lacZ* reporter fusion.

The 1020 bp fragment (amplified using primers 454P1 and 454P2 – arrows 10 and 9, respectively) was cloned into the expression vector pOT2 to create pBIO1946, for heterologous expression of *SPO0454* with its native promoter.

Blue striped bars indicate junction with *lacZ* of pBIO1878.

## 4.2.1.1 Construction and assay of a dddW-lacZ fusion

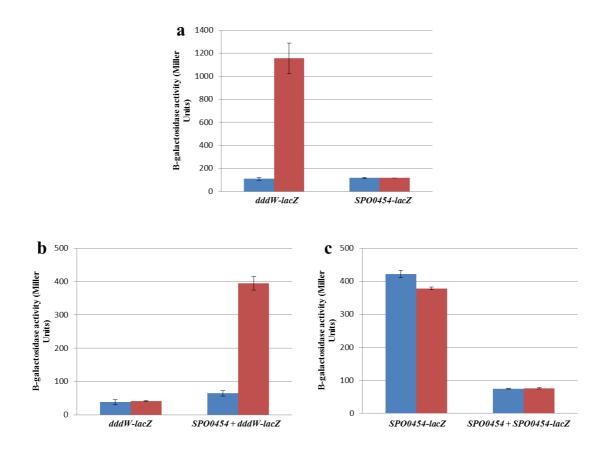
First, a *dddW-lacZ* transcriptional fusion was made by amplifying the intergenic region between *SPO0453* and *SPO0454*, using primers Wprom1 and Wprom2 to amplify the corresponding fragment from genomic DNA. This was cloned upstream of the promoter-less *lacZ* reporter in the wide host-range promoter-probe vector pBIO1878, cut with *EcoR*I and *Pst*I, and orientated such that the reporter *lacZ* would be under the control of the *dddW* promoter. The resulting plasmid, pBIO1945, was mobilized via a triparental conjugational mating into strain J470 (*Ruegeria pomeroyi* DSS-3 Rif<sup>R</sup> mutant), selecting for Rif<sup>R</sup>, Spec<sup>R</sup>, Tet<sup>R</sup> transconjugants. Two of these were then assayed for  $\beta$ -galactosidase activity after growth in minimal media that either contained or lacked 5 mM DMSP. Consistent with the microarray data, the *SPO0453-lacZ* fusion was induced 10-fold, in the DMSP-amended media, compared to the control (Figure 4.2a).

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The difference in the ratios between the microarray and transcriptional fusion data could be due to the method by which expression is quantified. Microarray analyses take a snapshot of mRNA levels in a cell at a single moment in time, whereas  $\beta$ -galactosidase assays record the accumulation of the expressed reporter enzyme over a sustained period. Also, as noted previously, microarray data represents a ratio of uninduced:induced expression, rather than absolute levels of expression.

## Figure 4.2 Regulation of the *Ruegeria pomeroyi dddW-lacZ* fusion in response to DMSP and the putative regulator, SPO0454



β-galactosidase activity of the *dddW-lacZ* (a, b), or *SPO0454-lacZ* (a, c) fusion plasmid when expressed by cultures of *Ruegeria pomeroyi* DSS-3 (a) or *Rhizobium leguminosarum* strain 3841 (b, c) in minimal media in the presence or absence of 5mM DMSP (blue and red bars, respectively). Error bars calculated from triplicate experiments.

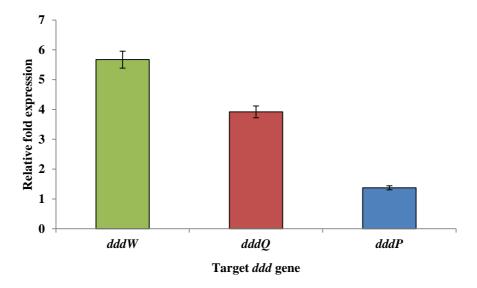
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### 4.2.1.2 qRT-PCR analysis of *dddW* transcription

DMSP-dependent induction of *SPO0453* (*dddW*) by growth in the presence of DMSP was also ratified by <u>quantitative real-time reverse transcriptase PCR</u> (qRT-PCR). To do this, *Ruegeria pomeroyi* DSS-3 was grown in the same manner as for the microarrays, overnight incubation followed by a 2 hour induction with/without 5 mM DMSP. The cultures were stopped for growth using phenol:ethanol and the cells frozen in liquid nitrogen. The RNA was harvested, purified and quantified using a Nanodrop 3000 (Thermo), and also checked for DNA or phenol contamination (based on the absorbance ratio at 260 nm: 280 nm and 260 nm: 230 nm, respectively). The RNA was then reverse-transcribed to cDNA and PCR-amplified using primers RTdddWF/RTdddWR, RTdddQF/RTdddQR and RTdddPF/RTdddPR, specific to *dddW, dddQ* and *dddP*, respectively (designed using Primer<sup>3</sup> software – Rozen and Skaletsky, 2000). Detection of amplification was achieved by using SYBR-green fluorescent label in conjunction with a C1000 Thermal Cycler and CFX Real-Time PCR detection system (BioRad).

The relative fold change in expression was calculated in relation to *SPO2904*, which encodes a serine/threonine protein phosphatase/nucleotidase. This gene is thought to be constitutively expressed in the conditions tested, so acts as a control gene for the results to be normalised (Bürgmann *et al.*, 2007). Despite *SPO2904* being slightly down-regulated in the array (-1.36 fold) in the presence of DMSP, there was a *ca*. 6-fold increase in *SPO0453* transcript levels over *SPO2904* in cells that had been grown in DMSP (Figure 4.3). Transcript levels were also measured for the genes encoding other lyases present in *R. pomeroyi*, namely *dddQ* and *dddP*. The qRT-PCR results showed that *SPO0453* is more responsive to DMSP than either *dddQ* or *dddP*. The microarray data and qRT-PCR do not corroborate each other as well as would be hoped, but nonetheless, an increase in transcription of *dddW* is observed in both experiments.

Figure 4.3 qRT-PCR showing the increase in transcription of the primary *ddd* genes in *Ruegeria pomeroyi* DSS-3, in response to DMSP



RNA extracted from *R. pomeroyi* cultures grown in minimal (MBM) media overnight, followed by a 2 hour induction by 5 mM DMSP, or no addition, was used to carry out qRT-PCR. Primers RTdddWF/RTdddWR, RTdddQF/RTdddQR and RTdddPF/RTdddPR were used to amplify *dddW* (*SPO0453*), *dddQ* (*SPO1596*) and *dddP* (*SPO2299*), respectively. Expression ratios are shown, using the reference gene, *SPO2904*, as a baseline control. Error bars calculated from duplicate readings.

### 4.2.2 Demonstration that SPO0453 encodes a novel DMSP lyase - DddW

Having confirmed that *SPO0453* was specifically induced by DMSP, the function of its product was examined directly, as follows.

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## 4.2.2.1 DddW confers a Ddd<sup>+</sup> phenotype when expressed in *E. coli*

Using primers Wpet1 and Wpet2, the intact *SPO0453* gene was PCR-amplified from *R*. *pomeroyi* DSS-3 genomic DNA (Figure 4.1), then ligated to the expression vector pET21a, cut with *Bam*HI and *Nde*I. The ligation mix was used to transform *E. coli* strain BL21 and the cells were plated onto media containing X-gal, IPTG and ampicillin, these three compounds being used, for blue-white screens to detect inserts (X-gal), induction of the *lacZ* promoter for expression of the cloned gene (IPTG) and to screen for the transformants (ampicillin). Plasmid DNA was isolated from six white transformant colonies and these were all confirmed to contain the correct insert, as judged by *Bam*HI / *Nde*I analytical restriction digests. One of these (with a plasmid termed pBIO1948) was ratified by sequencing.

A culture of an *E.coli* transformant containing pBIO1948 was then assayed for its Ddd phenotype. This was done by sonicating the cells and incubating the cell lysate for 1 hour with 5 mM DMSP. Protein levels were calculated using a Bradford assay. The DMS product, measured by gas chromatography (GC), was made at a rate of 35 pmol<sup>-1</sup>  $\mu$ g<sup>-1</sup> protein<sup>-1</sup> min<sup>-1</sup>. This activity is similar to that of *E. coli* harbouring plasmids that contained either the cloned *dddP* or the cloned *dddL* genes of *R. pomeroyi* DSS-3 and *Rhodobacter sphaeroides* 2.4.1., respectively (Todd *et al.*, 2012a; S Newton-Payne, personal communication).

## 4.2.2.2 DddW releases acrylate from DMSP, as confirmed by NMR

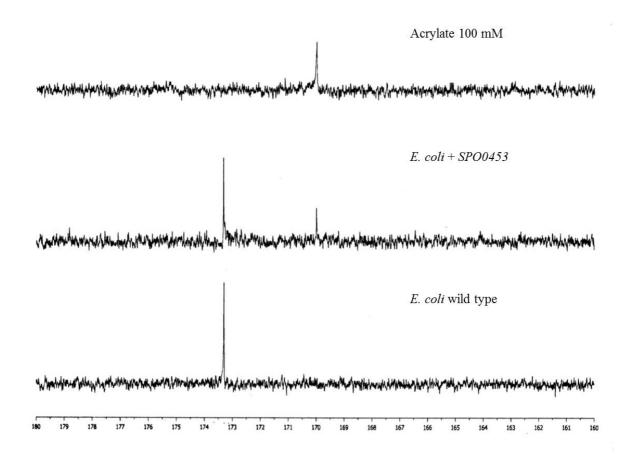
Thus, it seemed that the *SPO0453* gene encoded a DMSP lyase, which liberates DMS from DMSP. To identify the C3 cleavage product, cultures of the *E. coli* strain containing the cloned *SPO0453* were grown overnight at 37°C in LB then diluted  $10^{-2}$  into 1 ml M9 minimal medium (made up in deuterium oxide (>99.9%)], containing glycerol and 10 mM [1-<sup>13</sup>C]DMSP and 0.2 mM IPTG to induce expression. Following incubation at 28°C overnight, perchloric acid (5% v/v final concentration) was added to lyse the cells. The supernatant was added to NMR tubes and proton-decoupled <sup>13</sup>C-nuclear magnetic resonance (NMR) spectra were measured at 75 MHz with a Varian Gemini 2000 in D<sub>2</sub>O (as in Todd *et al.*, 2010). This

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was used to show that a proportion of the <sup>13</sup>C was converted to acrylate (Figure 4.4; J D Todd, personal communication). Thus, the *SPO0453* gene encodes a newly identified DMSP lyase, termed DddW, which cleaves DMSP into acrylate plus DMS.

## Figure 4.4 Nuclear Magnetic Resonance (NMR) spectra of *Escherichia coli* containing *dddW*, fed with [1-<sup>13</sup>C]DMSP



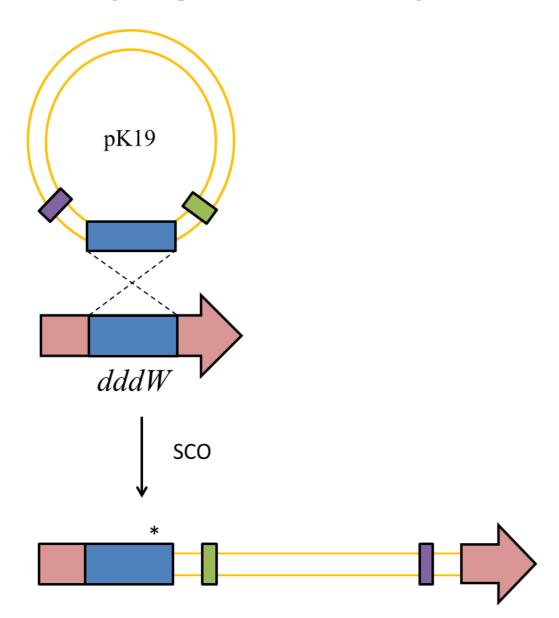
NMR spectra for pure sample of 100 mM acrylate is indicated, with a chemical shift peak at 69.9 p.p.m. The spectra below the acrylate reference, from top to bottom, shows *E. coli* expressing *SPO0453* from the pET21a plasmid, and *E. coli* wild type strain. The peak at 173.22 p.p.m. represents DMSP, added at 10 mM. Average scans recorded was 300 (J D Todd, personal communication).

#### 4.2.2.3 An insertion mutation into *dddW* affects DMS production

To examine the role of *dddW* in DMSP catabolism in *Ruegeria pomeroyi* itself, a 341 bp internal fragment of the gene was amplified using primers Wmut1 and Wmut2, and cloned into the suicide plasmid pBIO1879, cut with *BamH*I, creating plasmid pBIO1949 (Figure 4.1). The pBIO1879 plasmid is derived from pK19*mob* (Schäfer *et al.*, 1994), and contains an extra, selectable antibiotic resistance gene (spectinomycin, Spec<sup>R</sup>) to facilitate counterselection in the Roseobacters, several of which are resistant to tetracycline (Todd *et al.*, 2011). The original plasmid pK19 *mob* confers resistance to kanamycin and is mobilisable by conjugation at high frequency into a wide range of host bacteria, but fails to replicate in hosts other than enterics, such as *E. coli*. Therefore, if a fragment internal to a gene of interest is cloned into the polylinker of pK19*mob* and the resultant plasmid is mobilised into the corresponding host strain, then, by selecting for an antibiotic resistance marker on the vector, transconjugants should arise by a single crossover event in which the plasmid has integrated into, and hence disrupted, the chromosomal version of the gene (Figure 4.5).

The recombinant plasmid, pBIO1949, was mobilised into strain J470 (*Ruegeria pomeroyi* DSS-3 Rif<sup>R</sup>) via tri-parental mating (see materials and methods) and the transconjugants screened for those that were Rif<sup>R</sup>, Kan<sup>R</sup>and Spec<sup>R</sup>. These arose at *ca*.  $5.0 \ge 10^3$ cfu/ml and six colonies were used for genomic DNA preparation to determine if the plasmid had indeed inserted into the target gene, *dddW*. This was ratified by both PCR amplification and by Southern blotting of genomic DNA from the mutant strain, using the PCR product that was amplified to create pBIO1948 (J D Todd, personal communication).

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Blue bar indicates internal fragment of *dddW* cloned into the suicide vector pBIO1879 (pK19*mob* spec<sup>R</sup>), and the corresponding region in the genomic *dddW* of *Ruegeria pomeroyi* DSS-3 (red arrow). Green and purple bars indicate orientation of inserted plasmid DNA. SCO; <u>s</u>ingle <u>crosso</u>ver event via homologous recombination.\* It should be noted that any part of this region may be the starting point for the SCO event, and as such, may result in the insertion of pK19 at any point along this homology.

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The DddW<sup>-</sup> mutant strain was then assayed for DMSP-dependent DMS production, by growing in minimal medium containing 5 mM DMSP overnight, then assessing DMS levels using GC. It was found that the DddW<sup>-</sup> mutant released DMS at a rate ~50% lower than wild type cells. Thus, DddW makes a contribution to the overall Ddd<sup>+</sup> phenotype of *R. pomeroyi*, DSS-3, as do the other two lyases, DddQ and DddP, since mutations that abolish each of these lyases reduce, but do not abolish DMS production (Todd *et al.*, 2012a).

### 4.2.3 Regulation of *dddW*

As noted above, *dddW* is separated by 92 bps from the downstream gene, and is predicted to be in a single gene transcriptional unit (see Figure 4.1). The downstream *SPO0452* gene was up-regulated by 3.3-fold in the presence of DMSP, but its loose homology to a tellurite resistance protein does not appear to relate to DMSP in any significant manner. Thus, the function of this gene product is unknown.

The divergently transcribed *SPO0454* gene encodes a member of the widespread family of LysR-type transcriptional regulators (LTTRs). These proteins are responsible for regulating a diverse range of phenotypes, such as virulence, motility, quorum sensing and metabolism (Maddocks and Oyston, 2008). Another feature of many LTTRs is that they are auto-regulatory, repressing their own expression, even in the absence of the cognate co-inducer molecule. Given its location, relative to *dddW*, an attempt was made to show if the SPO0454 gene product was responsible for the DMSP-dependent regulation of *dddW*.

### 4.2.3.1 Difficulty in constructing a SPO0454<sup>-</sup> mutant *R. pomeroyi* strain

To do this, attempts were made to create a pK19 insertion mutant of *SPO0454* in the same way as described for *dddW*. Although transconjugants that were  $Rif^R$ ,  $Kan^R$ ,  $Spec^R$  were obtained, when these were checked by PCR amplification, all of them had an intact *SPO0454* 

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gene. A possible explanation for this is that the gene immediately downstream (separated by 29 bps) of *SPO0454* is predicted to encode a lysyl-tRNA synthetase, LysS, (Figure 4.1). Aminoacyl-tRNA synthetases are essential for all living cells, because they attach ("charge") amino acids to their cognate RNA molecules, prior to their incorporation by the ribosome into peptide chains during translation. Therefore, the failure to obtain an insertion into *SPO0454* may have been due to polar effects of such a mutation on the transcription of this downstream gene (Todd *et al.*, 2012a).

### 4.2.3.2 Effect of cloned SPO0454 on expression of dddW-lacZ in R. leguminosarum

I therefore adopted a different approach to test the regulatory role (if any) of the *SPO0454* gene. To do this, a "surrogate" host bacterium was employed to examine the expression of *dddW* in the presence and absence of *SPO0454*. The chosen bacterium was *Rhizobium leguminosarum* strain 3841, which was chosen for reasons already described.

First, the intact *SPO0454* gene, plus its native promoter, was cloned as a 1020 bp fragment into the wide host-range plasmid vector, pOT2, using primers 454P1 and 454P2 (see Figure 4.1), using *R. pomeroyi* genomic DNA as the template. This formed the recombinant plasmid, pBIO1946, which was then ratified by *SalI/ PstI* analytical restriction digests and by DNA sequencing of the insert. The *dddW-lacZ* fusion plasmid, pBIO1945, was transferred from *E. coli* into *R. leguminosarum* 3841 by triparental mating and then the *SPO0454*-containing plasmid, pBIO1946, was also introduced into the *R .leguminosarum* 3841/pBIO1945 by conjugation. Note that the two cloning vectors are compatible with each other and determine different antibiotic resistances (Gent<sup>R</sup> for pOT2 and Tet<sup>R</sup>/ Spec<sup>R</sup> for pBIO1878).

The two *Rhizobium* strains, each with the *dddW-lacZ* fusion plasmid and which either contained, or lacked, the cloned *SPO0454* gene, were each grown in the presence or absence of 5 mM DMSP and the cells were assayed for  $\beta$ -galactosidase activity. In the absence of the cloned *SPO0454* gene, the *dddW-lacZ* fusion was not expressed at significant levels in

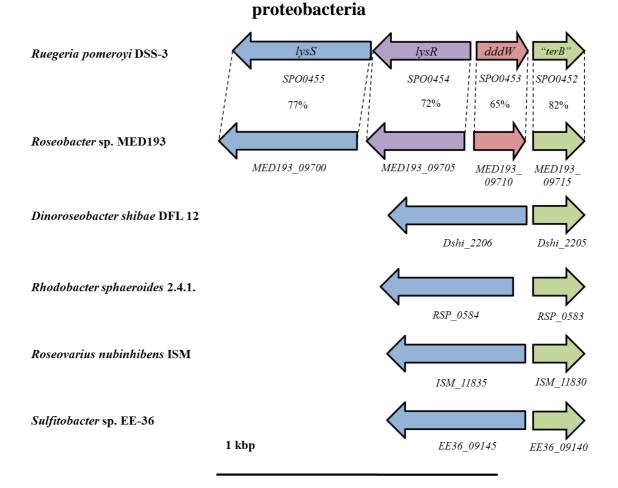
### Chapter 4: DddW

minimal media, either in the presence or absence of DMSP. However, in the *Rhizobium* strain containing both the fusion and the heterologously expressed SPO0454,  $\beta$ -galactosidase was expressed at high levels, but only when DMSP was present in the medium (Figure 4.2b). This provides strong evidence that the LysR- like *SPO0454* gene product is indeed a regulatory protein that activates transcription of *dddW* in the presence of DMSP.

As described previously, many LTTR proteins are auto-regulatory (Maddocks and Oyston, 2008). This was shown to be the case for SPO0454, as follows. A *SPO0454-lac* fusion was created by amplifying 1055 bp of the promoter region of *SPO0454*, using primers 454Prom1 and 454Prom2, and cloning the fragment into the *lacZ* reporter plasmid, pBIO1878, cut with *EcoR*I and *Pst*I, to make pBIO1947 (Figure 4.1). This was then mobilised into *R*. *leguminosarum* 3841. The *SPO0454-lacZ* fusion expressed  $\beta$ -galactosidase constitutively in *R. leguminosarum* itself. However, when pBIO1946 (pOT2 containing *SPO0454* and its native promoter) was present, the fusion was repressed ~4 fold, independent of the presence or absence of DMSP (Figure 4.2c).

### 4.2.4 Distribution and characteristics of DddW homologues

A BLASTp survey of the NCBI database (as at September, 2012) using DddW from *Ruegeria pomeroyi* DSS-3 as the *in silico* probe found one close homologue (65% identity at the amino acid level,  $E = 3e^{-63}$ ) to a polypeptide (MED193\_09710) in the  $\alpha$ -proteobacterium, *Roseobacter* sp.MED193. In this strain, *dddW* is also transcribed divergently from a gene that encodes a polypeptide that closely resembles that of *SPO0454* (72% identity at peptide level,  $E = 1e^{-148}$ ) (Figure 4.6, see later).



### Figure 4.6 Comparison of the dddW gene clusters in several $\alpha$ -

Approximate scale provided. Gene numbers are given below, and gene names for the *R*. *pomeroyi* homologues are given in the arrows. Matching colours indicate homologous gene products. Identity between the gene products of *R. pomeroyi* and *Roseobacter* sp. MED193 are given between dashed lines (%).

SPO0452, Putative tellurite related, TerB-like protein;

SPO0453, DMSP lyase, DddW;

SPO0454, LysR-like transcriptional activator;

SPO0455, Class I lysyl-tRNA synthetase.

### Chapter 4: DddW

There were no other convincing DddW homologues in any other known organisms whose genome sequences are available in current databases, so DddW is rare, compared to the other Ddd lyases that have been described.

Homologues of *R. pomeroyi* DddW were also sought in metagenomic data bases, including those in the Global Ocean Survey (GOS) (Rusch *et al.*, 2007). Only four convincing homologues (40–50% identical, probability  $<e^{-22}$ ) were found, a much lower number than DddQ and DddP, but about the same as DddL, and more than DddY. These homologues were all from the same location, a hypersaline lagoon site at Punta Cormorant in the Galapagos, the same site that contained the only DddL homologues in the GOS data set (Curson *et al.*, 2008; Todd *et al.*, 2012a).The significance, if any, of this observation remains to be determined and it also needs to be established if such divergent gene products have functional DMSP lyase activity.

Interestingly, when the *dddW* regions of the genomes of *R. pomeroyi* and *Roseobacter* sp. MED193 are compared, it appears that the *dddW-SPO0453 / MED193\_09710* genes have been inserted into a region of their genomes that is conserved in other Roseobacters, as follows.

A ~4kbp region containing genes *SPO0452- SPO0455* is well conserved in both *R. pomeroyi* and *Roseobacter* sp. MED193. Of the Roseobacters interrogated, other species possessed only divergently transcribed *SPO0455* and *SPO0452* genes, without the *SPO0453-SPO0454* genes (Figure 4.6). This is supportive of a horizontal gene transfer (HGT) event involving acquisition of the *dddW* gene pair. The distribution and function of the 2 ancillary and single co-acquired genes surrounding *dddW* in this cluster will now be discussed.

### 4.2.5 SPO0454 and SPO0455

There were a significant number of homologues of the divergently transcribed SPO0454 detected in the NCBI database, due to the ubiquity of LysR-like regulators (Maddocks and Oyston, 2008). Many of the closest matches were to *Burkholderia* spp. and *Pseudomonas* spp. ( $\beta$ - and  $\gamma$ - proteobacteria, respectively). SPO0454 also exhibits 72% identity at the peptide level with the *Roseobacter* sp. MED193 protein, MED193\_09705. The SPO0454 protein has been ratified as the regulator of *dddW* and itself in this study, and this fits well with the predicted LysR-like function of the gene product. The LysR-type SPO0454 is also very different from the LysR-type regulator, DddR in *Marinomonas* sp. MWYL1. This transcriptional activator of *dddD* is only 26% identical at the peptide level to SPO0454 (Todd *et al.*, 2007).

The gene directly downstream of *SPO0454* is thought to be co-transcribed, based on the polar effects of a mutation in *SPO0454* and a DOOR operon prediction (Mao *et al.*, 2008). The gene encodes a protein, SPO0455, which is 77% identical (E = 0.0) to the MED193\_09700 peptide, a predicted Class I lysyl-tRNA synthetase, LysS. This sequence elicited a large number of homologues, due to its essential nature as a tRNA synthetase. SPO0455 has a 24 amino acid N-terminal extension, compared to the other peptides, but a SignalP analysis failed to identify a leader sequence in the peptide (Petersen *et al.*, 2011). It is unknown why this extension is present in SPO0455 at this time.

The presence of two *lys* genes in close proximity perhaps reflects the acquisition of genes to clusters that complement their activity.

### 4.2.6 The cupin motif in DddW

Like DddL and DddQ, the DddW polypeptide contains a predicted cupin domain (see introduction). As shown in Figure 4.7, DddW shares several of the catalytically important residues with these two other lyases, notably, the two histidines and the glutamate within

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motif 1, and the histidine in motif 2, all of which are implicated in metal-binding. In preliminary studies (S Newton-Payne, personal communication), the DMSP lyase activity of semi-purified DddW protein was inhibited by addition of the metal-chelating agent ethylenediaminetetra-acetic acid (EDTA), providing circumstantial evidence that DddW is indeed a metallo-protein.

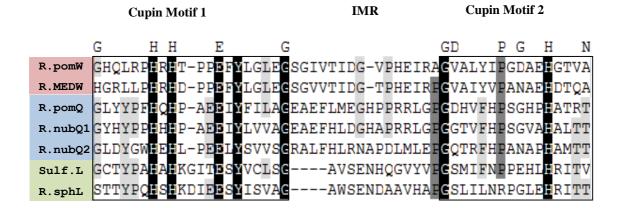


Figure 4.7 Alignment of the cupin domains of DddW, DddL and DddQ

Sequences of the predicted cupin motifs sequences of two DddW polypeptides are compared with representatives of the DddQ and DddL DMSP lyases. The amino acids that are highly conserved in all cupins are shown above the sequences (Dunwell *et al.*, 2004). *Ruegeria pomeroyi* DSS-3 (R.pomW; *SPO0453*) and *Roseobacter* sp. MED193 (R.MEDW; *MED193\_09710*;), shown in pink, are aligned with the DddQ lyases from *R. pomeroyi* (R.pomQ; *SPO1596*) and *Roseovarius nubinhibens* ISM (R.nubQ1, *ISM\_14090* and R.nubQ2 *ISM\_14085*), shown in blue, and the DddL lyases from *Sulfitobacter* sp. EE-36 (Sulf.L *EE36\_11918*) and *Rhodobacter sphaeroides* 2.4.1 (R.sphL*RSP\_1433.*), shown in green. Residues highlighted in black, dark grey or light grey indicate 100%, >80% or >60% identity between sequences, respectively. IMR; Inter-Motif Region. The work described illustrates the power of microarrays in the identification of genes of interest, in this case dddW, which encodes a novel DMSP lyase although one whose product has some similarity to those of the previously identified products of the dddL and dddQ genes. This new DMSP lyase has some features that are of interest for a number of reasons, as follows.

### 4.3.1 DddW is comparatively rare

The only convincing orthologue of DddW is in the closely related strain *Roseobacter* sp. MED193, which was isolated from the surface seawaters in the Northwest Mediterranean Sea (Roseobase; http://www.roseobase.org/). Few, if any, were in the GOS or any other metagenomic database. It is possible that it is more abundant in niches that have not been studied in detail – isolated or unsampled sediments for example - or it may be more frequent in clades of organism that are difficult to culture, so are under-represented in genome sequences. Or, DddW may just be rare.

### 4.3.2 A surfeit of enzymes that act on DMSP in Ruegeria pomeroyi DSS-3

The finding of a novel DMSP lyase in *Ruegeria pomeroyi* DSS-3 means that this bacterium possesses a demethylase, DmdA, an inactive DddD (see introduction) plus no less than three functional DMSP lyases. As such, *R. pomeroyi* DSS-3 has the most diversity of any single strain (to date) in its repertoire of DMSP catabolic enzymes. There are other examples of bacteria that harbour multiple Ddd<sup>+</sup> genes. – for example, *Roseovarius nubinhibens* ISM, contains *dddP*, *dddQ* and *dmdA* in its genome - but none has more than strain DSS-3 (see chapter 7). A recent, novel example of a non-Roseobacter with multiple DMSP-degrading

systems is the  $\gamma$ -proteobacterium, *Oceanimonas doudoroffii*, which contains functional *dddP* and *dddD* (Curson *et al.*, 2011b).

There are several putative reasons why *R. pomeroyi* (and other strains) may have multiple DMSP lyases:

- Each enzyme may be tailored to different cellular concentrations of DMSP, for example, by having a different K<sub>m</sub> for DMSP. This would allow a flexible, adaptive response to local environmental fluctuations. This could relate to events such as phytoplankton blooms, where large amounts of DMSP would be readily leaked into the milieu, and an enzyme with a more appropriate specificity would be required (Merzouk *et al.*, 2008);
- •
- The three enzymes also differ in size, having predicted molecular weights of 16, 22 and 50kDa for DddW, DddQ and DddP, respectively and may also require different cofactors, such as metal ions. Studies have not yet conclusively determined the metalbinding abilities of DddW and DddQ, with DddP having been found to bind no known metal ions. Therefore, in conditions where there is limited availability of the necessary metal cofactors, DddP may act as the primary DMSP lyase. Then, when conditions are favourable, DddW or DddQ become dominant;
- The "response times" of the enzymes may also differ. The microarray data recorded levels of RNA present in the cells at exactly 2 hours post induction. Therefore, each of the enzymes may be induced more, or less, quickly when exposed to DMSP, or are constitutively expressed. Again, this would be important for an efficient response to short-term events such as phytoplankton blooms.
- Lastly, the different DMSP lyases might even act on different substrates, with DMSP catabolism being a secondary function.

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Recent work in our laboratory has shown that far from being a model for the catabolism of DMSP, *R. pomeroyi* DSS-3 is something of an exception regarding its response to DMSP. In a survey of several different Roseobacters, it was the only one that grew well on DMSP and on acrylate as sole C sources (E Fowler, personal communication). It remains to be seen if this is connected with its unusual trio of DMSP lyases, a subject that is investigated more thoroughly in chapter 5.

### 4.3.3 Future work

- 1. Assaying the DMS production of a triple mutant strain of *R. pomeroyi* (DddW<sup>-</sup>/DddQ<sup>-</sup>/DddP<sup>-</sup>) would confirm that no other DMSP lyases exist in this bacterium.
- 2. The purification of DddW would allow ratification of the proposed cupin domain and identification of any bound metal.
- 3. The regulatory targets of the SPO0454 protein may extend beyond *dddW*, and it would be of interest to see if other genes are affected by this regulator in *R. pomeroyi*.

# Chapter 5

### The role of the acuI - dmdA operon in

### Ruegeria pomeroyi DSS-3

### **5.1 Introduction**

The previous chapter described the discovery of a novel DMSP lyase, DddW. This chapter will examine the *dmdA* gene and its enzymatic product, which is responsible for the initial step in the DMSP demethylation pathway. Just as with *dddW*, the initial observations on *dmdA* were obtained in the microarray analyses but were extended to yield new insights into the function and regulation of the *dmdA* gene. Additionally, an adjacent gene was found to provide novel insights into the connection between DMSP demethylation and cleavage in *R*. *pomeroyi*.

As shown in Figure 5.1, the *dmdA* gene (*SPO1913*) of *Ruegeria pomeroyi* DSS-3 is upstream of, and in the same orientation as, *SPO1914* from which it is separated by 77 bp. As mentioned previously, *SPO1914* encodes an enzyme that is a homologue (54% identical,  $E = 3e^{-112}$ ) of the so-called AcuI zinc dependent oxidoreductase from *Rhodobacter sphaeroides* 2.4.1, in the medium chain reductase superfamily, and which is involved in acrylate catabolism (Sullivan *et al.*, 2011). In addition, *acuI*-like genes are closely linked to several *ddd* genes that encode very different DMSP lyases and which occur in very different bacteria – *dddD* in *Halomonas, dddP* in *Oceanimonas* and *dddY* in *Alcaligenes*.

The finding of *acuI* next to a gene involved in the very different, demethylation, pathway was particularly provocative, especially since this was the case for many different Roseobacters (see below) and there are no *acuI*-like genes near the various *ddd* genes in *Ruegeria pomeroyi* or the other Roseobacters.

The work in this chapter therefore set out to uncover any links – both functional and regulatory – between the *acuI* and the adjacent *dmdA* genes in *R. pomeroyi*.

### **5.2 Results**

# 5.2.1 Induction of expression of the *dmdA-SPO1914* transcriptional unit in response to DMSP and acrylate

The microarray data (appendix A3) show that the expression of both *dmdA* and *acuI* have similar responses; both are induced by 5 mM DMSP (23.3-fold for *dmdA* and 16.4-fold for *acuI*). It was striking that the *dmdA* and *acuI* also responded to the DMSP catabolite acrylate (added at 2.5 mM), their levels of expression in the microarray being enhanced 23.8- and 17.3-fold compared to the no-addition controls.

The genes that are known to be involved in the downstream steps that convert MMPA to methane thiol (MeSH), labelled *SPO2045* (*dmdB*), *SPO3804* (*dmdC*) and *SPO3805* (*dmdD*) (see introduction) were also examined. The *dmdB* gene does not appear to be induced under any condition, but *dmdC* and *dmdD* were increased in expression in +DMSP (4.6-fold and 5.3-fold, respectively). Reisch *et al.* (2011b) observed that transcripts of all these genes were increased when cells were grown with MMPA or DMSP as a sole C source. Thus, *dmdB*, *dmdC* and *dmdD* transcripts were increased by 2-, 5- and 6-fold in MMPA, and 5-, 2- and 3-fold in DMSP, respectively.

The gene divergently transcribed from *dmdA*, *SPO1912*, was also up-regulated in the presence of DMSP and acrylate, being increased in expression by 4.6- and 6.1-fold, respectively (see below).

The responses of *dmdA* and *acuI* to DMSP resemble previous findings that detected a potential (but not significant) up-regulation of both genes in response to DMSP (Bürgmann *et al.*, 2007). The smaller response that was reported by those authors is likely due to their use of a lower concentration ( $80\mu$ M) of DMSP and/or the fact that they added DMSP to cells that were approaching stationary phase, rather than the exponentially growing cells used in the present study (see introduction; chapter 3). Bürgmann *et al.* (2007) did not investigate the effects of acrylate on the transcriptome of *R. pomeroyi*.

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The catabolite-mediated induction seen in the microarray is not unique. As described above, 3HP, a product of the DddD-mediated catabolism of DMSP in *Halomonas* HTNK1, induced transcription of the *ddd* genes *in situ* (Todd *et al.*, 2010). The regulation of *acuI* by acrylate also resembles the findings of Sullivan *et al.* (2011) in *Rhodobacter sphaeroides* 2.4.1. However, in the case of *Ruegeria pomeroyi dmdA-acuI*, acrylate is not an immediate, obvious catabolite of DmdA. The reasons for this unusual regulation are investigated below.

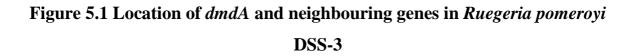
### 5.2.1.1 Constructing and assaying *dmdA-lacZ* and *acuI-lacZ* fusions

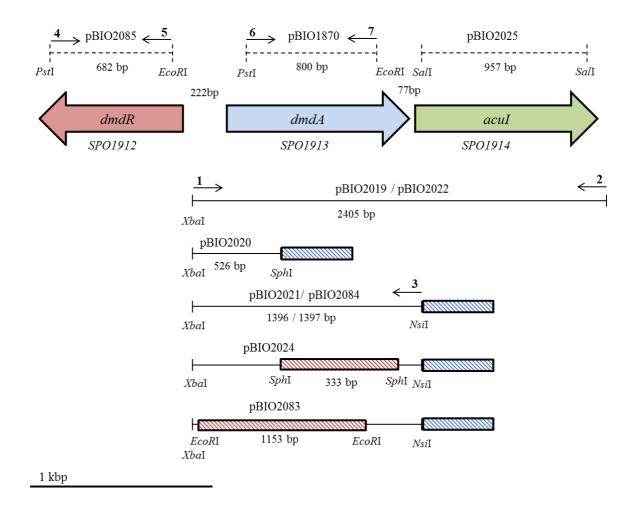
The microarray data and their close linkage provided good *prima facie* evidence that *dmdA* and *acuI* were co-regulated and both responded to the substrate DMSP and one of its catabolites, acrylate. To confirm this, a wholly independent measure of gene expression, namely *lacZ* reporter transcriptional fusions, was used.

In general terms, this was done by cloning fragments that spanned the *dmdA* and *acuI* promoter region into a wide host-range promoter probe plasmid, pBIO1878, and mobilising the resultant plasmids into *Ruegeria pomeroyi* DSS-3. The transconjugants were then grown in different conditions and the gene expression was measured by assaying  $\beta$ -galactosidase activities. The particular constructs were made as follows.

A 2405 bp fragment, which included the intact *dmdA* and *acuI* genes, plus 216 bps upstream of *dmdA* which spanned the predicted promoter region for these genes, was amplified from *Ruegeria pomeroyi* DSS-3 genomic DNA using primers SPO1913/14\_XbaF and SPO1913/14\_BamR (Figure 5.1). This fragment, cut with *XbaI* and *Bam*HI, was ligated into the expression vector, pBluescript SK-, digested with the same enzymes and the ligation mix was used to transform *E. coli* strain JM101. Transformants arose at *ca.* 5 x  $10^2$  cfu/ml and six colonies were selected for plasmid DNA preparations, followed by diagnostic restriction digests. One recombinant plasmid, termed pBIO2019, was confirmed as containing the correct fragment, cloned in the appropriate orientation, and was used as a starting point to create two individual *lacZ* reporter transcriptional fusions.

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Genes are shown to scale, illustrating gene numbers and intergenic regions.

A 2405 bp fragment of DNA was amplified using primers SPO1913/14\_XbaF (1) and SPO1913/14\_BamR (2) and cloned into pBluescript SK- to create pBIO2019, or pBIO1878 (*lacZ* reporter plasmid) to form pBIO2022. This PCR product was also used as a probe for Southern blotting of *dmdA*<sup>-</sup> and *acuI*<sup>=</sup> mutants.

A 526 bp fragment of DNA was released from the pBIO2019 plasmid by digestion with *Xba*I and *Sph*I and cloned into pBIO1878 to form the *dmdA-lacZ* fusion plasmid, pBIO2020.

A 1396 bp fragment of DNA was released from the pBIO2019 plasmid by digestion with *Xba*I and *Nsi*I and cloned into pBIO1878 to form the *acuI-lacZ* fusion plasmid, pBIO2021.

Primers SPO1913/14\_XbaF (1) and acuI+1\_NsiR (3) were used to amplify the same fragment as for pBIO2021, with the addition of a C nucleotide upstream of the *Nsi*I cut site, in order to create a frame shift mutation reporter plasmid, pBIO2084.

A 333 bp fragment of DNA was removed from pBIO2021 (red striped bar) by restriction digestion with *Sph*I to create the *acuI-lacZ* fusion plasmid, pBIO2024.

An 1153 bp fragment of DNA was removed from pBIO2021 (red striped bar) by restriction digestion with *Eco*RI to create the fusion plasmid, pBIO2083.

Primers 1912\_EcoF (5) and 1912\_PstR (4) were used to PCR amplify a 682 bp internal fragment of *dmdR* to clone into the suicide vector, pBIO1879 (a pK19*mob* derivative) to make pBIO2085. This was used to create a DmdR<sup>-</sup> strain of *R. pomeroyi*.

Primers SPO1913F\_Pst (6) and SPO1913R\_Eco (7) were used to PCR amplify an 800 bp internal fragment of *dmdA* to clone into pBIO1879 to make pBIO1870. This was used to create a DmdA<sup>-</sup> strain of *R. pomeroyi*.

The PCR product amplified using primers SPO1913/14\_XbaF (1) and SPO1913/14\_BamR (2) was digested with *Sal*I to release a 957 bp internal fragment of *acuI*, cloned into pBIO1879 to create pBIO2025. This was used to create an AcuI strain of *R. pomeroyi*.

Blue striped bars indicate the junction point with *lacZ* of pBIO1878.

above for pBIO2020.

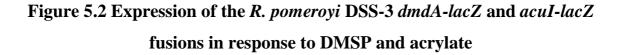
#### Chapter 5: *dmdA* and *acuI*

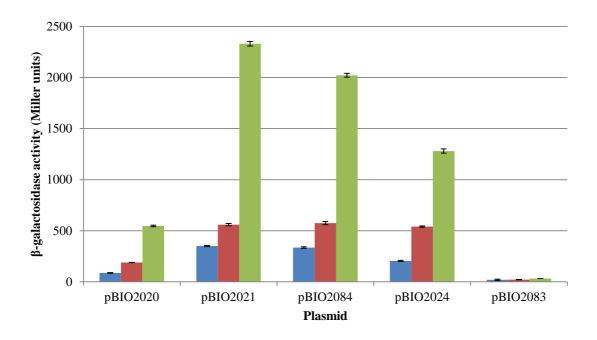
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A *dmdA-lacZ* transcriptional fusion was created by digesting pBIO2019 DNA with XbaI plus SphI to release a 526 bp fragment comprising 216 bp 5' upstream, and 310 bp downstream from the start of the *dmdA* gene and which therefore spans the predicted *dmdA-acuI* promoter region. This fragment was gel-extracted and ligated into the reporter plasmid pBIO1878 (see chapter 4), also digested with XbaI and SphI, before transforming E. coli strain 803, selecting Tet<sup>R</sup> (specified by pBIO1878) transformants. Restriction digests and then DNA sequencing of the plasmid DNA in six of the resulting transformants confirmed that they each contained the 526 bp fragment in the correct orientation and one of these plasmids, termed pBIO2020, was chosen for further study. A similar method was used to make the *acuI-lacZ* fusion plasmid. Thus, a 1396 bp fragment containing 216 bp upstream of *dmdA*, the entire *dmdA* gene plus 8 bp of the 5' end of acul was released from pBIO2019 by restriction digest with XbaI and NsiI, a naturally occurring restriction site within acuI. The NsiI site was used to sub-clone into pBIO1878, which possessed XbaI and PstI restriction sites, because the cutting of an NsiI restriction site leaves sticky ends complementary to PstI, an isoschizomer of NsiI. The recombinant plasmid was termed pBIO2021 and its integrity was confirmed as described

The two fusion plasmids pBIO2020 and pBIO2021 were each introduced into Ruegeria *pomeroyi* strain J470 in triparental conjugational crosses, selecting for Tet<sup>R</sup> transconjugants. For each cross, one transconjugant was purified and was then used to examine the expression of the corresponding fusion, as follows.

Cultures of R. pomeroyi J470/pBIO2020 and J470/pBIO2021 were grown in unsupplemented 1/2 YTSS medium and also in 1/2 YTSS that contained either 5 mM DMSP or 2.5 mM acrylate, before assaying the cultures for  $\beta$ -galactosidase activity. Note the use of the complete  $\frac{1}{2}$ YTSS medium, rather than the minimal MBM medium. This was because cell growth was considerably greater in the former and the levels of expression correspondingly greater than in the minimal medium. As shown in figure 5.2, the patterns of expression of the two fusions, as reflected in the resultant  $\beta$ -galactosidase activities were similar, but not identical to those seen in the microarrays.





Derivatives of *Ruegeria pomeroyi* DSS-3 (J470) containing one of the various fusion plasmids as indicated were grown in  $\frac{1}{2}$  YTSS medium (blue bars) or in the same medium supplemented with 5 mM DMSP (red bars) or 2.5 mM acrylate (green bars). Results of triplicate assays of  $\beta$ -galactosidase activities are shown as Miller Units, with standard errors.

Figure 5.1 shows the dimensions of the cloned DNA in the fusion plasmids, which are:

dmdA-lacZ (pBIO2020);

### acuI-lacZ (pBIO2021);

*acuI-lacZ* +1 (pBIO2084), a single base insertion, to create a frameshift in *lacZ* (see below); *acuI-lacZ* with a fragment of *dmdA* deleted but with an intact promoter region (pBIO2024);

acuI-lacZ with the majority of dmdA and its promoter region deleted (pBIO2083).

#### Chapter 5: *dmdA* and *acuI*

Thus, both fusions were induced by pre-growth in the presence of DMSP and of acrylate, compared to the values in the unsupplemented medium, and, for both fusions, acrylate was a more effective co-inducer than was DMSP. However, the ratios of increase were considerably less than those seen in the microarrays. Thus, DMSP caused a *ca*. 2-fold and 1.5-fold increase in  $\beta$ -galactosidase activity from, respectively, the *dmdA-lacZ* and the *acuI-lacZ* fusions compared to the control, whereas the corresponding factors of enhancement in the microarrays were, respectively *ca*. 20-fold and 15-fold. Similarly, the potency of acrylate-dependent induction was *ca*. 6-fold and 7-fold increase in  $\beta$ -galactosidase activity, and a *ca*. 20-fold and 15-fold in the microarray, for the *dmdA-lacZ* and *acuI-lacZ* fusions, respectively.

Whereas microarray data only present the *ratios* of the corresponding RNA molecules in the different samples, other methods, such as qRT-PCR and, as used here, the assaying of reporter fusion activities, indicate the *absolute* levels of expression of the genes under study. In the present study, the data presented in figure 5.2 show that (a) the *dmdA* and *acuI* genes are expressed at very high levels when induced by either acrylate or DMSP, since  $\beta$ -galactosidase activities > 1,000 Miller Units are, in our experience, towards the top end of gene activity as measured in this way and (b) perhaps most strikingly, the *acuI-lacZ* fusion was expressed at significantly greater levels than was the *dmdA-lacZ* fusion, under all three growth conditions (control, +DMSP or +acrylate).

### 5.2.2 Unusual expression of the *dmdA-lacZ* and *acuI-lacZ* transcriptional fusions

This expression pattern was surprising, since it had been predicted that *dmdA* and *acuI* are in a single transcriptional unit, and, while it is not unusual for downstream genes in an operon to be expressed at *lower* levels than the promoter-proximal genes, the converse is unusual. Three possible explanations for this observation were considered, and tested as follows.

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### 5.2.2.1 The *acuI* gene has another, internal promoter that responds to DMSP and to acrylate

To examine this possibility, a new *acul*–*lacZ* fusion plasmid, pBIO2083, was made (Figure 5.1). Compared to pBIO2021, the original plasmid that contains this fusion, pBIO2083 was deleted for the DNA upstream of *dmdA* and therefore lacks the promoter that is located in that region, as shown via the  $\beta$ -galactosidase assays. This new *acul-lacZ* fusion plasmid was made by digesting pBIO2021 with *Eco*RI and re-ligating to remove an 1153 bp fragment that contained the majority (975 bp) of *dmdA* and its 178 bp upstream promoter region. The ligation mix was transformed into *E. coli* 803, and plasmid DNA was isolated from 10 of the resulting transformants. These plasmids were then analysed by restriction digest with *Eco*RI to identify those that had lost the appropriately sized *Eco*RI fragment. One such plasmid, termed pBIO2083, was mobilised into *R. pomeroyi* J470 and a culture derived from a purified Tet<sup>R</sup> transconjugant was assayed for  $\beta$ -galactosidase activities, as described above. It was found that the "promoter-less" fusion was almost totally abolished in its activity, in all three conditions. Thus the promoter that drives the expression of *acuI* does indeed seem to be upstream of the *dmdA* gene.

### 5.2.2.2 The presence of an intact *dmdA* gene in the fusion plasmid may affect the expression of *acuI-lacZ*

The original *acuI-lacZ* fusion plasmid, pBIO2021, contains an intact version of *dmdA*, but the *dmdA-lacZ* fusion plasmid pBIO2020 does not. In light of the interactions between the DmdA-mediated DMSP demethylation pathway and the various Ddd-mediated cleavage pathways (see below), it may be that the presence of the extra copies of *dmdA* in pBIO2021 affect the expression of the *acuI-lacZ* fusion. To examine this, another *acuI-lacZ* fusion plasmid was made, which was very similar to pBIO2021, but which lacked an intact *dmdA* gene. Thus, the new fusion would be under the control of its promoter, but would not contain a functional *dmdA* gene. This plasmid was made by digesting pBIO2021 with *Sph*I and religating to remove a 333 bp fragment within *dmdA*, towards its 3' terminus (Figure 5.1). The

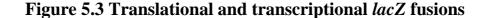
### Chapter 5: *dmdA* and *acuI*

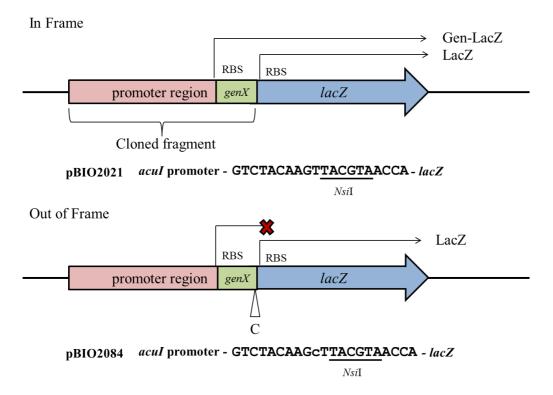
ligation mix was transformed into *E. coli* 803 and 10 colonies were used for plasmid DNA extraction. These plasmids were then analysed by restriction digest with *Xba*I and *Bam*HI to identify any that were appropriately truncated. One such plasmid, termed pBIO2024, was mobilised into *R. pomeroyi* J470 and a culture derived from a purified Tet<sup>R</sup> transconjugant was assayed for  $\beta$ -galactosidase activities, as described above. The new fusion was found to exhibit essentially the same pattern of induction as the *acuI-lacZ* fusion, with a *ca.* 2-fold and 5-fold increase in activity in the presence of DMSP or acrylate, respectively (Figure 5.2). Activity in acrylate was not as markedly high as in the *acuI-lacZ* fusion, but it is uncertain whether this is due to the reduced levels of DmdA in the cell, or if the removal of this region affected the stability or availability of the mRNA (see chapter 7). The latter seems unlikely, given the activity of a similarly truncated transcript, in the form of pBIO2083 (see above). In any event, this result showed that the intact *dmdA* gene is likely not the sole reason for the unexpectedly high  $\beta$ -galactosidase activity of the *acuI-lacZ* fusion.

# 5.2.2.3 The *lacZ* fusion plasmid, pBIO1878, contains its own, efficient, ribosomal binding site

The promoter-probe plasmid pBIO1878 was designed to construct transcriptional fusions (the reporter *lacZ* retains its own ribosomal binding site {RBS}). However, in some cases, translational fusions can be made if the junction between the cloned gene and *lacZ* is inframe. Therefore, in such cases, the level of  $\beta$ -galactosidase may be affected by the relative strength of the target gene's RBS, compared to that of *lacZ*. To examine this possibility, the junction sequences of the *acuI-lacZ* and the *dmdA-lacZ* fusions were deduced, first *in silico* and then directly by DNA sequencing. It was found that *dmdA* was not in frame with *lacZ*, but that the *acuI-lacZ* fusion in pBIO2021 did form an inframe fusion. To determine whether this translational fusion affected  $\beta$ -galactosidase activity, a +1 frame shift mutation was introduced into the junction point of the *acuI-lacZ* gene fragment. The mutagenic primer, acuI+1\_NsiR, was used with SPO1913/14\_XbaF to PCR amplify the fragment of the *dmdA-acuI* region from pBIO2019 used to create pBIO2021

(*acuI-lacZ*). The acuI+1\_NsiR primer introduced an additional C nucleotide 2 base pairs 5' of the *Nsi*I cut site, creating a +1 frame shift in the sequence, shown in Figure 5.3.





When the promoter and start of a gene (*genX*) are cloned in frame with the downstream *lacZ* gene of pBIO1878, the ribosome can translate the mRNA transcript from two ribosomal binding sites (RBS), resulting in a LacZ protein product and a Gen-LacZ hybrid enzyme, because of read-through. Here, the binding efficiency of the inserted RBS may affect the outcome of a  $\beta$ -galactosidase assay. This was the case for pBIO2021, an *acuI-lacZ* transcriptional fusion plasmid, as indicated below the "In Frame" region.

However, if the promoter region is fused out of frame (indicated by the insertion of a C nucleotide in the figure), the RBS for this promoter region will soon hit a nonsense codon and lead to a non-functional, deleted  $\beta$ -galactosidase gene product. This was the case for pBIO2084, whereby a C nucleotide (lower case in the sequence) was inserted in the sequence prior to the *Nsi*I restriction site, which represents the junction point with the plasmid bound *lacZ*.

This mutagenised fusion was cloned and transformed as described for pBIO2021. A single successful transformant was confirmed by DNA sequencing as a *bona fide* +1 frame shift mutant. The resulting plasmid, pBIO2084 (Figure 5.1), was transferred via triparental conjugation to *R. pomeroyi* J470 and assayed for  $\beta$ -galactosidase activity, as above. The frameshift mutation had no significant effect compared to its progenitor (Figure 5.2). Therefore, the additional RBS present in pBIO1878 does not appear to affect  $\beta$ -galactosidase activity.

A terminator prediction using TransTermHP (Kingsford *et al.*, 2007) located a stem-loop-tail terminator motif (GGCCCCCGG ACATAGCG CCGGGGGGCC, score 5.2), located 19 bps downstream of *acuI*. This was the only such predicted terminator within the entire *acuI-dmdA* region, again supporting the notion that *acuI* and *dmdA* are co-transcribed, from a promoter upstream of *dmdA*.

Given these results, it is still unclear why the *acuI-lacZ* reporter fusion exhibited higher  $\beta$ -galactosidase activity than the *dmdA-lacZ* reporter fusion.

### 5.2.3 Insertion mutations into *dmdA* and *acuI*

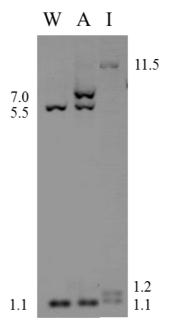
Having shown that *dmdA* and *acuI* are co-transcribed and co-regulated, the functions of these two genes were examined in some detail. This first involved the isolation of insertional mutations into each of these genes followed by phenotypic characterisation of these mutants.

### 5.2.3.1 Construction of DmdA<sup>-</sup> and AcuI<sup>-</sup> R. pomeroyi strains

To make these mutations, fragments internal to *acuI* and to *dmdA* were cloned, separately, into the suicide plasmid vector pBIO1879. The *dmdA* fragment was made by amplifying an 800 bp region from *R. pomeroyi* DSS-3 genomic DNA using primers SPO1913F\_Pst and

SPO1913R Eco (Figure 5.1), which contain *PstI* and *EcoR*1restriction sites, respectively, prior to cloning into a similarly digested pBIO1879 vector, to form pBIO1870. A 957 bp acuI internal DNA fragment was made by digesting pBIO2019 (see above) with SalI, which cuts twice within acul, and this fragment was then cloned into SalI-digested pBIO1879 to form pBIO2025. To recap, pBIO1879 is a pK19mob suicide vector derivative with an additional Spec<sup>R</sup> cassette, which inserts into the host genome via homologous recombination, based on homology to an inserted fragment of DNA, and disrupts the function of the respective gene. This method was used previously in the mutagenesis of the *dddW* gene (see chapter 4). The two plasmids pBIO1870 and pBIO2025 were then each introduced by conjugation into R. pomeroyi J470 in triparental matings, selecting Rif<sup>R</sup>/Spec<sup>R</sup>/Kan<sup>R</sup> transconjugants, which should arise via a single cross-over event within the corresponding genes (See Figure 4.5, chapter 4). Each cross generated  $\sim$ 50 transconjugants with the selected resistances; genomic DNA was isolated from four of these, and was digested with *Eco*RI. The fragments were separated on an agarose gel, which was then blotted to nitrocellulose and probed with a PCR product that extended from 216 bp upstream of *dmdA* to the 3' end of *acuI*. This had been generated from R. pomeroyi genomic DNA, using primers SPO1913/14\_xbaF and SPO1913/14\_BamR (Figure 5.4) and would label a ~1.1 kbp and ~5.5 kbp fragment in wild type *R. pomerovi*. The DmdA<sup>-</sup> mutant exhibited fragments of *ca*. 1.1, 5.5 and 7 kbp in size, representing the insertion of the ~6 kb pBIO1879 plasmid into the 1.1 kb fragment. The unusual nature of the insertion of pK19 into host genomes (as described in chapter 4) makes accurately calculating the insertion difficult at times. Thus, primers located external to the *dmdA* gene were used to PCR amplify it from both the wild type and DmdA<sup>-</sup> strains. As expected of an insertion of a 6 kb fragment into *dmdA*, a PCR product for *dmdA* was amplified from the wild type, but not the DmdA<sup>-</sup> strain. The AcuI<sup>-</sup> mutant strain exhibited fragments of ca. 1.1, 1.2 and 11.5 kbp, consistent with the insertion of pBIO1879 into the acul gene, and introduction of an EcoRI site from the plasmid ~1.2 kb from an existing EcoRI site in the genome. These mutant Ruegeria pomeroyi DSS-3 strains were termed J527 (AcuI) and J471 (DmdA<sup>-</sup>).

### Figure 5.4 Southern blot of Ruegeria pomeroyi DmdA<sup>-</sup> and AcuI<sup>-</sup> mutants



Genomic DNA from wild type (W), *dmdA*<sup>-</sup> (A), *acuI*<sup>-</sup> (I) strains of *Ruegeria pomeroyi* J470, digested with *Eco*RI and probed with the PCR product that was obtained using primers SPO1913/14\_XbaF and SPO1913/14\_BamR (see Figure 5.1.) which spans both *dmdA* and *acuI* genes. Numbers represent approximate DNA length in kilo base pairs (kbp) of the labelled bands.Note the appearance of a larger band in the DNA from the mutant strains, consistent with the insertion of the pBIO1879 plasmid into the genome.

Prior to examining these two mutants for their phenotypes, a series of plasmids was made (using the methods described above) in which the *acuI* and *dmdA* genes were each cloned alone, and also together, in the wide host-range plasmid pBIO1878, thus allowing the correction of any mutant phenotypes by the corresponding wild type genes to be determined.

To make these constructs, the PCR product (Figure 5.1) that included both *dmdA* and *acuI*, with their native promoter was cloned as a 2045 bp fragment to form pBIO2022. Then, pBIO2024 (*acuI* alone) was created by cloning pBIO2023 (itself a copy of pBIO2019 with a

*Sph*I fragment of *dmdA* deleted –see above) into pBIO1878. Plasmid pBIO2021 comprises the *acuI-lacZ* fusion, and so contains functional *dmdA* but not *acuI*.

### 5.2.3.2 Growth of DmdA<sup>-</sup> and AcuI<sup>-</sup> strains of *R. pomeroyi* on DMSP or acrylate

The following phenotypes were examined for the original DmdA<sup>-</sup> and Acu<sup>-</sup> mutants and for the transconjugants in which the cloned, wild type genes had been introduced into these mutants by conjugational triparental matings.

- (i) Ability to grow on acrylate as sole carbon source;
- (ii) Ability to grow on DMSP as sole carbon source;
- (iii) Tolerance to toxic effects of acrylate;
- (iv) Levels of DMSP-dependent DMS production.

Cultures of the various strains were grown in MBM to an  $OD_{600} \sim 0.8$  before being washed in fresh MBM and spotted in 20µl aliquots onto solid MBM media, in which either DMSP or acrylate (each at 5 mM) was the sole C source. In addition, aliquots of the cultures were plated on MBM containing each of these compounds, plus the "regular" C source, succinate (10 mM) to test the toxicity of DMSP and of acrylate. The results are summarized in Table 5.1.

### Table 5.1 Summary of growth of wild type, DmdA<sup>-</sup> and AcuI<sup>-</sup> strains of *Ruegeria pomeroyi* J470 on DMSP or acrylate, and phenotypic correction with intact *dmdA* and/or *acuI*

R. pomeroyi strain	Growth on 5 mM	Growth on 5 mM	Growth on 10 mM
	DMSP as sole C	acrylate as sole C	succinate in the presence
	source	source	of 5 mM acrylate
J470	+	+	+
J471 (DmdA <sup>-</sup> )	-	-	-
J527 (Acul <sup>-</sup> ) <sup>-</sup>	-	-	-
J471 :: pBIO2019 (cloned	+	+	+
<i>dmdA</i> plus <i>acuI</i> )			
J471 :: pBIO2021 (cloned	-	-	-
dmdA)			
J471 :: pBIO2024 (cloned	+	+	+
acuI)			
J527 :: pBIO2019	+	+	+
J527 :: pBIO2021	-	-	-
J527 :: pBIO2024	+	+	+

+ or – indicates growth or no growth on the tested conditions.

### Data summarised from this study and Todd et al., 2012b.

It was striking that both the Acu<sup>T</sup> mutant (J527) and the DmdA<sup>-</sup> mutant (J471) failed to grow on either DMSP or acrylate, whereas the wild type grew well on both compounds. Not only that, but both mutants failed to grow in the presence of either of DMSP or acrylate (each at 5 mM) even when the alternative C source succinate was supplied.

In view of these observations, an attempt was made to quantify the sensitivity of the DmdA<sup>-</sup> and Acu<sup>-</sup> mutants by spotting aliquots of these strains onto succinate-containing MBM plates

to which varying levels of DMSP and of acrylate, from 0.2 mM to 10 mM were added. Strikingly, the wild type grew well on each of these at concentrations as high as 10 mM (DMSP) or 5 mM (acrylate). In contrast, both the mutants were inhibited for growth at concentrations of 1 mM DMSP and 0.5 mM acrylate. These initial results imply that both *dmdA* and *acuI* are involved in the catabolism of both these substrates.

It was noted that the release of DMS, the other product of the cleavage pathway, was also enhanced 5-fold, when the DmdA<sup>-</sup> mutant was assayed for DMSP lyase activity. This could be explained by the channeling of more of the substrate DMSP into one or more of the DMSP lyase systems, since the demethylation pathway was blocked. This corroborates findings by Bürgmann *et al.* (2007) that a DmdA<sup>-</sup> mutant of *Ruegeria pomeroyi* accumulated acrylate (the other product that is made by cleavage of DMSP by the lyase[s]) from DMSP than the wild type.

### 5.2.3.3 DmdA<sup>-</sup> and AcuI<sup>-</sup> complementation assays

To further investigate the growth phenotypes, a series of complementation tests was done. Thus, the plasmids that contained *acuI* alone (pBIO2024), *dmdA* alone (pBIO2021) or both *dmdA* plus *acuI* (pBIO2022) were introduced into the AcuI<sup>-</sup> and the DmdA<sup>-</sup> mutant strains and the transconjugants were examined for their growth characteristics. As expected, pBIO2022 corrected both mutants for all the defects (growth on and tolerance to DMSP and acrylate). However, both of the plasmids that contained each of the genes, alone, gave some very surprising results.

Thus, the cloned *dmdA* gene, in pBIO2021 did not correct any of the phenotypes, whereas the cloned *acuI* gene, alone in pBIO2024, corrected all of them, even the ability of the DmdA<sup>-</sup> mutant to grow on DMSP as sole C source (Table 5.1). These results can be explained as follows.

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First, it is likely that the insertion of pBIO1879 into *dmdA* has a polar effect on the transcription of the downstream *acuI* gene; therefore the DmdA<sup>-</sup> mutant would actually be defective in both DmdA and AcuI. This would explain the acrylate sensitivity and the failure to grow on this substrate in the DmdA<sup>-</sup> mutant, since the *acuI* gene of other bacteria (*Rhodobacter sphaeroides*, see above) had already been implicated in conferring acrylate resistance and, perhaps in acrylate catabolism (and see also below). It would also explain why neither of these phenotypes was corrected by the cloned *dmdA* gene, alone, but could, as expected, be corrected by the cloned *acuI* gene. The real surprise was that the failure of the DmdA<sup>-</sup> mutant to grow on DMSP could also be corrected by the cloned *acuI* gene, but *not* by the cloned *dmdA*. The implications of this are important and unexpected.

First, it means that growth of *Ruegeria pomeroyi* on DMSP can occur even in the absence of a functional demethylation pathway, indicating that growth on this compound may involve one or more of the DMSP lyases (DddP, DddQ and/or DddW). Furthermore, it seems that this system requires a functional AcuI, either to confer resistance to the acrylate that is generated as the product of these DMSP lyases, and/or to initiate a productive catabolic pathway that uses acrylate and leads to growth of the bacteria.

Taken together, the work described above provides a genetic and perhaps a physiological link between AcuI and both the DMSP demethylation and DMSP cleavage pathways. When regarding the function of the Dmd enzymes (which mediate the demethiolation pathway of DMSP catabolism, see introduction), the growth of a DmdB<sup>-</sup> mutant strain of *R. pomeroyi* on DMSP as a sole C source (Reisch *et al.*, 2011a; 2011b) provides compelling evidence that DMSP cleavage is able to sustain cellular growth by utilising acrylate via the actions of the AcuI enzyme (also see chapter 7). It was therefore important to explore the role of AcuI more thoroughly.

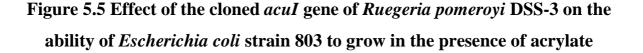
### 5.2.4 A role for AcuI

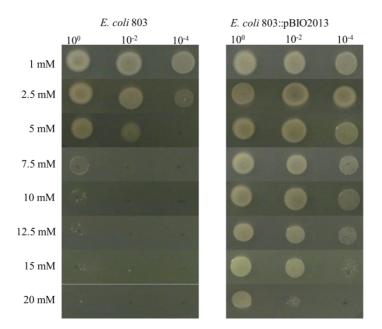
Having shown that AcuI conferred increased tolerance to acrylate and was required for growth of *Ruegeria pomeroyi* on this substrate, the effects of this gene product in other bacteria were investigated. Specifically, its effects in *Escherichia coli* were investigated, as follows.

### 5.2.4.1 Ectopically expressed AcuI confers acrylate resistance to E. coli

Primers SPO1913/14\_BamR and SPO1914\_NdeF were used to amplify a 1017 bp fragment containing the intact *R. pomeroyi* DSS-3 *acuI* gene (0 bp upstream of its start and 24bp downstream from its end). This was ligated into the over-expression vector, pET21a and the ligation mix was added to competent cells of *E. coli* 803, before plating for  $Amp^R$  (encoded by pET21a) transformants. These arose at *ca*. 1.0 x 10<sup>3</sup> cfu/ml and 6 of these were used for plasmid isolation. These plasmids were digested with *Bam*HI and *Nde*I to identify the appropriately sized insert. One of these, termed pBIO2013, was confirmed by DNA sequencing and was used to determine the effects of the cloned gene on acrylate tolerance in this heterologous host.

The *E. coli* 803::pBIO2013 strain and a control lacking the introduced plasmid were grown in LB broth to an OD <sub>600</sub> nm of 1.0, serially diluted and spotted as 20  $\mu$ l aliquots onto solid LB media amended with acrylate, (1 mM – 20 mM) and incubated for 24 hours at 37°C (Figure 5.5).





Cultures of wild type *E. coli* strain 803 with or without the cloned *acuI* gene from *R. pomeroyi* DSS-3, in pBIO2013, were grown in LB medium to an OD <sub>600</sub> nm of ~1.0. The cultures were serially diluted as indicated and 20  $\mu$ L aliquots were spotted onto LB agar plates, supplemented with various levels of acrylate as indicated. Plates were incubated for 24 hours at 37<sup>o</sup>C.

The wild type *E. coli* 803 cells showed signs of growth inhibition at 5 mM acrylate, whereas the presence of the cloned *acuI* gene in pBIO2013 made cells significantly more resistant (Figure 5.5). Cell extracts obtained from this strain were analysed by HPLC following incubation with acrylate, as in Sullivan *et al.* (2011) and Todd *et al.* (2010). The cultures of *E. coli* 803::pBIO2013 were grown overnight at 37°C in LB media, before diluting  $10^{-2}$  into 1 ml M9 minimal medium containing 10 mM glycerol and 2.7 kBq [1- <sup>14</sup>C] acrylate, with unlabelled acrylate added to a final concentration of 2.5 mM. The cells were lysed and pelleted and the supernatant was resolved on a Dionex (Sunnyvale, CA, USA) ICE-AS6 HPLC column. The amount of acrylate dropped by 3-5% in both the wild type and pBIO2013 expressing cells. Also, no new peaks that might be attributed to a catabolite of acrylate, and

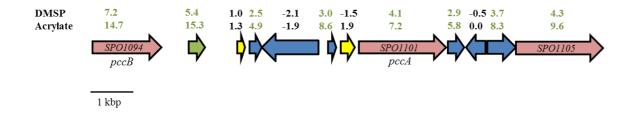
mediated by the presence of the cloned *acuI* gene were seen (M J Sullivan, personal communication).

These observations resemble those of Sullivan *et al.* (2011) who were unable to detect a known catabolite released from acrylate by cell extracts of *Rhodobacter sphaeroides* 2.4.1, which harbours an AcuI homologue. Most importantly, Schneider *et al.*, (2012) showed that the AcuI of *Rhodobacter sphaeroides* 2.4.1. (the very same strain studied by Sullivan *et al.*, [2011]) could catalyse the reduction of acryloyl-CoA to propionyl-CoA. It is thought that acryloyl-CoA can act as a strong electrophile once inside the cell, making it a potentially toxic compound to any organisms that produce it (Herrmann *et al.*, 2005), and this is likely to explain the toxic effects of exogenous acrylate on AcuI mutants.

### 5.2.4.2 Downstream catabolism of acryloyl-CoA

Assuming that AcuI has the same function in *Ruegeria* as in *Rhodobacter*, there must be a route by which propionyl-CoA can be further catabolised in these species. Interestingly this pathway relates directly to a cluster of several closely linked genes whose expression was much elevated by acrylate in the microarrays. As shown in Figure 5.6 and Figure 5.7 these genes were *SPO1094*, *SPO1101*, *SPO0932* and *SPO1105*.

### Figure 5.6 The SPO1094-SPO1105 region in Ruegeria pomeroyi DSS-3



Approximate scale provided. Arrows in red are confirmed to be involved in the catabolism of acrylate:

SPO1094, PccB propionyl-CoA carboxylase B sub-unit;

SP01101, PccA propionyl-CoA carboxylase A sub-unit;

SPO1105, methylmalonyl-CoA mutase.

Green arrow represents a hypothetical polypeptide containing a cupin domain.

Yellow arrows represent putative lipoproteins.

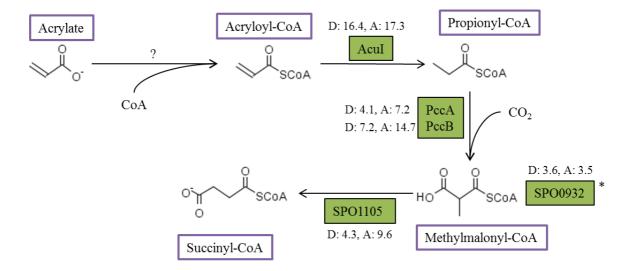
Blue arrows represent hypothetical gene products.

The regulation of each gene in the cluster by DMSP or acrylate is represented by the numbers above the genes, as indicated. Green text illustrates where there is a >2.5 fold increase in transcription over uninduced conditions.

Some of the genes that follow are only potentially (as indicated by a large variance or P-value >0.05) up-regulated in the presence of 5 mM DMSP or 2.5 mM acrylate, by up to *ca.* 15-fold. Three of these genes are strongly predicted to be involved in the conversion of propionyl-CoA to succinyl-CoA (see Figure 5.7).

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The PccA and PccB gene products (encoded by *SPO1101* and *SPO1094*, respectively) are the two polypeptides in the heterodimeric enzyme propionyl carboxylase (EC 6.4.1.3), which is responsible for the conversion of propionyl-CoA to *S*-methylmalonyl-CoA. The SPO0932 (EC:5.1.99.1) gene product is annotated as a methylmalonyl-CoA epimerase, and this enzyme is responsible for the inter-conversion of the *R* and *S* forms of methylmalonyl-CoA (McCarthy *et al.*, 2001). Finally, the *SPO1105* gene encodes a methylmalonyl-CoA mutase (EC:5.4.99.2), which is responsible for the conversion of *R*-methylmalonyl-CoA to succinyl-CoA. This can then be incorporated into the citric acid cycle.



### Figure 5.7 Acrylate catabolism via AcuI in Ruegeria pomeroyi DSS-3

A CoA group is first attached to acrylate to form acryloyl-CoA, mediated by an as-yet unknown enzyme (?). Acryloyl-CoA is reductively converted to propionyl-CoA by AcuI, and then carboxylated by the PccAB/SPO1101 heteromer to *S*-methylmalonyl-CoA. A predicted methylmalonyl epimerase, encoded by *SPO0932*, converts *S*-methylmalonyl-CoA to *R*-methylmalonyl-CoA (\*). This is then acted upon by SPO1105, a methylmalonyl-CoA mutase (EC 5.4.99.2) to produce succinyl-CoA, which may be incorporated into central metabolism via the citric acid cycle.

Green boxes indicate that the genes encoding these enzymes are up-regulated in the presence of DMSP and acrylate, and the associated numbers are the fold change in expression over uninduced cells. D, DMSP; A, acrylate.

Gene numbers in R. pomeroyi DSS-3:

AcuI, SPO1914;

PccA, SPO1101;

PccB, SPO1094.

The gene product responsible for the initial transfer of a CoA group onto acrylate is unknown. However, an acetyl-CoA ligase (EC 6.2.1.1) from *E. coli* has been shown to exhibit low specificity, being capable of attaching CoA to acrylate and other substrate analogues (Patel and Walt, 1987). Thus, an orthologue of this gene in *R. pomeroyi* may act to carry out the initial step in this reaction.

Preliminary work by E Fowler (personal communication) has corroborated some of the microarray data. She showed that a *pccB-lacZ* transcriptional fusion was up-regulated *ca*. 4- and 7- fold in the presence of DMSP and acrylate, respectively, indicating that DMSP may have to be catabolised to acrylate to act as a co-inducer.

Importantly, she also made strains of *R. pomeroyi* with insertional mutations into the *pccA*, *pccB* and *SPO1105* genes and found that they were unable to grow on acrylate or, significantly, on DMSP as sole C source.

### 5.2.4.3 The SPO1094-SPO1105 region

Despite encoding two sub-units of the same enzyme, the *pccA* and *pccB* genes are separated by several genes and large intergenic regions in *R. pomeroyi* (see Figure 5.6). These include 2 large apparently non-coding regions (~1kbp) between *SPO1094*, *SPO1095* and *SPO1096*. There are several genes encoding hypothetical proteins or lipoproteins. The two lipoproteins are encoded by *SPO1096* and *SPO1100*. SPO1096, at 45 amino acids long, has 41%-60% identity (>e<sup>-9</sup>) to putative lipoproteins from other Roseobacter species only. The SPO1100 polypeptide has between 46%-60% identity (>e<sup>-4</sup>) to lipoproteins from other Roseobacters also.

Two other genes that have at least some homology with proteins of known function, or contain known functional domains, are *SPO1095* and *SPO1098*. The *SPO1095* gene is predicted to be in a one-gene operon, with an intergenic region of approximately 1 kbp between its flanking genes (Figure 5.6). A BLASTp analysis retrieves weak homology to

#### Chapter 5: *dmdA* and *acuI*

cupin domain-containing proteins sporadically distributed, such as in a SAR116 cluster bacterium (72%,  $1e^{-37}$ ) and *Azospirillum brasilense* Sp245 (66%,  $9e^{-32}$ ). Due to the possession of a cupin-containing domain like other DMSP lyases (DddL, DddQ and DddW – see introduction and chapter 4) and the induction by DMSP, an *E. coli* pET21a::*SPO1095* clone was created and tested for a Ddd<sup>+</sup> phenotype (J D Todd, personal communication). No DMS was produced when cells were amended with 5 mM DMSP overnight. However, in light of the recent findings that genes present in the same region have effects on acrylate metabolism, and this gene's increased expression in the presence of acrylate over DMSP, it would be worth revisiting this mutant in the future to assay its effect on acrylate and other related compounds.

Finally, the *SPO1098* gene encodes a hypothetical protein in *R. pomeroyi*, which has homologues in a few other Roseobacter species (>65% identity at the peptide level). There are also several considerably less well conserved matches (~26% identity, ~ $e^{-24}$ ) in  $\beta$  and  $\gamma$ -proteobacteria, with most also being annotated as hypothetical. However, in one particular group of  $\gamma$ -proteobacteria, *Acinetobacter* spp., the homologues are annotated as DcaP-like proteins. In *Acinetobacter baumannii* AYE this gene product is described as a porin precursor in catabolism of dicarboxylic acids (hence, DiCarboxylicAcid, DcaP). Porins allow the passive diffusion of hydrophilic molecules through the outer membrane (Benz and Bauer, 1988). The polypeptide was interrogated using the SignalP signal sequence search tool, and a signal peptide was predicted (mean value, 0.882), with a putative cleavage site at position 25. This implies that the gene product is indeed a porin-like protein. Interestingly, succinic acid is a dicarboxylic acid, so perhaps this relates to the other enzymes that are coded by genes in this cluster, converting propionyl-CoA to succinyl-CoA.

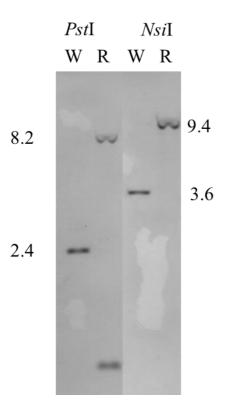
This unusual interruption of the propionyl-CoA carboxylase (PCC) genes is somewhat conserved between members of the Roseobacters, with a little variation. For example, *SPO1095* is only present in *R. pomeroyi*. One Genus in which the 3 "core" PCC genes are found clustered together with no interference is the  $\alpha$ -proteobacterial *Brucella* spp. Perhaps this is indicative of their roles as progenitors for the genes found in the Roseobacters.

#### 5.2.5 A putative *dmdA* regulator, SPO1912 (DmdR)

Separated by 222 bps from the translational start of *acuI* is the ATG of the divergently transcribed gene, *SPO1912* (now designated *dmdR*), which likely comprises a one-gene operon (Figure 5.1). The microarrays showed that the expression of *dmdR* was significantly induced by DMSP and acrylate, by 3.6 and 6.1-fold, respectively. The *dmdR* gene product belongs to the large family of GntR-like repressors, the originally named GntR being found in *Bacillus subtilis*, where it represses the *gnt* genes, unless relieved by addition of the co-inducer gluconate (Fujita and Fujita, 1987). These observations suggest that *dmdR* may be a regulatory gene, controlling the expression of the *dmdA-acuI* operon. Indeed, homologues of the *SPO1912* gene have been noted to exist upstream of 42% of *dmdA* homologues within the Roseobacter clade (Howard *et al.*, 2008).

#### 5.2.5.1 Construction of a DmdR<sup>-</sup> mutant R. pomeroyi strain

To test the regulatory function of *dmdR*, an insertion mutation into the gene was created, in essentially the same way as had been done for the *acuI* and *dmdA* mutations (see above). In this case, primers SPO1912\_EcoF and SPO1912\_PstR were used to amplify a 682 bp fragment, internal to the *dmdR* gene, which was then cloned into the suicide plasmid pBIO1879 (see above, Figure 5.1), creating pBIO2085. This was mobilised into *R. pomeroyi* J470 via triparental mating, selecting for Rif<sup>R</sup>/Spec<sup>R</sup>/Kan<sup>R</sup>. Transconjugants appeared following 2 days incubation ( $1.0 \times 10^2$  cfu/ml). Four of these were re-inoculated to fresh media and genomic DNA extracted. DNA was digested with *PstI* and *NsiI* and the *dmdR* PCR product was used as a probe for a Southern blot, as carried out for J471 and J527, above. This confirmed an insertion into *dmdR*, with the loss of the original *ca*. 2.4 kbp and *ca*. 3.6 kbp wild type fragments, and the appearance of mutated fragments at *ca*. 8.2 kbp and 9.4 kbp for *PstI* and *NsiI* digested DNA, respectively (Figure 5.8); the mutant, strain J530, was retained for further study.



#### Figure 5.8 Southern blot of a DmdR<sup>-</sup> mutant of *Ruegeria pomeroyi* J470

Genomic DNA samples from wild type (W) and a DmdR<sup>-</sup> mutant strain of *Ruegeria pomeroyi* J470 (R), were digested with *Pst*I or *Nsi*I and probed with the *dmdR* PCR product, amplified with primers SPO1912\_EcoF and SPO1912\_PstR. Numbers represent approximate DNA length in kilo base pairs (kbp) of the labelled bands. Note the larger 8.2 kb and 9.4 kb fragments in the DmdR<sup>-</sup> mutant, compared to the 2.4 kb and 3.6 kb wild type fragments, due to the insertion of the *ca*. 6 kb pBIO1879 plasmid into *dmdR*.

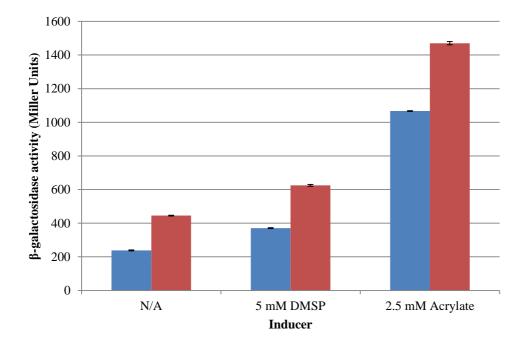
#### 5.2.5.2 Effects of an insertion mutation into *dmdR* on expression of *acuI*

To determine whether a mutation in *dmdR* affected expression of the *dmdA-acuI* transcriptional unit, the *acuI-lacZ* fusion plasmid, pBIO2021, was conjugated via tri-parental mating into the newly created DmdR<sup>-</sup>*R. pomeroyi* (J530), selecting for Rif<sup>R</sup>/Spec<sup>R</sup>/Kan<sup>R</sup>/Tet<sup>R</sup> transconjugants. One of these was purified and cultures were grown in ½ YTSS medium to

which either 5 mM DMSP or 2.5 mM acrylate had been added, with unsupplemented medium acting as the control. These cells were then assayed for  $\beta$ -galactosidase activity.

As shown in Figure 5.9, the mutation in *dmdR* had a little or no effect on the expression of the *acuI-lacZ* fusion compared to the wild type, under any of the growth conditions tested. Thus, DMSP and, to a greater extent, acrylate acted as co-inducers for the expression of the *acuI-lacZ* fusion and there was no evidence that DmdR was a positively or negatively acting transcriptional regulator.

#### Figure 5.9 Expression of an *acuI-lacZ* fusion in wild type and DmdR<sup>-</sup> mutant *Ruegeria pomeroyi* J470



Wild type *Ruegeria pomeroyi* J470 (blue bar) and the DmdR<sup>-</sup> mutant derivative J530 (red bar) containing the *acuI-lacZ* fusion plasmid pBIO2021 was grown in ½ YTSS medium (N/A) or in the same medium supplemented with 5 mM DMSP or 2.5 mM acrylate. Results of triplicate assays of  $\beta$ -galactosidase activities, in Miller Units, with standard error are shown.

### 5.2.5.3 The presence of a cloned *dmdR* gene affects expression of *dmdA-lacZ / acuI-lacZ* in the heterologous host, *Rhizobium leguminosarum*

Despite this result, preliminary work using *Rhizobium leguminosarum* indicated that DmdR may indeed act a transcriptional repressor of the *dmdA-lacZ* fusion (J D Todd, personal communication), in keeping with its homology to the GntR family of transcriptional repressors. This was shown by the fact that when the *dmdA-lacZ* and *acuI-lacZ* fusions were transferred to this heterologous host, both were expressed constitutively at high level. However when the cloned *dmdR* gene was also present, the expression of both fusions was substantially reduced by exposure of the *Rhizobium* cells to either DMSP or acrylate. Perhaps DmdR does indeed respond to these two molecules, but they may need to be converted to the *bona fide* co-inducer, perhaps by a function that is present in *Ruegeria* but not in *Rhizobium*.

#### 5.2.5.4 MMPA also affects expression of *acuI-lacZ* in *R. pomeroyi*

Very recently, the effects of MMPA on the expression of the two fusions were examined in *Ruegeria pomeroyi* and it was found that an *acuI-lacZ* fusion was enhanced, *ca.* 10-fold in cells that were pre-grown in the presence of this molecule (J D Todd, personal communication) in *Ruegeria pomeroyi*. So, here is yet another example in which the product of DMSP catabolism may act as a co-inducer.

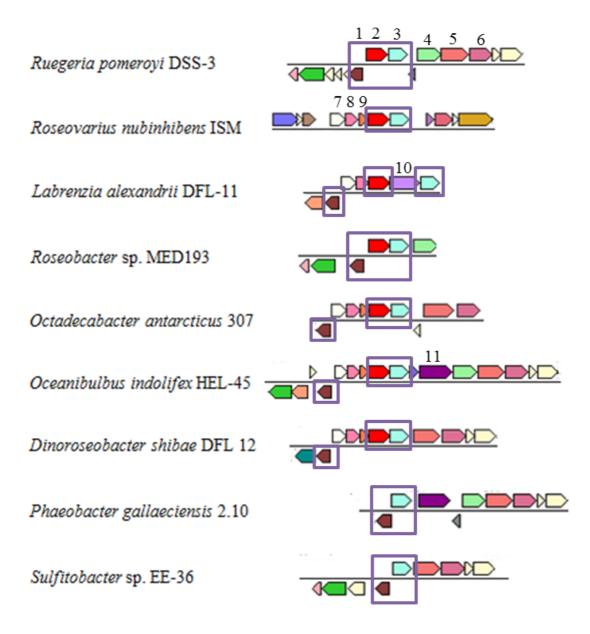
#### 5.2.6 Distribution of the *dmdA* and *acuI* genes

It had already been well-established that the DmdA demethylase was widespread (though not universal) among the Roseobacters; of the 37 sequenced Roseobacter strains, 26 harbour a homologue of DmdA (see introduction; Howard *et al.*, 2006; 2008; 2011; Reisch *et al.*, 2008; Varaljay *et al.*, 2010). Additional homologues occur in the SAR11 and SAR116 Clades and one or two other  $\gamma$ - Proteobacteria. The *acuI* gene has been detected in all of the Roseobacter clade species, including *R. pomeroyi*, that contain *dmdA*. Close homologues (> 70% identical) of DmdR are found in many of the Roseobacters.

#### 5.2.6.1 Close linkage of the *dmdA*, *acuI* and *dmdR* genes

The genes are nearly always closely linked, though there are differences in the precise genetic architecture, shown in figure 5.10.

#### Figure 5.10 Representative comparison of the *dmdA* and/or *acuI* regions



Gene homologues are indicated by coloured arrows. 1, DmdR; 2, DmdA; 3, AcuI; 4, aminotransferase; 5, PhrB deoxyribodipyrimidine photolyase; 6, cyclopropane-fatty-acylphospholipid synthase; 7, hypothetical protein; 8, hypothetical protein; 9, DinB family protein; 10, putative AMP-binding enzyme; 11, Betaine-Carnitine-Choline Transporter (BCCT family). The *dmdR*, *dmdA* and *acuI* homologues are highlighted by purple boxes. Created using IMG. (http://img.jgi.doe.gov) *Dinoroseobacter shibae* DFL12 and *Sulfitobacter* sp. EE-36 harbour intervening genes between the *dmdR* and *dmdA* genes, or no *dmdA* gene at all, respectively. Of the genomes analysed, there were a few cases, such as *P. gallaeciensis* 2.10, in which *dmdR-acuI* were divergently transcribed from one another without *dmdA*, but none in which *dmdR-dmdA* were found without *acuI*. In one case, in *Roseovarius nubinhibens* ISM, a homologue of *dmdR* was not detected at all, having only the *dmdA-acuI* gene pair.

#### 5.2.6.2 Other notable genes present in regions containing dmdR/dmdA/acuI

It was noted that two of the repeatedly encountered genes in the *dmdR/dmdA/acuI* clusters were concerned with DNA repair. Thus, *dinB*, found in *Oceanibulbus indolifex* HEL-45 and *Dinoroseobacter shibae* DFL 12 encodes a DNA polymerase that is important in protection from UV-induced mutagenesis in *E. coli* (Wagner *et al.*, 1999). The *phrB* gene, found in *Ruegeria pomeroyi* DSS-3 and *Octadecabacter anarcticus* 307 encodes a deoxyribodipyrimidine photolyase (EC 4.1.99.3), involved in the repair of pyrimidine dimers that occur from UV radiation (Dorrell *et al.*, 1993). With DMSP catabolising bacteria proliferating mainly in response to events such as phytoplankton blooms, there will be a significant chance that the cells utilising DMSP will be exposed to high levels of UV radiation actually increased cellular DMSP and its lysis to DMS in marine algal cultures. Thus, pairing systems for UV induced DNA repair with genes involved in catabolism of a likely substrate seems evolutionarily advantageous (Levine *et al.*, 2012).

Another gene in proximity to the *dmdA-acuI* genes is a BetT BCCT family choline transporter. The DddT DMSP transporter identified from *Halomonas* HTNK1 and *Marinomonas* MWYL1 (see introduction; Todd *et al.*, 2007; Sun *et al.*, 2011) is a member of this family, so the presence of another BCCT gene near DMSP catabolic genes is enticing.

The other main group of *dmdA*-containing bacteria, *Candidatus* Pelagibacter spp. are different. It is not known why, but they do not have Ddd enzymes so do not encounter

acrylate as is the case for the Roseobacters. However, all three genome-sequenced strains of Pelagibacter have homologues of AcuI (~40% identity at the peptide level to the *Rhodobacter sphaeroides* AcuI), but none is closely linked to *dmdA* (Todd *et al.*, 2012b).

#### **5.3 Conclusion**

The work described in this chapter has greatly enhanced our understanding of a gene that was, at one point, considered peripheral to the activities of *dmdA* (Howard *et al.*, 2008). Instead, *R. pomeroyi* harbours two genes, *dmdA* and *acuI*, each responsible for the activities of a different DMSP catabolic pathway, in a co-transcribed, co-regulated operon, pointing to an unexpected link between the demethylation and lyase pathways – considered more fully in Chapter 7.

#### 5.3.1 Future work

Much work remains to fully reveal the complexity of this system, including:

- 1. Investigate the actions, if any of the DmdR regulator, and identify the *bona fide* cognate coinducer molecule and its gene targets.
- 2. The activity of the accessory genes surrounding, and interrupting, the *dmdR-dmdA-acuI* operon could be established in regards to DMSP metabolism.

# Chapter 6

# Effects of DMSP, acrylate and DMS on the expression of selected genes involved in DMSP, methane thiol and carbon monoxide metabolism

#### **6.1 Introduction**

Previous chapters described genes that are directly related to the catabolism of DMSP (e.g. *dddW, dmdA*) or acrylate (*acuI, pccA, pccB, SPO0932, SPO1105*) in *Ruegeria pomeroyi* DSS-3. In many cases, the expression of these genes was affected by the addition of DMSP, and/or acrylate and/or other metabolically related compounds, as revealed by the microarrays. But, in addition to those genes whose products act directly on DMSP, or acrylate or some immediate downstream catabolites, several other metabolic pathways impinge (or have been proposed to impinge) on DMSP catabolism.

In this chapter, the microarray-based expression of a selection of the other genes that are predicted (though not ratified) by MetaCyc or KEGG (Caspi *et al.*, 2012; http://www.genome.jp/kegg/) to encode some of the enzymes in these "side" reactions is presented, and the implications of their differential expression are discussed.

By their very nature, microarrays are data-heavy. In this study, there were 3 separate growth conditions. This, in conjunction with the 4,252 genes present in the *Ruegeria pomeroyi* DSS-3 genome, resulted in the generation of 12,756 data points, rather too many to interpret for a single thesis. Therefore, in what follows, I will focus only on those cases that satisfy one or more of the following criteria:

- 1. A large difference in expression, >10 fold between one or more of the treatments
- 2. A predicted link with DMSP or DMS catabolism
- 3. Co-regulation of more than one, and ideally all, the genes in a predicted operon
- 4. And, in some cases, ratification of microarray data by the use of lacZ transcriptional fusions

There are some other caveats attached to these analyses, which represent a highly selective sample within the genome-wide response. Firstly, few pathways under consideration here have been verified by direct biochemical studies in *Ruegeria*, or indeed in any Roseobacter strain. Secondly, the approach relies heavily on the correctness of the microarray data, since, in some cases, no other criteria (e.g. qRT-PCR or  $\beta$ -galactosidase assays) were used to

confirm the differential levels of expression that were recorded in the microarrays. This may have been due to a number of factors; difficulties in creating *lacZ* transcriptional fusions, the rapid focus onto AcuI over all else, the isolation of a gene from other genes of interest, no predicted function, lack of regulation of other genes in a putative operon, or highly variable microarray data (see chapter 3).

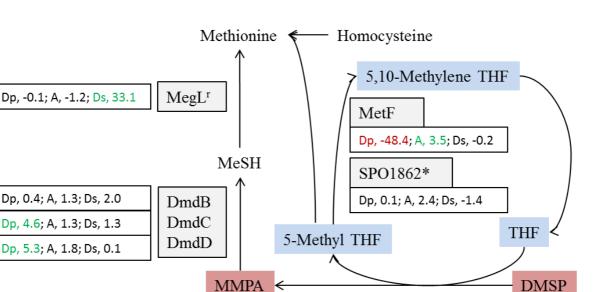
Despite these reservations, the genes discussed in this chapter help build a more complete picture of DMSP catabolism in *Ruegeria pomeroyi*, and in one particular case, open up avenues of research that were never expected.

#### **6.2 Results**

#### 6.2.1 Genes linked to THF and/or methionine

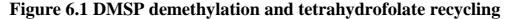
#### 6.2.1.1 The MetF-like enzymes, SPO3016 and SPO1862

The *SPO3016* and *SPO1862* genes have potentially close links to the recycling of the tetrahydrofolate (THF) cofactor that acts as a receptor of one-carbon units from donor molecules, including methyl groups during DMSP demethylation. This has connections to methionine biosynthesis, and the interactions of the relevant gene products are summarised in Figure 6.1.



DmdA

Dp, 23.3; A, 23.8; Ds, -1.4



Gene products are highlighted grey. Fold change in expression, as shown by the microarray data, is given next to each protein: Dp, DMSP; A, acrylate; Ds, DMS. Light blue boxes highlight the various intermediate forms of the co-factor, tetrahydrofolate (THF). Red boxes indicate DMSP and its catabolic product, MMPA, released by demethylation.

\* indicates that transposon disruption of the gene affected DMSP dependent MeSH production (Bürgmann *et al.*, 2007). <sup>r</sup> indicates a hypothesized reversal of the direction of the reaction compared to that which is normally annotated for the function of this gene.

DmdA: DMSP demethylase, SPO1913; DmdB: MMPA-CoA ligase, SPO2045; DmdC: MMPA-CoA dehydrogenase, SPO3804; DmdD: methylthioacryloyl-CoA hydratase, SPO3805;

MetF: 5,10-methylene-THF reductase, SPO3016; SPO1862: MetF-like enzyme;

MegL: methionine γ lyase, SPOA0318.

Adapted from Bürgmann et al., 2007; Reisch et al., 2011a.

2012

In *R. pomeroyi*, it was recently found that the methyl group from DMSP is transferred to THF, to form 5-methyl-THF, as shown in figure 6.1 (Schuller *et al.*, 2012). The production of 5-methyl-THF by glycine cleavage T-proteins (the family to which DmdA belongs – see introduction, chapter 5) is highly unusual and, indeed, had not been previously reported. Rather, the methylated folate products normally take the form of 5,10-methylene-THF (Schübert *et al.*, 2003; Schuller *et al.*, 2012).

Bürgmann *et al.*, (2007) noted that a mutation in a gene, *SPO1862*, abolished DMSPdependent MeSH production in *R. pomeroyi*, but MMPA-dependent MeSH production remained intact. The gene product was annotated as a MetF-like protein, with the canonical MetF (encoded by *SPO3016* in *R. pomeroyi*) being a predicted 5,10-methylene-THF reductase (E.C. 1.5.1.20). The MetF enzyme catalyses the reversible conversion of 5,10methylene-THF to 5-methyl-THF (Figure 6.1), which may then be used by methionine  $\gamma$ lyase to convert homocysteine to methionine (Matthews *et al.*, 1998). The product of the *SPO1862* gene is only 29% identical (with a rather high E value of 2e<sup>-7</sup>, over only 162/300 amino acid sequence coverage) to the SPO3016 polypeptide, and the gene encoding this protein was not induced notably in the microarray under any conditions used here. However, the *SPO3016* gene was down-regulated, 48.4-fold in the +DMSP medium, but was upregulated, 3.5-fold, in acrylate. The *SPO3016* gene product is 43% identical (E = 1e<sup>-83</sup>) to the canonical MetF enzyme in *E. coli* K12 and contains a well-conserved methylene-THF reductase domain (E = 8.6e<sup>-91</sup>). Nearly all of the sequenced Roseobacters contained a gene product that was very similar (>51% identity).

A possible explanation for this regulation is as follows. The MetF (SPO3016) is downregulated in DMSP, causing accumulation of 5-methyl-THF, a by-product of DmdAmediated DMSP demethylation. This cofactor would then be available for the methylation of homocysteine to form methionine, and methionine has been shown to repress *metF* in *S*. *typhimurium* (Cowan *et al.*, 1993).Thus, MetF might mediate a negative feedback loop via the action of DmdA. The phenotype observed in the SPO1862<sup>-</sup> mutant strain may relate to a similar function to MetF, involving the cycling of THF.

#### 6.2.1.2 The methionine γ-lyase, SPOA0318

The *SPOA0318* gene, located on the *R. pomeroyi* DSS-3 megaplasmid, was markedly enhanced (59.6-fold) in one set of data, and slightly enhanced (6.5-fold) in another set, in the presence of DMS, but not of DMSP or acrylate. The gene encodes a protein with a conserved methionine  $\gamma$ -lyase (EC 4.4.1.11) domain (E = 0.0). This enzyme catalyses the conversion of methionine to MeSH and 2-oxobutanoate, in many *Enterobacteriaceae* (Manukhov *et al.*, 2005). The MegL, methionine  $\gamma$ -lyase is an important connection between methionine synthesis and DMSP catabolism, indicated in Figure 6.1. The expression of this gene should be examined by *lacZ* fusions to ascertain whether DMSP is a true inducer or not, as this would fit perfectly with the predicted role it plays.

#### 6.2.1.3 The trimethylamine methyltransferase-like SPO2134

The *SPO2134* gene, located in a single gene unit, was up-regulated 49-fold in the presence of DMSP. This gene encodes a putative trimethylamine methyltransferase (EC 2.1.1.-), with a highly conserved MTTB domain (E =  $2.28e^{-147}$ ). The SPO2134 protein shares 30% identity at the peptide level with the MttB from the archaeon, *Methanosarcina barkeri* (E =  $3e^{-53}$ ). In methanogens such as *Methanosarcina barkeri*, the MttB enzyme catalyses the conversion of trimethylamine to dimethylamine, transferring the methyl group onto a corrinoid protein, Coenzyme M (Paul *et al.*, 2000). The enzymes involved in methionine biosynthesis contain corrinoid domains and utilise the simultaneous binding of a corrinoid, THF and homocysteine to transfer the methyl moiety between intermediates, to form methionine (Matthews *et al.*, 2008). Thus, the MttB-like enzyme in *R. pomeroyi* may play a role in transferring methyl groups between corrinoid proteins for use in methionine biosynthesis, but its function is unknown at present. However, the catabolism of DMSP through to methionine is certainly allusive of this function.

#### 6.2.2 SPO0759

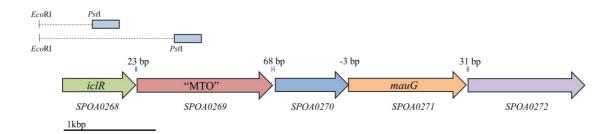
Another gene that was up-regulated by 10.6-fold and 23.6-fold in DMSP and acrylate, respectively, was *SPO0759*. This gene was also down-regulated in the presence of DMS by 3.8-fold. However, the gene encodes a short (107 amino acid) peptide that has no predicted function, with only a few weak homologues (<57% identity, low sequence coverage and  $E>5e^{-12}$ ) in other  $\alpha$ -proteobacteria. Nearby genes include an acetyl-CoA acetyltransferase (*SPO0758*) and a 4-carboxymuconolactone carboxylase domain containing gene (*SPO0760*). However, none of the neighbouring genes is conserved between species.

#### 6.2.3 Ratified DMSP-regulated genes

Following discussion of the genes that had no further experimental evidence for their role in DMSP catabolism, I will discuss two sets of genes that were discovered from the microarray that have been ratified in some way. They have less direct connections to DMSP catabolism. Indeed, the latter set is a complete surprise, with no obvious connection to the pathways described up until this point.

#### 6.2.3.1 Methane thiol oxidase genes

A cluster of genes on the resident megaplasmid was up-regulated by DMSP (*ca.* 2-8 fold increases, but with large variance). A DOOR analysis (Mao *et al.*, 2008) predicted two separate operons, comprising *SPOA0268-9* and *SPOA0270-2* (Figure 6.2).



#### Figure 6.2 The SPOA0268-SPOA0272 region of Ruegeria pomeroyi DSS-3

Scale indicated. Gene names and/or numbers are shown. Sizes of intergenic regions are shown in base pairs (bp). Dashed lines indicate the dimensions cloned to form the *lacZ* transcriptional fusions of *SPOA0268* and *SPOA0269*, with blue boxes representing the *lacZ* junction (J D Todd, unpublished data). MTO, methane thiol oxidase.

Despite these weakly convincing microarray data, subsequent experimental validation confirmed that at least one of these operons was indeed induced by DMSP. J D Todd (Personal communication) constructed a *SPOA0269-lacZ* fusion plasmid (containing an intact *SPOA0268* gene, with its putative promoter, and the 5' region of *SPOA0269*-Figure 6.2) and found that its expression was increased, ~10-fold in DMSP-grown cells.

The *SPOA0269* gene is thought to encode a methane thiol oxidase (MTO) which is 54% identical to the corresponding Hden\_0743 gene product in *Hyphomicrobium* (J D Todd, personal communication). MTO catalyses the reversible conversion of MeSH to formaldehyde, as discovered by Suylen *et al.*, 1987:

 $MeSH + H_2O + O_2 \Leftrightarrow CHO + H_2O_2 + H_2S$ 

DMSP catabolism in *R. pomeroyi* leads to the production of MeSH through DMSP demethylation and possibly DMSP cleavage, so it was decided to test the effects of MeSH itself as a co-inducer. It was found that pre-growth in MeSH of *Ruegeria pomeroyi* containing the *SPOA0269-lacZ* plasmid did indeed induce expression of the fusion, but the ratio of

increase was rather less than with DMSP (see above) indicating that the effect of DMSP was not mediated by its conversion to MeSH.

The related function of the neighbouring genes supports the role of SPOA0269 as a MTO, as it would act to shuttle the formaldehyde produced by the associated enzymes into less toxic intermediates. Several of the genes in this region encode hypothetical proteins, but annotated examples include SPOA0268 as an IclR transcriptional regulator, SPOA0271 as MauG, a methylamine utilisation protein, and SPOA0272 as a glutathione dependent formaldehyde dehydrogenase. The IclR family regulators are diverse, controlling the expression of genes involved in carbon metabolism, aromatic compound degradation and solvent tolerance, to name but a few (Krell *et al.*, 2006). MauG assists in maturation of the methylamine dehydrogenase required to convert formaldehyde to methylamine (van der Palen *et al.*, 1995) and SPOA0272 is a predicted *S*-(hydroxymethyl)-glutathione dehydrogenase (EC 1.1.1.284), carrying out the reaction of *S*-hydroxy-methyl-glutathione to *S*-formyl-glutathione, as described in *Rhodobacter sphaeroides* (Barber *et al.*, 1996). *S*-hydroxy-methyl-glutathione is the spontaneous product of glutathione and formaldehyde.

To investigate the role of the IcIR-like regulator, J D Todd created a transcriptional fusion to the promoter region of *SPOA0268*, this time omitting the intact *SPOA0268* gene (Figure 6.2). When expressed in *Rhizobium leguminosarum* J391, the *SPOA0268-lacZ* fusion plasmid was constitutively expressed under all conditions, at *ca.* 900 Miller Units, compared to *ca.* 100 Miller Units for the *SPOA0269-lacZ* fusion, which raised to *ca.* 400 Miller units in the presence of MeSH. Therefore, SPOA0268 is a MeSH dependent repressor of the MTO in *Ruegeria pomeroyi*. A number of investigations remain, including confirming the operonic structure of this gene cluster and assaying the *Rhizobium* for MeSH removal.

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In addition to those genes with some connection with DMSP and DMS catabolism or formation, the microarrays revealed some wholly unexpected sets of genes with no known link to these compounds, yet whose expression was massively affected by DMS. Among these, the *cox* genes, involved in carbon monoxide catabolism, are described here.

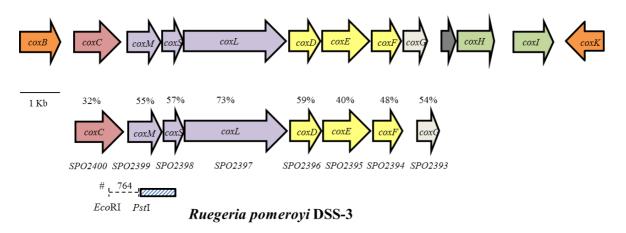
In total, a set of 7 contiguous R. pomerovi cox genes, SPO2399- SPO2393 inclusive, were markedly up-regulated in response to DMS addition, the factors of increase of the individual genes ranging from a remarkable 80.1-fold (coxM – SPO2399) to 3.5-fold (coxE – SPO2395), as summarised in figure 6.3 and table 6.1. To confirm this totally unexpected finding, a transcriptional fusion plasmid was made, in which lacZ was fused to the coxM (SPO2399) in a manner similar to those described in chapters 4 and 5. Thus, primers SPO2399\_ecoRIF and SPO2399\_PstIR were used to amplify a 760 bp fragment that spanned the 3' end of the upstream gene, *coxC*, the *coxC-coxM* intergenic gap and the 5' end of *coxM* (Figure 6.3). Plausible promoter sites had been identified 58 nucleotides upstream of *coxM* in a highly similar cluster in another bacterium, O. carboxidovorans (see below; Santiago et al., 1999) and so this fragment was predicted to contain the *coxM* promoter. The fragment was PCRamplified from *Ruegeria pomeroyi* J470 genomic DNA, and cloned into the *lacZ* reporter plasmid, pBIO1878 (Figure 6.3). The resultant recombinant plasmid was introduced into E. *coli* 803 and transformants arose at a frequency of  $\sim 5 \times 10^3$  cfu/ml. Six colonies were selected for analytical digestions using EcoRI and PstI, with the plasmid DNA from one of the transformants that contained an insert of the expected size being sent for DNA sequencing to confirm that the correct fragment had been cloned. This plasmid, pBIO2086, was then introduced in a triparental mating from the E. coli host into Ruegeria pomeroyi J470 as a recipient. A purified transconjugant was then grown overnight in MBM minimal media, before splitting the culture, with one aliquot being grown for a further 5 hours in the presence of 10 mM DMS and the other being used as the control. The  $\beta$ -galactosidase activity was *ca*. 4-fold greater in the cultures exposed to DMS than in the control, consistent with the microarray data. The absolute level of  $\beta$ -galactosidase activity itself was quite low, at 145  $\pm$  2

Miller Units for +DMS conditions,  $35 \pm 10$  Miller Units for un-induced. Nevertheless, it was confirmed that at least one of the *cox* genes is induced in DMS-grown cells.

#### 6.2.3.3 The cox gene cluster in Ruegeria pomeroyi and Oligotropha carboxidovorans

The *cox* genes have been studied in most detail in another  $\alpha$ -proteobacterium, *Oligotropha carboxidovorans* OM5 (Meyer *et al.*, 1993). As shown in Figure 6.3, there is remarkable conservation of the *cox* gene order and of the sequences of the Cox gene products in *O*. *carboxidovorans* and in *R. pomeroyi*.

### Figure 6.3 The *cox* gene clusters of *Oligotropha carboxidovorans* OM5 (pHCG3) and *Ruegeria pomeroyi* DSS-3



Oligotropha carboxidovorans OM5, pHCG3

Scale bar given. Gene names given, with gene numbers for *R. pomeroyi* provided below arrows. Coloured arrows indicate known or predicted gene functions: red, putatively regulatory; purple, CO dehydrogenase enzyme; yellow, maturation; green, cytoplasmic; orange, putative transmembrane; light grey, unknown; dark grey, pseudogene. The %age identities between the gene products in the two bacteria are shown between genes. # indicates region used to create pBIO2087, with the blue striped bar indicating fusion to the *lacZ*.

By a mixture of biochemistry and genetics it had been shown for *O. carboxidovorans* that the

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carbon monoxide dehydrogenase (CODH) enzyme itself is a heterotrimer, consisting of: CoxL, the large subunit; CoxM, the medium, FAD-containing polypeptide; and CoxS, the small subunit, which contains an FeS cluster; these are encoded by the corresponding genes *coxL, coxM* and *coxS*. Other Cox polypeptides (CoxD, CoxE and CoxF) are also required for the ability of CODH to oxidise CO to  $CO_2$  and may be involved in some form of "maturation" of the enzyme. This may involve the correct insertion of a Mo-containing co-factor that is found in CODH, and also in the related enzymes aldehyde oxidase and xanthine dehydrogenase (Pelzmann *et al.*, 2009). There is also a *coxG* gene following *coxDEF*. A CoxG<sup>-</sup> mutant strain of *O. carboxidovorans* retained CO oxidising activity, however, so it is not clear why this should be called a *cox* gene (Schübel *et al.*, 1995; Santiago *et al.*, 1999).

There are striking similarities with the arrangement and composition of the gene clusters, and a high level of sequence conservation of the polypeptides in the two species, ranging from 32% to 73% identity. However, there are also some significant differences in the Cox systems of *Ruegeria pomeroyi* and *O. carboxidovorans*, as follows.

Many Roseobacters, such as *R. pomeroyi*, lack the ribulose-1,5-bisphosphate carboxylase/oxygenase required for carbon fixation, so cannot assimilate it (Moran *et al.*, 2004; Tolli *et al.*, 2006; Newton *et al.*, 2010). Also, these Roseobacters lack a homologue of an unusual, carbon monoxide-resistant cytochrome b<sub>561</sub>, which allows *O. carboxidovorans* OM5 to respire even in the presence of CO. CODH generates a proton gradient by channelling electrons into a CO-insensitive respiratory chain from this cytochrome (Meyer *et al.*, 1990). The combination of these genes allows the organism to grow with H<sub>2</sub> plus CO<sub>2</sub> under chemolithoautotrophic conditions (Santiago and Meyer, 1997), but Roseobacters cannot. However, it is thought that the Roseobacter species may use CO as a potential supplementary energy source, providing some reducing equivalents and perhaps accounting for the widespread distribution of the *cox* genes in this clade (Moran *et al.*, 2004; 2007; Newton *et al.*, 2010; Cunliffe, 2010). The *R. pomeroyi cox* gene cluster itself lacks several genes, namely *coxB*, *coxH*, *coxI* and *coxJ*, that *O. carboxidovorans* harbours (see Figure 6.3). The CoxHI proteins are thought to be involved in interaction of the CODH with the cytoplasmic membrane in *O. carboxidovorans* (Meyer *et al.*, 1990). No homologues of *coxHI* were detected within the Roseobacter Clade, and this fits with the absence of any proteins that require a proton gradient to be maintained between the CODH and the cytoplasmic membrane (see above). The CoxB and CoxK enzymes have no confirmed function at present, and are merely labelled as putative transmembrane proteins. A summary of these gene products is given in Table 6.1.

## Table 6.1 Comparison of cox genes from Oligotropha carboxidovorans andRuegeria pomeroyi

Gene <sup>a</sup>	Predicted function of	Identity to R. pomeroyi	Factor of
	gene product <sup>b</sup>	homologue, E value <sup>c</sup>	DMS
			Induction <sup>d</sup>
coxB	Putative	No significant match in <i>R</i> .	-
	transmembrane protein	pomeroyi genome	
coxC	Transcriptional	SPO2400, 32%, 6e <sup>-54</sup>	0.2
	regulator		
coxM	CODH medium subunit	SPO2399, 55%, 6e <sup>-108</sup>	80.1
coxS	CODH small subunit	SPO2398, 57%, 3e <sup>-67</sup>	31.9
coxL	CODH large subunit	SPO2397, 73%, 0.0	21.2
coxD	CODH active site	SPO2396, 59%, 4e <sup>-112</sup>	16.9
	maturation		
coxE	CODH active site	SPO2395, 40%, 8e <sup>-91</sup>	3.5
	maturation		
coxF	CODH active site	SPO2394, 48%, 6e <sup>-78</sup>	8.0
	maturation		
coxG	Unknown	SPO2393, 54%, 2e <sup>-42</sup>	7.3
coxH	Possible interaction	CoxC, 27%, 9e <sup>-16</sup>	-
	with cytoplasmic		
	membrane		
coxI	Interaction with	SPO2640 <sup>#</sup> , 31%, 3e <sup>-38</sup>	-
	cytoplasmic membrane		
coxK	Putative	No significant match in <i>R</i> .	-
	transmembrane protein	pomeroyi genome	
lon*	Regulatory protease	SPO2613, 69%, 0.0	-

Name of gene in *Oligotropha carboxidovorans* OM5, column (a), predicted protein function (b), homology between peptides from *R. pomeroyi* and *O. carboxidovorans* OM5 (Locus tag, % identity and e value) (c) and -fold induction by DMS (d) are given. The SPO2640 (<sup>#</sup>) protein shares low identity with CoxI and the gene is not located in the *cox* operon. The *lon* gene (\*) is not in the *cox* gene cluster (Santiago *et al.*, 1999). Grey-shaded entries indicate the genes in the *cox* cluster in *R. pomeroyi*.

#### 6.2.3.4 Carbon monoxide dehydrogenase (CODH) activity

Despite these differences, it is nevertheless clear that the *cox* genes identified here in *Ruegeria* are critical for CODH activity. Recent work by S Newton-Payne (personal communication) isolated a SPO2397<sup>-</sup> (CoxL) mutant and M Cunliffe (personal communication) showed that it was unable to oxidise CO to CO<sub>2</sub>. However, in initial experiments, there was no evidence that pre-growth of *R. pomeroyi* in DMS-containing medium caused any increase of the actual CODH activity, as measured by the removal of CO that had been added to the cultures (M Cunliffe, personal communication). Interestingly, CODH activity is not induced by the substrate CO itself in *R. pomeroyi* (Johnson *et al.*, 2007), whereas in *O. carboxidovorans* CO does act as a co-inducer of the *cox* genes, but these are repressed if other carbon sources are available (Santiago *et al.*, 1999).

#### 6.2.3.5 A cox cluster regulator

From its location relative to the structural *cox* genes, *coxC* is an *a priori* candidate for a *cox* regulatory protein. A BLASTp analysis of CoxC showed that it has two identifiable domains. The N-terminal signalling domain, contains two trans-membrane helices and is weakly predicted ( $E = 2.3e^{-7}$ ) to be in the MHYT superfamily. It was proposed that this domain senses an exogenous signal, with some members of the family having been shown to respond to Cu or NO (Galperin *et al.*, 2001). The other, C-terminal domain, is in the LytTR

superfamily ( $E = 3.32e^{-21}$ ) of non-helix-turn-helix DNA-binding domains found in a variety of bacterial transcriptional regulators (Galperin, 2008). Experimental evidence shows that polypeptides possessing LytTR domains act as transcriptional activators, and often exist in two-component response regulator systems (Gao *et al.*, 2007; Galperin *et al.*, 2008).

There are also conserved homologues of *R. pomeroyi coxC* gene products in 12 other Roseobacter genomes, (>42% identity,  $E < 3e^{-82}$ ), several *Bradyrhizobium* spp. (~30% identity) and *Sinorhizobium meliloti* strains (~28%), in the same locations relative to the other *cox* genes. This implies a similar (regulatory?) role for this gene in a range of different COoxidising bacteria, even those with no known link to DMS, as in *Bradyrhizobium* spp. It should be noted that the CoxC protein was the least conserved of all the Cox peptides, with only 32% identity, perhaps implying divergent functions in the different species.

There may well be another regulatory protein involved in the control of this gene cluster, namely the Lon protease which is encoded by the *lon* gene. This is located on the 133 kbp megaplasmid (pHCG3) of *O. carboxidovorans* OM5 (Santiago and Meyer, 1997; Santiago *et al.*, 1999; Fuhrmann *et al.*, 2003), as is the *cox* gene cluster and the ancillary *cbb* and *hox* genes for CO respiration. The Lon protein is an ATP-dependent serine protease, present throughout all life forms, with targets including the regulatory proteins such as the transcriptional activator, SoxS (Chung and Goldberg, 1981; Shah and Wolf, 2006). Mutations in the *lon* gene in pHCG3 abolished growth on CO (Santiago and Meyer, 1997; Santiago *et al.*, 1999). A BLASTp of the Lon protein from *O. carboxidovorans* retrieved a homologue (SPO2613) in *R. pomeroyi* with 69% identity at the peptide level (E = 0.0). In *R. pomeroyi*, the *lon* gene orthologue is induced by 2.2-fold by DMS, but not by any other co-inducer tested.

It is striking that several bacteria, such as *Bradyrhizobium*, as well as the model *Oligotropha* itself are almost certainly never exposed to DMS, yet, these too contain the strongly predicted CoxC and Lon regulators. Although unlikely, it needs to be established if a similar induction of their *cox* genes is also mediated by exposure to DMS. If so, then this suggests that this observation is some sort of chance molecular aberration, with no adaptive value. But, if

DMS-dependent induction of the *cox* operon is found only in the Roseobacters, then this points to some sort of functionality, and would prompt a search for the mechanism(s) involved.

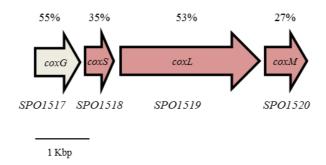
This would involve the following types of experimentation:

- 1. Isolation and characterisation of CoxC<sup>-</sup> mutants.
- 2. If CoxC is involved, establish (by {e.g.} two-hybrids) if it interacts with other polypeptides, such as Lon.
- 3. Confirm by biochemical assays if Lon does indeed act as a protease on CoxC.
- 4. Use *in vitro* binding studies to examine CoxC binding to possible operator sequences and establish if DMS affects this interaction.
- Locate the promoters that are regulated by CoxC (and/or other transcriptional regulators) by RACE or primer extensions.

This story is further complicated by the fact that *R. pomeroyi* DSS-3, and several other Roseobacter strains contain not just one set of *cox* genes, but may have a second cluster, as described below.

#### 6.2.3.6 Distribution of cox gene clusters

*Ruegeria pomeroyi* DSS-3 possesses another predicted operon that contains genes that encode Cox polypeptides that are homologues of CoxG (SPO1517), CoxS (SPO1518), CoxM (SPO1519), and CoxL (SPO1520), which are 35%, 27% and 53% identical to the corresponding products of the genes in the *SPO1517-SPO1520* cluster (Cunliffe, 2010; Figure 6.4). The genes in this second cluster encode the so-called form II CODH, and, strikingly, these too were shown in the microarrays to be up-regulated by 3.2-5.2 fold in the presence of DMS.



#### Figure 6.4 The cox form II operon present in Ruegeria pomeroyi

Scale indicated. Coloured arrows show carbon monoxide dehydrogenase (CODH) form II gene products. Numbers below gene names gives corresponding gene number. The %age identities to the respective form I CODH polypeptides present in *R. pomeroyi* are given above arrows.

This second group of genes is most commonly found in members of the Roseobacter Clade, Rhizobiales Order (containing *Oligotropha carboxidovorans*) and  $\beta$ -proteobacteria, including species of *Burkholderia* (see also Cunliffe, 2010). Many of these homologues are labelled as xanthine dehydrogenase, and *R. pomeroyi* also possesses several other xanthine dehydrogenases in its genome, including *SPO3019*, whose gene product resembles the SPO2397 (CoxL) enzyme (31% identity, 8e<sup>-107</sup>). The function of the second form of CODH is putative in *Oligotropha* (Cunliffe, 2010), although *Mesorhizobium loti* USDA 3471, which contains only the second form of CODH, is known to oxidize CO (King, 2003). The relation to xanthine dehydrogenase may stem from the similarities in the enzymes' molybdenum binding domains and active sites, although at present this is conjecture only. It would be of great interest to see if this cluster, expressed in the *Rhizobium* strain that contains the other *cox* cluster (see above), is able to restore CO oxidising activity.

#### 6.2.3.7 Conserved promoter motifs

A MAST/MEME motif search (Bailey *et al.*, 2009) using the regions 200 bp upstream of the ATG of *R. pomeroyi* form I *coxM* and form II *coxS* (the beginning of each respective *cox* gene cluster, excluding *coxC*) did not retrieve any significant motifs that could be regarded as regulatory, e.g. inverted tandem repeats. Nor was a significant motif identified upstream of the start of the *coxG*, from *O. carboxidovorans* or *R. pomeroyi*. Note that this was the gene for which there was a considerable intergenic gap preceding it in *R. pomeroyi*, as to indicate a possible separately transcribed gene.

#### **6.3** Conclusion

The work carried out in this chapter has touched upon a number of disparate systems in *R*. *pomeroyi*, with indirect and sometimes unfathomable links to DMSP and MMPA catabolism. From THF recycling, to methane thiol oxidation, to carbon monoxide oxidation, the work described opens up several highly interesting avenues of research, despite the "handicap" of high variance and inherent unreliability of the microarray data.

# **Chapter 7**

### **General Discussion**

#### 7.1 Preamble

For all the (somewhat disappointing, and frustrating) problems of the high variances in the microarray data, they nonetheless revealed some novel and wholly unexpected genes whose expression was affected by one or more of the DMSP-related molecules that were tested.

In what follows, I present a brief overview of some of these newly discovered genes and/or links between previously described genes and DMS(P), reserving the most detailed discussion for a consideration of the ways in which this work, together with that of others, has thrown fresh light on the interactions between the DMSP demethylation and cleavage pathways.

#### 7.2 The DMSP lyase, DddW

I have shown that the novel DddW lyase is rare compared to most of the other DMSP lyases, both in the deduced proteomes of known bacteria and in bacterial metagenomic data sets. Collaborative work, with Mishto Dey (University of Iowa) is now underway to obtain the crystal structure of DddW, which would be the first DMSP lyase to be studied in such detail and be informative in showing if and how this cupin-like protein interacts with a metal (if that is in fact the case).

One other striking feature of DddW is that its expression is massively induced by the DMSP substrate, this being mediated by the transcriptional regulator that is encoded by the divergently transcribed *SPO0454*. It will be of interest to show exactly how this regulatory gene product interacts with its *cis*-acting DNA target(s) and with (presumably) the co-inducer DMSP. Electrophoretic mobility shift assays would ratify the binding of this regulator to the putative gene(s) in the neighbouring region. Also, a pan-genomic study (such as chromatin immunoprecipitation or chIP-on-chip) would show if this regulator acted on any other genes in *R. pomeroyi*.

#### 7.3 The cox genes

The novel and completely unexpected DMS-mediated regulation of the *cox* genes is of interest for a number of reasons, and raises a number of interesting questions, ripe for future study.

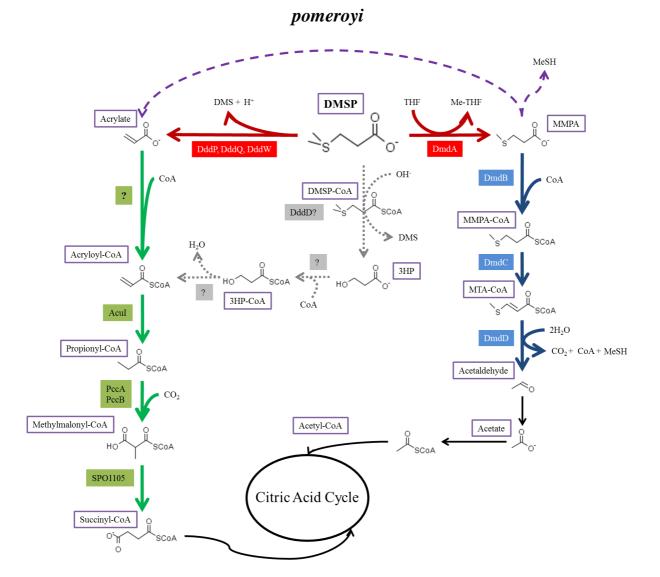
In cooperation with Michael Cunliffe (Plymouth), we will confirm the function of these gene products in *R. pomeroyi* by characterising the phenotypes of the various  $Cox^-$  mutants that have been made. Given that, unlike *Oligotropha carboxidovorans*, the Roseobacters are unable to grow on carbon monoxide – for reasons set out in chapter 6 – it will be informative to determine if the CODH enzyme(s) in these bacteria contribute some input to the energy generation, though not enough to support stand-alone growth.

There is still much to do to understand the mechanisms that underlie the DMS-dependent regulation of the *cox* genes – how widespread is this phenomenon?, what is the transcriptional regulator?, and how does this respond to DMS?, are but some of the questions to arise from this microarray based finding.

Possibly the most far-reaching set of observations that stem from the work in this thesis relate to those that impinge on the relationship between the demethylation and the DMS-emitting cleavage pathways. These are discussed in the following section.

## 7.4 Interplay between demethylation and cleavage pathways for DMSP catabolism

The findings and models set out below build on the pathways – ratified and conjectured – shown in figure 7.1.



#### Figure 7.1 Biochemical pathways for DMSP degradation in Ruegeria

Enzymes and predicted gene products that are involved in the dimethylsulfoniopropionate (DMSP) cleavage (green boxes) and demethylation (blue boxes) pathways in *R. pomeroyi* are shown, with the primary enzymes that act on DMSP itself highlighted in red boxes.

The grey boxes highlight the putative DddD-mediated reaction that is yet to be confirmed in *R*. *pomeroyi*, based on known, functional DddD enzymes and a putative pathway predicted by Schneider *et al.*, 2012 (indicated by ?).

The purple, dashed arrow indicates a putative breakdown of methylmercaptopropionate (MMPA) to acrylate (or possibly propionate) and methane thiol (MeSH) – see text. MTA, methylthioacryloyl.

Gene numbers in R. pomeroyi DSS-3:

*dddP*, *SPO2399*; *dddQ*, *SPO1596*; *dddW*, *SPO0453*; *dddD*, *SPO1703*;

acuI, SP01914; pccA, SP01101; pccB, SP01094;

dmdA, SPO1913; dmdB, SPO2045; dmdC, SPO3804; dmdD, SPO3805.

Adapted from Curson et al., 2011a; Reisch et al., 2012a; 2011b.

#### 7.4.1 Co-regulation of the *dmdA* and *acuI* genes

Importantly, we now know that the *dmdA* and *acuI* genes of *R. pomeroyi* are co-regulated, not only by DMSP, but by its cleavage catabolite, acrylate. Indeed, acrylate was a more potent inducer for both these genes than DMSP, implying a previously unconsidered link between DMSP cleavage and DMSP demethylation. Until now, *acuI* was relatively ignored in *R. pomeroyi*, with previous work concentrating almost solely on the *dmdA* gene (*e.g.* Howard *et al.*, 2006, Reisch *et al.*, 2008).

The genes neighbouring *dmdA* had only been touched upon prior to this thesis. Howard *et al.*, (2006) noted a predicted regulatory gene that was upstream of *dmdA* not only in *R. pomeroyi* but in several other Roseobacters, and speculated that its GntR-like product might control *dmdA* expression, but this was not checked experimentally. This surmise has now been confirmed, as shown in chapter 5. The gene that is known to be *acuI*, was described as a generic "dehydrogenase" by Howard *et al.* (2006).

#### 7.4.2 Growth of AcuI and DmdA mutants on DMSP or acrylate

The phenotypes of the *acuI* and *dmdA* mutants raised some intriguing questions about the roles of these genes in the catabolism of DMSP and acrylate. This work showed that wild type *R. pomeroyi* could grow effectively on both DMSP and on acrylate as a sole C source.

Intriguingly, a mutation in *acuI* abolished growth on both compounds, yet (essentially) a mutation in *dmdA* did not. Despite the plethora of DMSP catabolic enzymes, most Roseobacters do not grow well – or at all – with DMSP or acrylate as sole C sources, at least under the laboratory conditions tested to date (E Fowler, unpublished). Indeed, for reasons that are not clear, González *et al.* (1999) originally stated that *R. pomeroyi* did not grow on acrylate, but then came to the opposite conclusion in 2003 (González *et al.*, 1999; 2003).

In my hands, not only were Acu<sup>T</sup> strains unable to utilise acrylate as a sole C source, but they were sensitive to its toxic effects. Thus, AcuI appears to protect the cells against acrylate or its catabolites (most likely acryloyl-CoA, see below) as well as allowing growth on this substrate.

#### 7.4.3 A possible link between DMSP demethylation and cleavage

At first sight, the above observation on the mutant phenotypes might provide an explanation for a surprising result presented by Reisch *et al.* (2011b). They noted that *R. pomeroyi* DmdB<sup>-</sup>, DmdC<sup>-</sup> and DmdD<sup>-</sup> mutants still grew on DMSP as sole C source, which rather flies in the face of the previous belief that demethylation is the primary route of C assimilation from DMSP. They apportioned this to the fact that the cleavage pathways, mediated by DddP, DddQ and/or DddW might supply sufficient acrylate for growth.

However, Reisch *et al.* also noted that the DmdC<sup>-</sup> and DmdD<sup>-</sup> mutants failed to use MMPA as sole C source. The growth of DmdB<sup>-</sup> mutant was explained (but not checked) by their suggestion that there are two copies of *dmdB* in *R. pomeroyi*. One other striking observation by Reisch *et al.* was that not only did the DmdC<sup>-</sup> and DmdD<sup>-</sup> strains fail to grow on MMPA but they were sensitive to this compound. In light of the work presented in this thesis, this inhibition might be due to the build-up of potentially toxic MMPA- and/or MTA-CoA intermediates in such mutants, rather than a failure to use MMPA as a C source (see Figure 7.1). If true, this raises some doubts as to whether this pathway is indeed the most important even for the degradation of MMPA, let alone DMSP.

A closer examination of the experimental approach taken by Reisch *et al.* also casts some doubt on the nature of their conclusions. Firstly, the pathway was totally restricted to the identification of enzymes (and hence the corresponding genes) that generated catabolites that included a CoA moiety, and not the full panoply of catabolites *in toto*. Also, they did not measure the flux through this pathway, compared to that through cleavage, and/or any other routes for DMSP/MMPA catabolism, so it is not possible to assess their relative importance.

#### 7.4.4 Consideration of an alternative MMPA degradation pathway

Prior to the identification of any of the relevant genes and enzymes, it had been thought that the demethylation pathway was the main route by which the Roseobacters <u>grew</u> on DMSP, with the cleavage pathways providing only a minor contribution to nutrition (Kiene, 1996; Kiene *et al.*, 2000). This was based on <sup>35</sup>S DMSP tracer experiments tracking the accumulation of MeSH following DMSP addition. However, the recent descriptions of the phenotypes of the mutations in the various *dmd* genes (Reisch *et al.*, 2011b), in addition to the findings in this thesis, do not appear to support this view. However, there is another, non-ratified demethylation pathway described many years ago by Kiene *et al.* (2000) who predicted that MMPA might be cleaved directly, either to MeSH and acrylate, or by a reductive cleavage to MeSH and propionate. The recent discovery of the *dmdBCD*-mediated pathway does not necessarily overturn the veracity of either of these pathways

Let us suppose that there is an (as yet unidentified) enzyme that removes the MeSH from MMPA, to form acrylate. This would provide a second, link between DMSP and acrylate, but this time, this acrylate product is generated as part of the demethylation pathway (Figure 7.1). Thus, MeSH would be seen to accumulate through DMSP demethylation, but not necessarily through the actions of the DmdB, DmdC or DmdD enzymes. This might provide an even more convincing reason why *dmdA* and *acuI* are so closely linked in the Roseobacters and why the *dmdA-acuI* operon in *Ruegeria pomeroyi* responds to acrylate as a co-inducer. Indeed, perhaps the reduced expression of the *dmdA-acuI-lacZ* fusion with *dmdA* removed (pBIO2024) in comparison to one with an intact *dmdA* (pBIO2021), observed in chapter 5, stems from the removal of an additional source of MMPA and thus, acrylate, in those cells. Likewise, the induction of *acuI-lacZ* by MMPA may relate to its eventual catabolism to

acrylate. I have shown through the microarrays that the *pccA*, *pccB* and *SPO1105* genes that are responsible for the downstream conversion of acryloyl-CoA to succinyl-CoA (see Figure 7.1) are up-regulated in the presence of acrylate. Importantly, E Fowler (personal communication) then used this information to make targeted mutations in each of these genes and found that these indeed abolished growth on acrylate as sole C source. Furthermore, she used *lacZ* fusions to confirm that these genes were induced by the presence of acrylate in the medium. Thus, any acrylate formed, by whatever mechanism, could be incorporated into central metabolism for growth.

Moran and colleagues have discussed the presence of a bacterial "switch", which may control the relative importance of the demethylation and DMS-producing cleavage pathways in response to the substrate DMSP, or perhaps UV radiation, or temperature, or any number of stimuli (Sunda *et al.*, 2002; Howard *et al.*, 2006; 2008; Levine *et al.*, 2012). The work described in this thesis shows that it is likely to be more complex than a simple on/off system. Clearly though, the elucidation of this switch, if it does exist, will be crucial to understanding the catabolism of DMSP by *R. pomeroyi*, the Roseobacters, and indeed, all DMSP catabolising organisms.

#### 7.4.5 Future work on the demethylase- cleavage interactions

Firstly, it is important to identify the (strongly predicted) CoA ligase that is responsible for the initial transfer of CoA onto acrylate to form acryloyl-CoA. There may be several genes that are regulated by acrylate in the microarray that perhaps have annotations as CoA interacting gene products (e.g. SPO1813). Growth on acrylate could be assessed in strains with mutations in these genes, to confirm their role in this pathway. Alternatively, random transposon mutagenesis could be employed to obtain mutants that fail to grow on acrylate as a sole C source. The proposed acrylate catabolic pathway also requires biochemical ratification, using radio-labelled <sup>14</sup>C-acrylate – this will be done in the near future by others.

Another goal is to examine the flux of DMSP and acrylate through the different pathways described above, which would involve the use of different, labelled substrates, metabolomics and comparisons of the wild type and the various diagnostic mutants.

It is also important to confirm – or refute – an alternative branch to the demethylation pathway, in which MeSH is cleaved directly from MMPA. This could be done by screening cosmids from gene libraries for any that could generate MeSH from MMPA in a heterologous bacterial background.

Further work on gene regulation is also required. The exact role and function of DmdR needs to be confirmed using (for example) gel-shifts and foot-printing to identify its targets and to establish if and how it interacts with its cognate co-inducer (MMPA? Acrylate?). It would also be instructive to see if the *dmdA* genes that are not located near to *acuI* in Roseobacters and other bacteria were regulated in a similar way to that of *R. pomeroyi*. For example, Pelagibacter spp., also contain *dmdA* but do not possess *acuI* or the DMSP lyases (Todd *et al.*, 2012b).

# 7.5 The DMSP catabolic genes in the "model" Roseobacter, *Ruegeria pomeroyi*, and other bacteria

The work in this thesis has concentrated on the DMSP catabolic systems in *Ruegeria pomeroyi*, which has become something of a model for genetic and biochemical studies on DMSP catabolism in the Roseobacter clade. In fact, our studies indicate that this strain may actually be atypical in regard to its ability to catabolise DMSP. This is because it is one of very few strains of this clade that can use DMSP or acrylate as sole C sources (E Fowler, personal communication). Also, compared to other genome-sequenced Roseobacters it holds the record number (four – DddD, DddP, DddQ, DddW) of different *ddd* genes, although most other Roseobacters also have more than one type of DMSP lyase (see Table 7.1 and Buchan *et al.*,2005; Wagner-Döbler and Biebl, 2006). Outside the Roseobacters, the  $\gamma$ -proteobacterium, *Oceanimonas doudoroffii* has very recently been shown to express functional DddD and DddP enzymes, making it the first non  $\alpha$ -proteobacterium to be able to catabolise DMSP in multiple ways (Curson *et al.*, 2011b).

Despite variation in the complement of Ddd genes throughout the Rosebacters, there is excellent conservation of the acrylate degradation pathway. As such, homologues of AcuI, PccA and PccB were found together in the same species, and all of these also contained DMSP lyases and DmdA homologues. Intriguingly, the  $\alpha$ -proteobacterium, *Candidatus* Puniceispirillum marinum IMCC1322 possesses a DmdA and AcuI homologue, but no Ddd enzyme that results in the production of acrylate. However, neither does it contain the downstream acrylate catabolic enzymes, so the importance of this finding is unknown.

		me	chani	SIII						
					Ddd					
Species or strain	DmdA	Р	L	D	Q	W	Y	AcuI	PccA	PccB
<i>Ruegeria pomeroyi</i> str. DSS-3 (R)	$\mathbf{X}^1$	X <sup>1</sup>		X <sup>3</sup>	$X^1$	X <sup>1</sup>		$\mathbf{X}^1$	$\mathbf{X}^1$	X <sup>1</sup>
Dinoroseobacter shibae str. DFL12 (R)	Х		X <sup>1,2</sup>	X <sup>3</sup>				Х	Х	Х
Rhodobacteraceae bacterium str. KLH11 (R)	Х	Х		X <sup>3</sup>				Х	Х	Х
Rhodobacteraceae bacterium HTCC2150 (R)	Х	Х			Х			Х	Х	Х
Roseobacter litoralis str. OCh 149 (R)	Х	Х						Х	Х	Х
Phaeobacter gallaeciensis str. 2.10		Х						Х	Х	Х
Roseovarius nubinhibens str. ISM (R)	Х	X <sup>1</sup>			X <sup>1,4</sup>			Х	Х	Х
Silicibacter lacuscaerulensis str. ITI- 1157 (R)	Х	X			$X^1$			Х	X	X
Candidatus Puniceispirillum marinum str. IMCC1322 (SAR116) (α)	х			X				X <sup>1</sup>		
Oceanimonas doudoroffii (γ)		X <sup>1,5</sup>		X <sup>1</sup>						
<i>Alcaligenes faecalis</i> M3A (β)							X <sup>1</sup>	$\mathbf{X}^1$	X <sup>6</sup>	$X^6$

# Table 7.1 A selection of bacteria that catabolise DMSP by more than one mechanism

In the left hand column of strains; (R) = Roseobacter;  $(\alpha, \beta, \gamma) = \alpha, \beta, \gamma$ -proteobacteria. Crosses indicate the presence of the corresponding polypeptide, based on BLAST hits that satisfied the following stringency tests in comparison with functionally verified DmdA, Ddd and AcuI polypeptide sequences from the Roseobacters, as follows: DmdA, <e<sup>-85</sup>; DddP, <e<sup>-86</sup>; DddL, <e<sup>-52</sup>; DddD, 0.0; DddQ, <e<sup>-20</sup>; AcuI, <e<sup>-141</sup>; PccA, 0.0; PccB, 0.0 (PccA and PccB have e values of 63<sup>-147</sup> and 7e<sup>-80</sup> in *A. faecalis*, respectively, see below).

<sup>1</sup>: These proteins have been experimentally confirmed to be functional.

<sup>2</sup>: *Dinoroseobacter shibae* str. DFL 12 lacks a Ddd<sup>+</sup> phenotype under laboratory conditions examined to date, but its cloned *dddL* gene confers a Ddd<sup>+</sup> phenotype to *Escherichia coli*.

<sup>3</sup>: These three DddD homologues form an out-group from the known, functional DddD enzymes; DddD of *R. pomeroyi* str. DSS-3 lacks DMSP lyase activity under the conditions sued to date.

<sup>4</sup>: Roseovarius nubinhibens ISM has two DddQ homologues.

<sup>5</sup>: Oceanimonas doudoroffii has two DddP homologues.

<sup>6</sup>: *A. faecalis* has a methylcrotonyl-CoA carboxylase with 33% identity with PccB and an acetyl-CoA carboxylase with 52% identity to PccA in strain NCIB 8687 (there is no entry for M3A in the NCBI data base).

Adapted from Curson et al. 2011a; 2011b; Todd et al., 2012b.

#### 7.6 A widespread role for acrylate degradation enzymes

In a sense, this work started off in a rather parochial, specialised catabolic pathway for the breakdown of a rather esoteric substrate. However, it may now be addressing a phenomenon of much wider importance, namely the extreme toxicity of acrylate (or its CoA derivative) and the way that bacteria deal with it. Work by A R J Curson identified homologues of AcuI in many members of the Enterobacteria, such as *E. coli*. This homologue was termed YhdH, and is a medium chain reductase family protein (MDR012 class), that shares 54% identity with AcuI (Todd *et al.*, 2012b). An *E. coli* K12 YhdH<sup>-</sup> mutant was very markedly inhibited for growth by acrylate, and this could be relieved by introducing different *acuI* genes

(including that of *R. pomeroyi*) *in trans*. It is possible that despite not encountering acrylate in its natural environment, *E. coli* and other bacteria can convert acrylate to acryloyl-CoA, as does *R. pomeroyi*, and that this can be detoxified via AcuI-like enzymes. Additionally, O Burns (personal communication) screened metagenomic libraries of bacterial populations obtained from water-treatment pipes, soil and other, non-marine sources. Several recombinant plasmids were isolated on the basis that they corrected the acrylate sensitivity of the YhdH<sup>-</sup> *E. coli* mutant. Sequencing showed that some of these contained *acuI*-like genes, but others contained genes that encoded short chain dehydrogenase/reductase family proteins (EC 1.3.1.33) and were found in a range of bacterial taxa.

Perhaps all of these genes are merely protective measures independently evolved by different bacterial species to counteract the toxic effect of acrylate (and its CoA derivative), either sourced by DMSP in marine Roseobacters, or exogenously, as may be the case for species such as *E. coli*. Clearly, there are exciting discoveries ahead for the role of acrylate in the marine food web, and beyond, and hopefully this thesis has at least hinted at things to come.

# **Chapter 8**

Materials and Methods

#### 8.1 Strains, plasmids, growth and media

Bacterial strains and plasmids used in this work are shown in Tables 8.4 and 8.5.

#### 8.1.1 Preparation of culture media

All media were prepared using distilled water ( $dH_2O$ ), sterilised by autoclaving at 121°C for 20 minutes. Solid media contained 1.5% (w/v) agar, unless otherwise stated.

*Roseovarius nubinhibens* ISM was grown routinely at 28°C in Marine Broth 2216 Difco (MB).

*Ruegeria pomeroyi* was routinely grown at 28°C in complete ½ YTSS media: 1.25 g Tryptone, 2 g Yeast Extract, 20 g Sea Salts per litre of dH<sub>2</sub>O, adjusted to pH 7.0 (González *et al.*, 2003).

Minimal medium for *R. pomeroyi* DSS-3 was marine basal medium (MBM): 20 g Sea Salts, 250 ml basal media [150 ml 1M Tris HCl pH 7.5, 87 mg K<sub>2</sub>HPO<sub>4</sub>, 1.5 g NH<sub>4</sub>Cl, 375 ml dH<sub>2</sub>O], 700 ml dH<sub>2</sub>O, 10 mM succinate, prior to autoclaving and adding 50 ml FeEDTA [50 mg FeEDTA, 100 ml dH<sub>2</sub>O], 0.1% vitamin solution [2 mg biotin, 2 mg folic acid, 10 mg pyroxidine-HCl, 5 mg riboflavin, 5 mg thiamine, 5 mg nicotinic acid, 5 mg pantothenic acid, 0.1 mg cyanocobalamin, 5 mg p-aminobenzoic acid per 100 ml dH<sub>2</sub>O] following autoclaving (González *et al.*, 1997).

*Escherichia coli* was routinely grown at 37°C on Luria–Bertani (LB) broth: 10 g Tryptone, 5 g Yeast Extract, 5 g NaCl and 1.5 g D-glucose per litre of dH<sub>2</sub>O, adjusted to pH 7.2. (Sambrook *et al.*, 1989).

For growth in minimal media, *E. coli* was grown in M9: 200 ml 5x M9 salts [64 g Na<sub>2</sub>HPO<sub>4</sub>.7H<sub>2</sub>O, 15 g KH<sub>2</sub>PO<sub>4</sub>, 2.5 g NaCl, 5 g NH<sub>4</sub>Cl per litre dH<sub>2</sub>O], 0.1 mM CaCl<sub>2</sub>, 2 mM MgSO<sub>4</sub>, 30 mg Thiamine HCl, 10 mM succinate per litre dH<sub>2</sub>O (an additional 15 mg 187 methionine per litre dH<sub>2</sub>O was required for the Met<sup>-</sup> strain *E. coli* strain 803) (Sambrook *et al.*, 1989).

*Rhizobium leguminosarum* was routinely grown at 28°C in TY complete media: 5 g Tryptone, 3 g Yeast Extract and 0.9 g CaCl<sub>2</sub>.6H<sub>2</sub>O per litre of dH<sub>2</sub>O, adjusted to pH 6.8. (Beringer, 1974).

Liquid cultures were shaken at 200 rpm.

For the purpose of blue/white screens for cloned inserts, *E. coli* JM101 was grown on LB containing ampicillin (to screen for  $Amp^R$  provided by the plasmid e.g. pBluescript), 5-bromo-4-chloro-3-indolyl- $\beta$ -D-galactoside (X-gal – whose cleavage by  $\beta$ -galactosidase yields the blue product, 5,5'-dibromo-4,4'-dichloro-indigo), and isopropyl- $\beta$ -D-thiogalactopyranoside (IPTG – a gratuitous inducer of *lacZ*, located on the plasmid).

# Table 8.1 Antibiotics and supplement concentrations

Antibiotic Solvent		Stock (mg ml <sup>-1</sup> )	Final concentration (µg ml <sup>-1</sup> )		
Ampicillin (Amp)	dH <sub>2</sub> O	100	100		
Gentamicin (Gent)	dH <sub>2</sub> O	10	5		
Spectinomycin (Spec)	dH <sub>2</sub> O	50	200		
Streptomycin (Str)	dH <sub>2</sub> O	200	400		
Tetracycline (Tet)	70% Ethanol	5	5		
Supplement	Solvent	Stock (mg ml <sup>-1</sup> )	Final concentration (µg ml <sup>-1</sup> )		
5-Bromo-4-Chloro-3-Indolyl -β-D-galactoside (X-Gal)	2,2 Dimethyl- formamide	40	40		
isopropyl-β-D- thiogalactopyranoside (IPTG)	dH <sub>2</sub> O	200	200		
<i>Ortho</i> -Nitrophenyl-β- galactoside (ONPG)	dH <sub>2</sub> O	4	800		
Inducer	Solvent	Stock (mM)	Final concentration (mM)		
Dimethylsulfoniopropionate (DMSP)			5		
Acrylate	dH <sub>2</sub> O	100	2.5		
Dimethylsulfide (DMS)	dH <sub>2</sub> O	"Neat"	5		
Carbon source	Solvent	Stock (M)	Final concentration (mM)		
Succinate	dH <sub>2</sub> O	1	10		

#### 8.1.2 Long-term storage

For long-term storage, *E. coli* cultures were grown overnight to stationary phase, glycerol added to a concentration of 25%, then flash frozen on dry ice and stored at -80°C.

*R. pomeroyi* required 48 hour growth and the addition of 15% dimethylsulfoxide, and 15% glycerol before flash freezing and storage at -80°C (González *et al.*, 2003).

#### 8.1.3 Growth curves

For growth curve determination, 10  $\mu$ l of a 5 ml ½ YTSS starter culture of *R. pomeroyi* DSS-3 at OD<sub>600</sub> nm ~1.0 was inoculated to 10 ml MBM medium containing either; no added carbon, DMSP (5 mM) or acrylate (2.5 mM). Readings were taken at timed intervals for 72 hours.

#### 8.2 In vitro and in vivo genetic manipulations

General handling and manipulation of bacterial DNA was carried out as described by Wexler *et al.* (2001).

#### 8.2.1 Phenol:chloroform and midi plasmid preparations

#### 8.2.1.1 Preparation of plasmid DNA using alkaline lysis and phenol:chloroform step

This method of plasmid preparation was used for the small-scale isolation (< 5  $\mu$ g) of plasmid DNA from *E. coli* strains and yielded material of sufficient purity to be used for restriction digestion, PCR or transformation. Prior to DNA isolation, *E. coli* was cultured overnight in LB medium. All centrifugation steps were carried out at 13,000 rpm in a bench-top 5415 Microcentrifuge (Eppendorf).

The procedure is based around the optimised alkaline lysis method of Birnboim and Doly, (1979), followed by phenol: chloroform purification and elution of plasmid DNA. Buffers P1, P2, N3, PB and PE were supplied in the QIAGEN Plasmid Miniprep spin column kit and used according to the manufacturer's instructions.

The supernatant resulting from P1, P2 and N3 treatment steps (as indicated in manufacturer's instructions) was transferred to a clean 1.5 ml microfuge tube and an equal volume (400  $\mu$ l) of phenol:chloroform:isoamyl alcohol (25:24:1) added, then mixed by vortexing for 10 seconds and centrifuged for 2 minutes. The upper aqueous layer was transferred to a clean 1.5 ml microfuge tube (to remove protein from the lysate), then two volumes of room temperature 100% ethanol (700  $\mu$ l) was added and mixed to precipitate the plasmid DNA. This was then centrifuged for 30 minutes and the supernatant discarded. The DNA pellet was washed with 500  $\mu$ l 70% ethanol (v/v) to remove precipitated salts and centrifuged for 5 minutes. The pellet was air-dried and the DNA resuspended in 40  $\mu$ l dH<sub>2</sub>O, and stored at –20 °C.

#### 8.2.1.2 Midi-preparation of plasmid DNA using QIAGEN columns

The QIAGEN plasmid purification procedure was used when large amounts (> 5  $\mu$ g) of high purity DNA from *E. coli* strains were required for multiple procedures and sequencing. The procedure is also based around the optimised alkaline lysis method of Birnboim and Doly, (1979). It comprises three distinct steps: alkaline lysis, column purification using 'Qiagen resin', and precipitation of plasmid DNA. These preparations routinely yielded >70  $\mu$ g of plasmid DNA from *E. coli* cultures, which was free of RNA and protein. Buffers P1, P2, P3, QBT, QC, and QF were supplied in the QIAGEN Plasmid Midi kit and used according to the manufacturer's instructions.

#### 8.2.2 Preparation of genomic DNA

Wizard Mini columns (Promega) were used to prepare genomic DNA suitable for Southern blotting and PCR. The Nuclei Lysis, RNase, and Protein Precipitation solutions were supplied with the Promega kit and used according to the manufacturer's instructions for Gram Negative Bacteria.

#### 8.2.3 Restriction enzyme digestions

DNA was digested using restriction enzymes and buffers purchased from Roche or Invitrogen and were carried out according to the manufacturers' specifications and with supplied buffers. Approximately 1  $\mu$ g of plasmid DNA was routinely digested in a reaction containing 1x restriction buffer and ~10 U of restriction enzyme, made up to a total reaction volume of 20  $\mu$ l with dH<sub>2</sub>O. Digests, unless recommended otherwise, were routinely incubated at 37°C for 2-3 hours, and were stopped by heating samples to 80°C for 10 minutes.

#### 8.2.4 Dephosphorylation of cut vectors

Shrimp alkaline phosphatase (Boehringer Mannheim) was used to dephosphorylate digested vector DNA prior to its use in ligation reactions, according to the manufacturer's specifications, and with supplied buffers. The reaction mixture contained the digested vector DNA, 1  $\mu$ l shrimp alkaline phosphatase and 1X dephosphorylation buffer made up to a total volume of 20  $\mu$ l with dH<sub>2</sub>O. Reactions were incubated at 37°C for ~1 hour, and stopped by heating samples to 80°C for 10 minutes.

#### 8.2.5 Ligation of DNA fragments

T4 DNA ligase (Roche) was used to produce recombinant plasmids from digested vector and insert DNA, according to the manufacturer's specifications and with supplied buffers. The ligation reaction mixture contained ~ 200 ng of linearised vector DNA and insert DNA, 1  $\mu$ l T4 DNA ligase and 1x ligation buffer, made up to a total reaction volume of 20  $\mu$ l with dH<sub>2</sub>O. Ligation reactions were incubated overnight at 4°C.

#### 8.2.6 DNA electrophoresis

DNA fragments were separated by electrophoresis in 1% agarose gels containing 500 ng ml<sup>-1</sup> ethidium bromide (EtBr) and 1X TBE buffer (45 mM Tris-borate, 1 mM EDTA, pH 8.0). Before loading the DNA samples onto agarose gels, 0.2 volumes of 5x loading dye [0.25% bromophenol blue (v/v), 30% glycerol (v/v)] was added. Gels were run in SCIE-PLAS Mini or Midi horizontal gel tanks with 1X TBE buffer at 70-100 V for 1-2 hours with 1 kb Plus ladder (Invitrogen).

#### 8.2.7 Recovery of DNA from agarose gels

DNA separated on agarose gels was purified from the gels using a QIAquick Gel Extraction Kit (QIAGEN). Buffers QG and PE were supplied with the QIAGEN kit and used according to the manufacturer's instructions.

#### 8.2.8 DNA precipitation

Where necessary, DNA was precipitated from solution for sequencing. 1.1  $\mu$ l Na Acetate (pH 4.8) and 28  $\mu$ l 100% ethanol were added per 10  $\mu$ l DNA solution in a 1.5 ml microfuge tube and mixed well. This was incubated on ice for 15 minutes then centrifuged for 15 minutes. Then, the supernatant was discarded and 500  $\mu$ l 70% ethanol added and centrifuged for 5 minutes. The supernatant was carefully removed and the DNA pellet air-dried for 15 minutes.

#### 8.2.9 Bacterial Polymerase Chain Reaction (PCR)

A Primus 2596 thermocycler machine (MWG-Biotech) was used to perform PCR. Unless otherwise stated, standard PCR reactions were set up as follows:

5 μl miniprep DNA/genomic DNA or 1 μl midiprep DNA
1 μl primer 1 (20 pmol μl<sup>-1</sup>)
1 μl primer 2 (20 pmol μl<sup>-1</sup>)
10 μl 10x reaction buffer (Roche)
2 μl deoxynucleoside triphosphates (dNTPS) (10 mM for each nucleotide) (Roche)
1 μl Taq polymerase (Roche)
dH<sub>2</sub>0 up to a final volume of 100 μl

For the specific conditions used for standard PCR see tTable 8.2, and for site-directed mutagenesis PCR, see Table 8.3. PCR products were routinely sequenced to check that they contained the correct sequence.

Stage	Cycles		Temperature	Time (seconds)	
			(°C)		
1	1	(initial denature)	95	240	
2	30	(denature)	95	30	
		(anneal)	60	60	
		(extend)	72	90	
3	1	(final extend)	72	300	

### Table 8.2 PCR temperature cycles for standard PCR

### Table 8.3 PCR temperature cycles for site-directed mutagenic PCR

Stage	Cycles		Temperature	Time (seconds)	
			(°C)		
1	1	(denature)	94	240	
		(anneal)	50	120	
		(extend)	72	120/kb DNA	
				length	
2 18	18	(denature)	94	30	
		(anneal)	56	60	
	(extend)	72	60/kb DNA		
				length	
3	1	(final extend)	72	300	

Cycling parameters were from the Stratagene ExSite PCR-Based Site-Directed Mutagenesis Kit protocol.

#### 8.2.10 Methods for transforming E. coli with plasmid DNA

#### 8.2.10.1 Preparation of competent cells

Competent *E. coli* cells were prepared using CaCl<sub>2</sub>, as described by Sambrook *et al.* (1989). All centrifugation steps were carried out at 6,000 rpm in a bench-top centrifuge at  $4^{\circ}$ C.

A single colony of the desired *E. coli* strain was inoculated to 5 ml LB and incubated overnight at 37°C. Then, 1 ml of the overnight culture was inoculated to 100 ml LB and incubated at 37°C until OD<sub>600</sub> nm was between 0.3 and 0.4. Sometimes, 100 ml LB was inoculated directly from a loop of bacteria on an LB plate grown from a single colony. The cells were cooled on ice and pelleted by centrifugation for 15 minutes. The cells were kept on ice and the supernatant was decanted, before the cells were resuspended in 15 ml of 0.1 M CaCl<sub>2</sub> and incubated for 30 minutes. The cells were pelleted by centrifugation for 15 minutes, resuspended in 2 ml 0.1 M CaCl<sub>2</sub> and stored at  $-4^{\circ}$ C for at least 1 hour. Competent cells could be left at 4°C overnight if desired.

For site-directed mutagenic PCR, XL1-Blue supercompetent cells (Stratagene) were used for transformation and these required no preparation in-house.

#### 8.2.10.2 Transformation of competent cells

Competent *E. coli* cells were routinely transformed with plasmid DNA, as follows. Plasmid DNA or ligation mix was added to 100  $\mu$ l of competent cells in a clean 1.5 ml microfuge tube and incubated on ice for one hour. The cells were heat-shocked at 42°C for 3 minutes then transferred back to the ice for 2 minutes. 0.5 ml pre-warmed LB was added and incubated at 37°C for one hour, inverting the tube occasionally. The transformation mix was plated out on LB containing the appropriate antibiotics, X-gal plus IPTG as appropriate. In transformations involving pUC18-based clones, *E. coli* strain JM101 was used to facilitate blue–white screening of transformants that contained plasmids with inserts.

*E. coli* strain 803 used for the transformation of large (> 10 kb) plasmids.

*E. coli* strain BI-21 was the routine host for the over-expression plasmid, pET21a, used in protein purification experiments.

XL1-Blue supercompetent cells (Stratagene), used in site-directed mutagenesis, were transformed according to the manufacturer's instructions.

#### 8.2.10.3 Plasmid constructions

Construction of specific plasmids is described in the appropriate chapters. In general, the desired gene of an organism was PCR-amplified from genomic DNA with forward and reverse primers, then ligated into the relevant plasmid, digested with restriction enzymes complementary to the sites introduced into the PCR product.

For protein purification, the PCR products were cloned into pET21a, and for transcriptional *lacZ* reporter fusions, PCR products were cloned into the reporter plasmid, pBIO1878. Subcloning of fusions used pBluescript as the first plasmid. A list of plasmids constructed in this work is shown in table 8.4, table 8.5 and table 8.6.

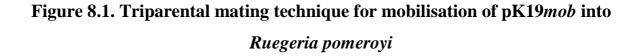
#### 8.2.10.4 Plasmid conjugation

Plasmids were transferred by conjugation to *Ruegeria pomeroyi* or *Rhizobium leguminosarum* from *E. coli* using a patch cross (Johnston *et al.*, 1978). Because the vectors pOT2 and pBIO1878 are not self-transmissible (lacking the *mob* genes), this was done by using triparental crosses with *E. coli* 803, containing pRK2013 as a helper plasmid (Figurski and Helinski, 1979; see Figure 8.1).

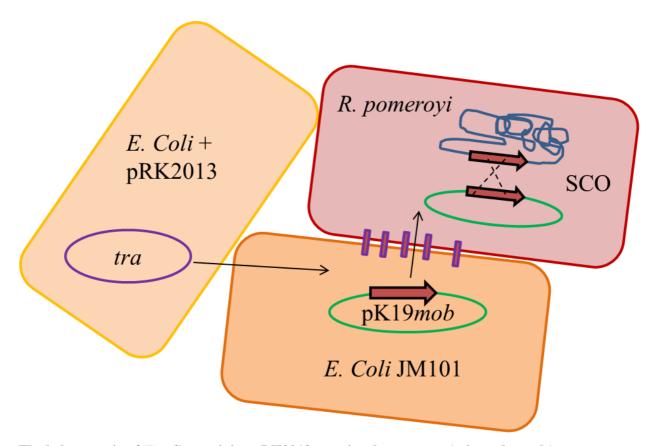
The *Ruegeria pomeroyi* DSS-3 or *Rhizobium leguminosarum* recipient, *E. coli* strain 803 *lacZ*::promoter fusion donor plus *E. coli* pRK2013 (the helper strain) were separately cultured on ½ YTSS (*R. pomeroyi*), TY (*R. leguminosarum*) or LB (*E. coli*) agar plates. Loopfuls of each strain were mixed on ½ YTSS (*R. pomeroyi*) agar, or TY (*R. leguminosarum*) agar. Following overnight incubation at 28°C, the cells were streaked on selective media (½ YTSS for *R. pomeroyi*, TY for *R. leguminosarum*) with the appropriate antibiotics for the plasmid and/or strain. These plates were incubated at 28°C for ~2 days to recover the desired transconjugants.

#### 8.2.11 Insertional mutants in Ruegeria pomeroyi

To generate targeted, insertional mutations into the genome of *Ruegeria pomeroyi*, a modified pK19*mob* system was used. Generally, PCR fragments internal to the gene were amplified from *R. pomeroyi* genomic DNA and then cloned into pBIO1879 using restriction enzymes complementary to the PCR primers used. The pBIO1879 plasmid is a derivative of the suicide insertion plasmid pK19*mob* (Kan<sup>R</sup>) (Schäfer *et al.*, 1994) into which a Spec<sup>R</sup> cassette was cloned as a 2 kb fragment from pHP45W (Prentki and Krisch, 1984) into the HindIII site of pK19*mob*. The resultant plasmid was transformed into *E. coli* strain JM101 and then mobilized by triparental conjugational mating (with pRK2013) into J470 (*R. pomeroyi* DSS-3 Rif<sup>R</sup>), selecting for transfer of Spec<sup>R</sup>, Kan<sup>R</sup> and the mutants selected for on ½ YTSS medium with spectinomycin, kanamycin, with the *E. coli* donor cells being eliminated by rifampicin.



2012



The helper strain of *E. coli* containing pRK2013 contains the *tra* genes (coloured purple), allowing conjugation of the pK19*mob*-derived plasmid (coloured green), containing a fragment internal to the target gene – coloured red, from a host, to the recipient *R. pomeroyi*. Then, selecting for the Kan<sup>R</sup> resistance encoded by pK19*mob spec*, a single cross-over event (SCO) can be chosen, in which the entire suicide plasmid is inserted into the target gene in the host genome (coloured blue), disrupting its function.

Because the recovery of insertional mutations was lower than that of the transfer of the plasmid itself, these conjugations were done as filter crosses, as follows (Beringer and Hopwood, 1976). *Ruegeria pomeroyi* cells were cultured overnight in ½ YTSS liquid at 28°C. *E. coli* containing the pBIO1879-derived mutagenic plasmid, and the helper *E. coli* pRK2013

were grown overnight in liquid LB at 37°C. Then, 0.5 ml of *R. pomeroyi*, 1.5 ml of *E. coli* pBIO1879 donor and 1 ml of *E. coli* pRK2013 was centrifuged, and re-suspended in 200  $\mu$ l of liquid ½ YTSS. This 200  $\mu$ l culture was spread on a sterile nitrocellulose filter on a non-selective ½ YTSS plate. Filters were incubated at 28°C for ~2 days, then the cells were washed off with ½ YTSS liquid, diluted as appropriate and spread onto ½ YTSS agar plates supplemented with spectinomycin, kanamycin and rifampicin. Transconjugants in which the plasmid had integrated into the gene were ratified by PCR and by Southern blot hybridizations (see below).

#### 8.3 Methods for in vitro analysis of DNA, RNA and protein

#### 8.3.1 Over-expression and purification of DddP protein.

100 ml cultures of *Escherichia coli* strain BL21 containing plasmid pBIO1658 were grown at 37 °C in LB broth containing 100 µg ampicillin ml<sup>-1</sup> to an OD<sub>600</sub> of 0.4–0.6. To induce *dddP* expression, 10 µM IPTG was added, followed by incubation at 25 °C until cells reached the stationary phase. Cells were pelleted by centrifugation and resuspended in 0.1 vols 20 mM Tris buffer (pH 8) at 4 °C. Cells were lysed by sonication (6×10 s) and cell debris was removed by centrifugation. Then, 25% (w/v) (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> was added to the lysate and the resulting precipitate was removed by centrifugation. The supernatant was loaded onto a phenyl sepharose high-performance column (xk16/20; GE Healthcare) equilibrated with 20 mM Tris containing 25% (w/v) (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>. Proteins were eluted using a 25–0% (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> gradient (flow rate 3 ml min<sup>-1</sup>). The 3 x 3 ml fractions that contained DddP polypeptide were then applied to a DEAE (HiTrap, 5 ml; GE Healthcare) column equilibrated with 20 mM Tris buffer, pH 8 (flow rate 5 ml min<sup>-1</sup>). Proteins were eluted in the same buffer with a linear gradient of 0–1 M NaCl. The 3 x 2 ml fractions containing DddP were concentrated to 2 ml using an Amicon Ultra 4 ml Centrifugal filter and loaded onto a Superdex 200 gel filtration column (10/300GL; GE Healthcare) equilibrated with 50 mM MES buffer, pH 6 (flow rate 1

ml min<sup>-1</sup>). The 2 x 5 ml DddP-containing fractions were pooled and stored at 4 °C. DddP concentration was estimated from  $A_{280}$  measurements, using  $\epsilon_{280}$  nm=76860 M<sup>-1</sup> cm<sup>-1</sup>, as calculated from the numbers of tryptophan, tyrosine and cysteine residues in the protein (Gill and von Hippel, 1989).

#### 8.3.2 Enzymic properties of DddP in vitro

To determine the pH optimum for DddP activity, a mixed buffer solutions (MBS) of 50 mM  $K_2$ HPO<sub>4</sub>, sodium citrate, Tris and *N*-cyclohexyl-2-aminoethanesulfonic acid was used to generate solutions in the pH range 2.0–9.0. HCl or NaOH were used to achieve the desired pH before assaying DMS production (section 8.5.2).

The effects of temperature on DddP activity were examined by incubating the reaction mixtures in 50 mM MES buffer, pH 6.0 over a range from 25 °C to 70 °C, at 5 °C intervals.

To examine the effects of metal availability on DddP function, the metal chelators 2,2'bipyridyl (2.5 mM) or EDTA (25 mM) were added to DddP in MES buffer pH 6 and incubated for 15 min at room temperature before adding DMSP substrate and assaying as in section 8.5.2.

 $K_{\rm m}$  and  $V_{\rm max}$  studies were done with ~0.3 µM DddP in MES buffer, pH 6, with DMSP concentrations ranging from 1 to 20 mM. Samples were incubated at 30 °C and DMS headspace measurements were taken at regular time intervals.

#### 8.3.3 Analytical ultracentrifugation

Analytical ultracentrifugation was done at 12000 r.p.m., 20 °C, in a Beckman Optima XL-I analytical ultracentrifuge, with absorbance optics and an An50Ti rotor. Partial specific

volumes were estimated from DddP amino acid sequences using SEDNTERP software, version 1.05 (Philo, 1997). Scans were recorded every 4 h to determine when protein samples had reached equilibrium, when five scans were recorded per sample. DddP ( $\sim$ 5  $\mu$ M) was in 20 mM Tris, 100 mM NaCl, pH 8. Data were analysed using Ultrascan (Demeler, 2005) and fitted to a one-component model.

#### 8.3.4 Southern blotting

#### 8.3.4.1 Probe design

To ratify pK19 insertional mutations, probes were designed for Southern blots that targeted a 5-10 kbp flanking region around the gene of interest, bordered by natural restriction sites. In some cases, probes were re-appropriated from existing PCR products, for example in the use of the *SPO193-1914* fragment originally cloned into pBluescript.

#### 8.3.4.2 Probe creation

This protocol used the DIG High Prime DNA labelling and detection starter kit (Roche). A 16  $\mu$ l aliquot of the probe DNA was boiled for 10 minutes to denature the DNA, followed by a quick cooling on a salt ice mixture. Then, 4  $\mu$ l of "dig1", DIG high prime, buffer was added, vortexed and incubated overnight at 37 °C for efficient random, primed labelling of the probe DNA. Following incubation, the probe was heat-killed at 65°C for 10 minutes and stored at - 20°C for use later.

#### 8.3.4.3 Southern blot buffers

The buffers prepared for this procedure are as follows:

Denaturation solution 0.5 M NaOH 1.5 M NaCl

Neutralisation solution 1 M Tris-HCl; pH8 1.5 M NaCl

## Depurination solution 0.2 M HCl

## 20X SSC (in 2 litres) 3 M NaCl 0.3 M Tri sodium citrate; pH7 To be diluted for use as 6X SSC

# Washing Buffer 0.1 M maleic acid 0.15 M NaCl; pH 7.5 0.3% (v/v) Tween 20

Maleic acid Buffer 0.1 M maleic acid 0.15 M NaCl; adjusted with solid NaOH to pH 7.5

Post Hybridisation Buffers (PHB) 1) 0.5x SSC, 0.1% SDS 2) 0.1x SSC, 0.1% SDS

Detection Buffer 0.1 M Tris-HCl 0.1 M NaCl; pH 9.5

#### **Blocking** solution

Dilute 10x blocking solution (vial 6 in the DIG kit) 1:10 in maleic acid buffer

Antibody solution

Centrifuge anti-digoxigenin-AP (vial 4 in the DIG kit) for 5 minutes at 13,000 rpm, and pipette from surface.

Dilute 1: 10,000 (75mU/ml) in blocking solution

#### 8.3.4.4 Southern blot protocol

Genomic DNA preparations were digested with the relevant restriction enzymes overnight at 37°C, and the fragments, separated by agarose gel electrophoresis were photographed, together with a ruler scale.

The gel was then rinsed in  $dH_2O$  and soaked in depurination solution for 15 minutes, then rinsed in  $dH_2O$  and soaked in denaturation solution for 30 minutes. It was rinsed again and finally soaked in neutralisation solution for 30 minutes. The gel was then set up for blotting as shown in figure 8.2 below, and left overnight.

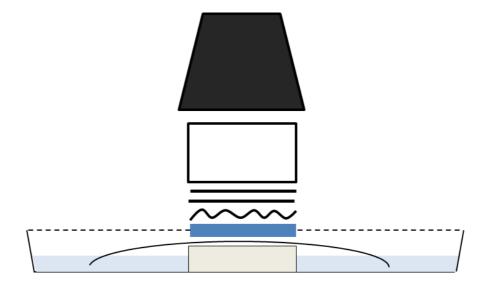


Figure 8.2 Cross section of a Southern blot hybridisation

Simplified representation of the Southern blot technique used in this study. A tray was partially filled with 6 x SSC blotting solution (light blue) and a filter paper "bridge" created (curved line) over a plastic base (light grey). Once soaked through, the agarose gel (dark blue) to be blotted was placed on top of the filter paper bridge and cling film (dashed lines) was used to seal the edges completely. This prevents the filter bridge from coming into contact with any of the other blotting components such as the blotting membrane and preventing efficient blotting. A Hybrid- N<sup>+</sup> nylon membrane (Amersham Biosciences), cut to identical dimensions as the agarose gel, was placed on top of the gel (waved line). Two similarly sized pieces of filter paper (straight line) were then placed on top of this, followed by a thick block of tissue (white rectangle). Finally, a heavy weight (black trapezium) was carefully placed on top and left overnight, to allow complete blotting to occur.

#### 8.3.4.5 Probe hybridisation

Following blotting, the nylon filter was removed and rinsed in 2X SSC solution, air-dried, and UV cross-linked at 254 nm. A 20 ml aliquot of DIG Easy Hyb solution (kit no. 7) was preheated to 42°C and added to the filter in a heat-sealed hybridisation bag (Amersham Biosciences) and incubated at 42°C for 30 minutes. The DNA probe was denatured at 100°C for 10 minutes, cooled in a beaker of salted ice for 2 minutes and 3  $\mu$ l was added to 3 ml of DIG Easy Hyb solution. This was added to the nylon filter in the bag, re-sealed and incubated at 42°C overnight.

#### 8.3.4.6 Post-hybridisation and detection

Following hybridisation, the filter was washed for  $2 \ge 5$  minutes in PHB 1 at room temperature, and  $2 \ge 15$  minutes in PHB 2 at  $68 \degree$ C.

The filter was then rinsed in washing buffer, and then incubated at room temperature for 30 minutes in blocking buffer. The DIG-Ap antibody mix was added to blocking buffer (4  $\mu$ l in 20 ml), and was then applied to the filter and incubated for 30 minutes at room temperature. The filter was then washed 2 x 15 minutes in washing buffer and 5 minutes in detection buffer, at room temperature.

A colour substrate solution was created by adding 200  $\mu$ l NBT/BCIP to 10 ml detection buffer; this was applied to the filter immediately, then stored in the dark. The filter was periodically assessed for development of bands.

#### 8.3.5 RNA extraction

Starter cultures of *R. pomeroyi* DSS-3 were grown overnight in  $\frac{1}{2}$  YTSS media, washed and diluted 1:200 in MBM minimal media containing 10 mM succinate, and incubated at 28°C for 16 h to an OD<sub>600</sub> nm ~ 0.4. Then, compounds whose effects on gene expression were to be

determined were added (see respective chapters). Then, cultures were incubated for a further 2 hours, before harvesting 2 x 50 ml aliquots of each growth condition. 20 ml of ice-cold 5% phenol, 95% ethanol (v/v) solution was added and the culture was incubated on ice for 1 hour to stabilize RNA and prevent degradation. Cells were then pelleted and RNA was extracted using SV Total RNA isolation kit and buffers (Promega). To achieve the quality and quantity of RNA required for the microarray analysis, an amended SV Total RNA protocol was used, as follows.

Pellets were re-suspended in 200 µl TE buffer containing 100 mg/ml lysozyme, and incubated at 37°C for 10 minutes. These were then freeze/thawed 5 times using dry ice. 135 µl lysis buffer (containing β-mercaptoethanol) and 630 µl RNA dilution buffer were added to the frozen pellet and mixed by inversion. The sample was then added to a 2ml screw cap vial containing 0.6 g Sigma ( $\leq 106 \,\mu$ M) acid-washed glass beads and broken open ("ribolysed") in a FastPrep (MP Bio) for 40 seconds at speed setting 6.0. Samples were then heated at 70°C for 3 minutes and centrifuged at maximum speed for 10 minutes. The supernatant was transferred to a clean microfuge tube containing 200 µl 100% ethanol and mixed by pipette, before being transferred to a spin column and centrifuged at max speed for 30 seconds. The eluate was discarded and the column washed with 600  $\mu$ l wash buffer before being spun again for 30 seconds. DNase mix was prepared in the following way: 5 µl 90 mM MnCl<sub>2</sub>, 40 µl DNase core buffer, 5 µl DNase. Then, 50 µl of the DNase mix was applied to the column matrix and incubated at 37°C for 40 minutes, before stopping with 200 µl DNase stop mix and centrifuged for 30 seconds at 13,000 rpm. Columns were washed with 600 µl wash buffer, and centrifuged at 13,000 rpm for 30 seconds. The eluate was discarded, and the column washed again, with 250 µl wash buffer and centrifuged at 13,000 rpm for 30 seconds. The eluate was discarded and the column spun again for 1 minute, to remove any residual wash buffer. Columns were transferred to clean microfuge tubes and 70 µl RNAse-free dH<sub>2</sub>O applied to the matrix. This was left to stand for 1 minute before being centrifuged at 4500 x g for 2 minutes and the column was discarded. The RNA sample was frozen at -80°C.

#### 8.3.5.1 RNA quality assessment

A Nanodrop was used on 1  $\mu$ l samples to estimate the DNA/RNA concentration (in ng/ $\mu$ l) and purity, using dH<sub>2</sub>O as a blank. Purity was estimated by the ratios of absorbance at 280 nm (DNA levels), 260 nm (RNA levels) or 230 nm (organic solvents) in the relevant sample.

An Experion automated electrophoresis platform (BioRad) was used to assess the integrity of the RNA for use in the microarray assay, as per the manufacturer's instructions for the Experion RNA StdSens kit and an RNA StdSens Chip (BioRad). Typically, 1  $\mu$ l of each sample was loaded into 1 of 11 wells on the StdSens chip. A virtual-gel output showed bands corresponding to 23S, 16S and 5S rRNA and the absence of any 'smearing' illustrated that the RNA had not degraded during the isolation process.

#### 8.3.6 Quantitative real-time RT-PCR

The absence of genomic DNA contamination in the RNA samples was confirmed by PCR amplification, using Taq PCR Master Mix (QIAGEN) according to manufacturer's specifications. For qRT-PCR, primers (Table 8.6) were designed using Primer3 software (Rozen and Skaletsky, 2000) and used to amplify genes from total RNA isolated from cells grown in the presence or absence of 5 mM DMSP, 2.5 mM acrylate or 5 mM DMS. The data were normalised against the RNA encoded by *SPO2904*, an *Ruegeria pomeroyi* gene that encodes a serine/threonine protein phosphatase/nucleotidase, and was used in the microarray protocol of Bürgmann *et al.* (2007). The iScript<sup>TM</sup> One-step RT-PCR Kit (Bio-Rad) with SYBR® Green was used for reverse transcription followed by PCR, as described in the manufacturer's manual. Master mix and RNA solutions were added to a final volume of 25  $\mu$ l, containing 50 ng of RNA, and quantification of mRNA transcripts was carried out using a CFX96 Real-time PCR Detection system (Bio-Rad).

#### 8.3.7 Microarrays

#### 8.3.7.1 Slide design

Agilent 8 x 15K gene expression microarray slides were used, and a GenePix Array List (GAL) file was created using eArray (https://earray.chem.agilent.com/earray/) based on the *Ruegeria pomeroyi* DSS-3 genome and megaplasmid sequences obtained from B Pearson (personal communication). Each gene was represented by 3 separate ~60 bp oligomer "probe" sequences, each designated to a particular "spot" on the array slide by eArray. Control spots were also assigned to the slide to allow calibration by the GenePix program.

#### 8.3.7.2 RNA Labelling

A Stratagene AffinityScript Multiple Temperature reverse transcriptase kit was used. RNA was extracted and assessed for quality and quantity, as above. A 10  $\mu$ g aliquot of RNA was dried using a rotary vacuum, then resuspended in 7.7  $\mu$ l of RNase-free water. The RNA labelling protocol was as follows.

Random priming reactions were set up with 7.7  $\mu$ l of RNA and 5  $\mu$ g (1.7  $\mu$ l) of random hexamers, and incubated at 70°C for 5 minutes, before being put on ice for 10 minutes. Samples were centrifuged briefly.

A reverse transcription (RT) reaction mix was prepared:

 $2 \ \mu l \ of \ 10 \ x \ RT \ buffer$ 

2 µl of 0.1 M Dithiothreitol (DTT)

 $0.6 \ \mu l$  of 50 x dNTPs (25 mM of dATP, dGTP, dTTP and 10 mM of dCTP, equivalent to 25  $\ \mu l$  of dA, dG, dT and 10  $\ \mu l$  of dCTP from Amersham Pharmacia 100 mM stock dNTP

2 µl of Cy3 or Cy5-dCTP (1 mM stock from Amersham Pharmacia)

4 µl of reverse transcriptase (AffinityScript, Stratagene)

Water, treated with 0.1% Diethylpyrocarbonate (DEPC) to a total volume of 20 µl.

Samples were mixed and incubated at 25°C for 10 minutes, incubated overnight at 42°C. The next day, 15  $\mu$ l of freshly made 0.1 M NaOH was added and the RNA hydrolysed at 70°C for 10 minutes. Then, 15  $\mu$ l of 0.1 M HCl was added to neutralise the reaction.

#### 8.3.7.3 Genomic DNA (gDNA) labelling reaction

A Gibco/BRL BioPrime DNA labelling kit was used. Firstly, a 2  $\mu$ g aliquot of chromosomal DNA from a QIAGEN Genomic DNA Kit was added and the volume brought to 21  $\mu$ l with DEPC-treated water. Then 20  $\mu$ l of 2.5 x Random primer/reaction buffer mix from the Gibco kit was added. The sample was boiled for 5 minutes, then put on ice for 5 minutes. Once on ice, 5  $\mu$ l of 10 x dNTP mix (1.2 mM each of dATP, dGTP, dTTP; 0.6 mM of dCTP; 10 mM Tris pH 8.0; 1 mM EDTA) plus 3  $\mu$ l of Cy3 or Cy5 dCTP (1 mM stock from Amersham) and 1  $\mu$ l of Klenow fragment from the kit was added. This was spun down at 13,000 rpm and incubated at 37°C overnight.

#### 8.3.7.4 RNA/gDNA Clean up

An Invitrogen PureLink PCR purification was used as follows. A 50  $\mu$ l aliquot of labelled RNA/gDNA reactions was added to 200  $\mu$ l of PureLink binding buffer B2 and mixed well by inversion. The sample was added to a spin column and centrifuged at 10 000X g in a microcentrifuge for 1 minute. The flow-through was discarded, and column washed with 650  $\mu$ l of Wash Buffer, then spun as before. The flow-through was discarded again, and the empty spin column was spun at full speed for 3 minutes to remove all traces of wash buffer. DNA was eluted using 50  $\mu$ L DEPC-treated water and spun after incubating for 1 minute at room temperature, elution was performed by centrifugation into a clean eppendorf for 2 minutes at 13,000 rpm.

#### 8.3.7.5 Array Hybridisation

An Agilent Hi-RPM gene expression hybridisation kit was used as follows. Samples for the conditions being compared in each array were dried as before, then resuspended in 10  $\mu$ l of DEPC-treated water, before being mixed. To this 20  $\mu$ l sample, 25  $\mu$ l of hybridisation buffer (Agilent) and 5  $\mu$ l of blocking reagent (Agilent) were added, and boiled for 3 minutes before being cooled to room temperature.

A 40  $\mu$ l aliquot of each sample was applied to a well on an Agilent hybridisation chamber, held in a clamp (Agilent). The array was carefully applied on top of the chamber, and clamped down firmly. This was incubated in a hybridisation oven at 65°C overnight.

#### 8.3.7.6 Washing the hybridised arrays

The hybridisation chamber and microarray, held in the clamp was disassembled in wash buffer 1 (Agilent) away from bright light. The array was loaded into a foil covered 50 ml Falcon tube filled with Wash Buffer 1 and rotated gently for 1 minute at room temperature. The array slide was transferred to another Falcon tube containing wash buffer 2, pre-warmed to 37°C and rotated again for 2 minutes. The slide was then immersed briefly in acetonitrile and air-dried, before being placed into stabilisation buffer, air-dried and stored in the dark.

#### 8.3.7.7 Scanning and normalisation

The array slide was scanned using a GenePix 4000B scanner, and data were acquired using GenePix 6.0 software (Axon Instruments). Laser excitation was provided by individual 635 nm and 532 nm lasers. These wavelengths correspond to the ideal wavelengths used to excite the fluorophores Cy3 and Cy5 (Amersham Pharmacia Biotech), respectively, used in this study. The intensity of each laser and the gain of the photomultiplier tube (detector) were manually calibrated, and an approximately equal fluorescence for both dyes was achieved, to

negate saturation of one wavelength. Spots were scanned at a resolution of 5 µm. Spots incorrectly aligned to the template GAL file were identified manually and corrected where possible. Data from the triplicate technical, and duplicate biological replicates were processed using GeneSpring GX software, and normalised using LOWESS and Marray packages, giving the fold change in fluorescence between conditions (Cleveland, 1979; Wang *et al.*, 2002; B Pearson, personal communication). Student's *T*-tests were used to calculate P values.

#### 8.4 Sequencing and bioinformatics

DNA sequencing was carried out at the John Innes Centre Genome Laboratory. Sequences were analysed with Artemis (http://www.sanger.ac.uk/Software/Artemis) and DNAStar (DNAStar Inc. Madison, WI) EditSeq and aligned using MegAlign and GeneDoc. Basic Local Alignment Search Tool analyses (BLAST) utilised the BLASTp (for proteins), BLASTn (for genes) or BLASTx (nucleotide into protein) function of the NCBI BLAST online program (www.ncbi.nlm.nih.gov/BLAST/). Genome alignment was carried out using the NMPDR database tool, the IMG database (Markowitz *et al.*, 2010) or PATRIC (Gillespie *et al.*, 2011). Operon prediction used the DOOR database (Mao *et al.*, 2008). Motif searches used MEME/MAST (Bailey *et al.*, 2009). PePPER was used to predict terminators (<u>http://pepper.molgenrug.nl/</u>), along with TransTermHP (Kingsford *et al.*, 2007).

#### 8.5 Gas chromatography

Gas chromatography used a flame photometric detector (GC 2010; Shimadzu, Milton Keynes, UK) and a 30 m x 0.53 mm ID-BP1 5.0 µm capillary column (SGE Europe, Milton Keynes, UK).

#### 8.5.1 Bacterial DMS assays

To measure DMS production, bacteria were first grown overnight in the appropriate medium. Cultures were then adjusted to the same  $OD_{600}$  nm, the cells were spun down and resuspended in complete medium containing a final concentration of 5 mM DMSP. Certain experiments involved washing three times in minimal media before re-suspending in minimal media. They were then incubated with shaking in 1ml vials (12 x 32 mm, Alltech Associates) at their appropriate growth temperature. After the necessary incubation time, DMS in the vial headspace was quantified by gas chromatography (see above). Activities are expressed as nmol DMS. min<sup>-1</sup>.  $\mu$ g protein<sup>-1</sup>. Protein concentrations were estimated using Bradford's assays (BioRad).

To measure the effects of pre-growth in the presence of potential co-inducer molecules (DMSP, acrylate, DMS) on the rates of DMS production, the relevant strains were grown in minimal medium supplemented with one of these compounds, or unamended. After overnight growth, the cells were washed, and were assayed for DMS production, using fresh DMSP substrate.

#### 8.5.2 Purified DddP DMS assays

For *in vitro* assays with purified DddP, the protein (~10  $\mu$ M) was added to an appropriate buffer solution containing different concentrations of the substrate DMSP in a sealed 1 ml vial (Alltech Associates). For routine assays, DMSP was used at a final concentration of 5 mM. Following incubation at 30 °C, the DMS in the headspace was quantified by gas chromatography as above.

#### **8.6** β-galactosidase assays

 $\beta$ -galactosidase activity was used to measure the transcription of *lacZ* reporter fusions, based on the reporter plasmid pBIO1878, similar to the methods described by Sambrook *et al.* (1989) and modified by Rossen *et al.* (1985).

Ruegeria pomeroyi or Rhizobium leguminosarum were inoculated from a single colony to 5 ml <sup>1</sup>/<sub>2</sub> YTSS, or TY medium, respectively, and grown for ~2 days at 28°C. 1 ml of this culture was inoculated to 100 ml MBM or TY and grown for 16 hours at 28°C (until OD<sub>600</sub> nm was between 0.3 and 0.6). Potential co-inducer molecules were added to 5 ml aliquots of R. pomeroyi or R. leguminosarum cultures in Sterilin 10 ml Universal containers and incubated at 28°C for a further 2 hours. Add 1 ml of culture to a 2 ml cuvette and measure the  $OD_{600}$  nm (blanked against MBM medium), to determine the cell density. Aliquots of the culture (0.1 - 0.5 ml depending on activity) were removed and added to a 2 ml microfuge tube and made up to 1 ml with Z buffer (see below). Then, 2 drops of chloroform plus 1 drop 0.1% SDS (w/v) was added to each microfuge tube and vortexed for 10 seconds, and the tubes incubated at  $28^{\circ}$ C for 5 minutes. 0.2 ml *O*-nitrophenyl- $\beta$ -D-galactopyranoside (ONPG) (4 mg ml<sup>-1</sup>) was added to each microfuge tube, a timer started and incubated at 28°C. When sufficient yellow colour had developed, 0.5 ml Na<sub>2</sub>CO<sub>3</sub> was added to stop the reaction and the time was recorded. The tubes were centrifuged for 3 minutes at 13,000 rpm to pellet the cell debris and 1 ml of the supernatant was added to a 2 ml cuvette, with the OD<sub>420 nm</sub> measured (blanked against Z buffer).

The  $\beta$ -galactosidase activity, in Miller units, was calculated using the following equation: Miller units =  $1000 \times OD_{420} / t \times V \times OD_{600}$ 

t = time

V = volume of culture used

Z-buffer contains 1 ml 3 M Na<sub>2</sub>HPO<sub>4</sub>.7H<sub>2</sub>O, 0.5 ml 4 M NaHPO<sub>4</sub>.7H<sub>2</sub>O, 0.5 ml 1 M KCl, 0.5 ml 0.1 M MgSO<sub>4</sub>.7H<sub>2</sub>O, 175  $\mu$ l mercaptoethanol, made up to a final volume of 50 ml with

 $dH_2O$ . ONPG was freshly prepared each time.  $OD_{600}$  nm and  $OD_{420}$  nm values were measured using a Unicam 8625 UV/VIS spectrophotometer.

#### 8.7 Detection of substrates, catabolites and metals

# 8.7.1 High Pressure Liquid Chromatography (HPLC) and Nuclear Magnetic Resonance (NMR)

#### 8.7.1.1 Detection of metabolites by HPLC

Labelled [1-<sup>14</sup>C]DMSP (2.7 kBq) was added to cell-free extracts of *E. coli* strain BL21 to a final concentration of 1 mM. Following 2 hours incubation, labelled products were identified by HPLC. The DMSP catabolites were resolved on a Dionex (Sunnyvale, CA, USA) ICE-AS6 column (250 mm x 9 mm id.) eluted isocratically with 0.4 mM HCl at a flow rate of 1 ml min<sup>-1</sup>. The column, suppressor and detection cell were maintained at 35°C. Metabolites were detected by tandem suppressed ion conductivity and UV detection at 210 nm. Ion suppression used a Dionex AMMS-ICE Micromembrane Suppressor with 5 mM tetrabutylammonium hydroxide as the solvent (flow rate of 2 ml min<sup>-1</sup>). Conductivity was measured with a Dionex ED50 conductivity detector and UV absorbance with a Jasco (Tokyo, Japan) UV-975 detector.

#### 8.7.1.2 Detection of metabolites by NMR

Cultures of *E. coli* containing cloned genes were grown overnight at 37°C in LB, adjusted to equivalent  $OD_{600}$  nm values, then diluted  $10^{-2}$  into 1 ml M9 [made up in deuterium oxide (> 99.9%)], containing glycerol (10 mM), and 10 mM [1-<sup>13</sup>C]DMSP or 5 mM [1-<sup>13</sup>C]acrylate and 0.2 mM IPTG to induce expression of the cloned genes. After incubating at 28 °C overnight, cells were lysed by adding perchloric acid (5% v/v final concentration) and samples were

incubated on ice for 10 min. Cell debris was spun down and the supernatant was added to NMR tubes.

For NMR of purified DddP, 50  $\mu$ l (~0.9  $\mu$ M) of protein was added to 5 mM [1-<sup>13</sup>C]DMSP and made up to 500  $\mu$ l in deuterium oxide.

Proton-decoupled <sup>13</sup>C NMR spectra were measured at 75 MHz with a Varian Gemini 2000 in  $D_2O$ . An average of 300 scans was recorded when DMSP was used and an average of 1000 scans when acrylate solutions were used. Note that <sup>13</sup>C NMR detects the <sup>13</sup>C isotope, whose natural abundance is only 1.1%.

#### 8.7.2 Inductively coupled plasma, optical emission spectrometry (ICP-OES).

ICP-OES was carried out on samples of 10  $\mu$ M DddP in 2.5% (v/v) nitric acid. A Varian Vista Pro CCD simultaneous ICP-OES, with axial torch, concentric seaspray nebuliser (Glass Expansion) and 50 ml cyclonic spray chamber was used to analyse triplicate samples. The power was 1.2 kW and the analysed wavelengths were: Co, 228.62 nm; Cu, 324.75 nm; Mn, 259.37 nm; Ni, 230.30 nm; Zn, 213.86 nm. The limits of detection for Co, Cu, Mn, Ni and Zn were 7.3, 5.3, 9.1, 206 and 10.0 nmol, respectively.

#### **8.8 Chemical syntheses**

These were carried out by Dr Y Chan (UEA chemistry department). DMSP was prepared by adding dimethylsulfide (15.3 ml, 0.21 mol) to aqueous HCl (100 ml, 2 M). Acrylic acid (10.0 g, 0.14 mol) was added, and the resulting mixture heated under reflux (95°C) for 2h. The reaction mixture was allowed to reach room temperature and concentrated under reduced pressure. The residue was triturated using a mixture of ethanol/diethyl ether, and the resulting solid was filtered, washed once with ethanol and twice with diethyl ether to yield DMSP (21.8 g, 92%). The identity and purity of DMSP was established by melting point, and by infrared and NMR spectroscopy [m.p. 134–135°C;  $v_{max}$  (solid/cm<sup>-1</sup>) 3013, 2621, 2549, 2478, 2426,

1787, 1691, 1414, 1396, 1247, 1183, 906;  $\delta_{\rm H}$  (400 MHz; D<sub>2</sub>O) 2.94 (6H, s, 2 × CH<sub>3</sub>), 2.98 (2H, t, J = 6.8 Hz), 3.53 (2H, t, J = 6.8 Hz);  $\delta_{\rm C}$  (75 MHz; D<sub>2</sub>O) 24.51, 27.84, 38.05, 173.17]. [1-<sup>14</sup>C]DMSP was made as described above except that <sup>14</sup>C-labelled acrylic acid (27 MBq in 0.75 ml dH<sub>2</sub>O) was also added, and the resulting mixture heated under reflux (95°C) for 2 hours. The reaction mixture was allowed to reach room temperature and concentrated under reduced pressure. The residue was triturated using a mixture of ethanol/diethyl ether, and the resulting solid was filtered, washed once with ethanol and twice with diethyl ether to yield [1-<sup>14</sup>C]DMSP. [1-<sup>13</sup>C]DMSP HCl was prepared by adding <sup>13</sup>C-acrylic acid (100 mg, 1.39 mmol) to aqueous HCl (5 ml, 2 M). DMS (2.78 mmol) was added and then treated as above, for [1-<sup>14</sup>C]DMSP, to yield [1-<sup>13</sup>C]DMSP (210 mg, 89%).

#### Table 8.4 Strains used in this study

Bacteria	Characteristics	Source
Escherichia coli 803	Met; used as host for transformation with	Wood (1966)
	large plasmids	
E. coli BL21	Used as host for expression from pET21a	Studier and Moffat
		(1986)
E. coli JM101	Used as host for expression of pBluescript and	(Yanisch-Perron et
	for blue white screens	al., 1985)
Rhizobium leguminosarum	Wild type; Strep <sup>R</sup> mutant	Young <i>et al.</i> (2006)
J391		
<i>Rhizobium leguminosarum</i> strain 3841	Wild type strain (Strep <sup>R</sup> ) Heterologous host for expression of $dddW$	Todd et al., (2011b)
Roseovarius nubinhibens	Wild type isolate	González et al.
ISM		(2003)
Ruegeria pomeroyi DSS-3	Wild type isolate	González et al.
		(2003)
Ruegeria pomeroyi J470	<i>R. pomeroyi</i> DSS-3 Rif <sup>R</sup> mutant	Todd <i>et al.</i> , (2011a)
Ruegeria pomeroyi J471	R. pomeroyi J470 with insertion in dmdA	This study; Todd et
		al., (2012)
Ruegeria pomeroyi J527	R. pomeroyi J470 with insertion in acul	This study; Todd et
		al., (2012)
Ruegeria pomeroyi J530	R. pomeroyi J470 with insertion in dmdR	This study; Todd et
		al., (2012)
Ruegeria pomeroyi J497	R. pomeroyi J470 with insertion in dddW	This study

#### 2012

Plasmid	Description	Features	Origin
pET21a	Used for expression of cloned <i>ddd</i> , <i>dmdA</i> and	Amp <sup>R</sup>	Novagen
	acuI genes		
pBIO1658	R. nubinhibens dddP cloned in pET21a	Amp <sup>R</sup>	Todd et al.,
			(2010)
pMP220	Broad-host-range promoter probe vector with	Tet <sup>R</sup>	Spaink et al.,
	promoterless <i>lacZ</i> reporter gene		(1987)
pBIO1878	Spc <sup>R</sup> cassette cloned into pMP220.	Tet <sup>R</sup>	Todd et al.,
	Used as <i>lacZ</i> -promoter fusion plasmid in	Spec <sup>R</sup>	(2012a)
	Ruegeria pomeroyi		
pRK2013	Used as mobilising plasmid in tri-parental	Kan <sup>R</sup>	Figurski &
	crosses		Helinski
			(1979)
pBluescript	Used for sub-cloning genes, prior to cloning	Amp <sup>R</sup>	Stratagene
	into larger, low-copy vectors		
pK19mob	Suicide insertion plasmid	Kan <sup>R</sup>	Schäfer et al.,
			(1994)
pBIO1879	pK19mob suicide vector with added Spec <sup>R</sup>	Spec <sup>R</sup>	Todd et al.,
	cassette	Kan <sup>R</sup>	(2011)
pBIO1887	<i>dddP</i> of <i>Ruegeria pomeroyi</i> DSS-3 cloned in	Amp <sup>R</sup>	This work
	pET21a		
pUC18	High copy number cloning vector	Amp <sup>R</sup>	Vieira et al.,
			(1982)
pBIO1945	pBIO1878 containing 307 bp fragment	Contains	This study
	spanning the <i>dddW</i> promoter	<i>lacZ</i> reporter	
		Tet <sup>R</sup> , Spec <sup>R</sup>	

## Table 8.5 Plasmids used in this study

M. Kirkwood Chapter 8: Materials and met		Interials and methods20	
pBIO1946	pOT2 containing <i>SPO0454</i> and its native promoter	Gent <sup>R</sup>	This study
pBIO1947	SPO0454 promoter cloned into pBIO1878	Amp <sup>R</sup> Spec <sup>R</sup>	This study
pBIO1948	pET21 containing R. pomeroyi dddW	Amp <sup>R</sup>	This study
pBIO1949	pBIO1879 containing internal <i>dddW</i> fragment	Spec <sup>R</sup> Kan <sup>R</sup>	Todd <i>et al.</i> , (2011)
pBIO1870	Internal fragment of <i>Ruegeria pomeroyi</i> DSS- 3 <i>dmdA</i> cloned into pBIO1879	Spec <sup>R</sup> Kan <sup>R</sup>	Todd <i>et al.</i> , (2012b)
pBIO2013	<i>Ruegeria pomeroyi</i> DSS-3 <i>acuI</i> cloned in pET21a	Amp <sup>R</sup>	Todd <i>et al.</i> , (2012b)
pBIO2019	<i>Ruegeria pomeroyi</i> DSS-3 <i>dmdA</i> and <i>acuI</i> with their native promoter, cloned in pBluescript	Amp <sup>R</sup>	Todd <i>et al.</i> , (2012b)
pBIO2020	Ruegeria pomeroyi DSS-3 dmdA-lacZ reporter fusion plasmid	Tet <sup>R</sup> Spec <sup>R</sup>	Todd <i>et al.</i> , (2012b)
pBIO2021	<i>Ruegeria pomeroyi</i> DSS-3 <i>acuI-lacZ</i> reporter fusion plasmid	Tet <sup>R</sup> Spec <sup>R</sup>	Todd <i>et al.</i> , (2012b)
pBIO2022	<i>Ruegeria pomeroyi</i> DSS-3 <i>dmdA</i> and <i>acuI</i> with their native promoter cloned in pBIO1878	Tet <sup>R</sup> Spec <sup>R</sup>	Todd <i>et al.</i> , (2012b)
pBIO2023	pBIO2019 with a deletion in <i>dmdA</i>	Amp <sup>R</sup>	Todd <i>et al.</i> , (2012b)
pBIO2024	<i>Ruegeria pomeroyi</i> DSS-3 <i>acuI</i> with its own promoter, sub-cloned from pBIO2023 into pBIO1878	Tet <sup>R</sup> Spec <sup>R</sup>	Todd <i>et al.</i> , (2012b)
pBIO2025	Internal fragment of <i>Ruegeria pomeroyi</i> DSS- 3 <i>acuI</i> cloned into pBIO1879	Spec <sup>R</sup> Kan <sup>R</sup>	Todd <i>et al.</i> , (2012b)

M. Kirkwoo	d Chapter 8: Materials and methods	2012	
pBIO2083	pBIO2021 digested with <i>EcoRI</i> to remove	Tet <sup>R</sup>	This study
	dmdA promoter and a large section of dmdA	Spec <sup>R</sup>	
pBIO2084	pBIO2021 with an introduced +1 frameshift	Tet <sup>R</sup>	This study
	prior to <i>lacZ</i>	Spec <sup>R</sup>	
pBIO2085	pBIO1879 containing internal <i>dmdR</i> fragment	Spec <sup>R</sup>	This study
		Kan <sup>R</sup>	
pBIO2086	pBIO1878 with <i>coxM</i> promoter	Spec <sup>R</sup>	This study
		Tet <sup>R</sup>	

## Table 8.6 Primers used in this study

Primer name	Primer sequence (5' – 3')	Use
DddPF	G <u>GAATTC</u> CATATGAACCAGCA	Cloning <i>dddP</i> of <i>Roseovarius</i>
	TTACAGCG	nubinhibens ISM into
		pET21a
DddPR	G <u>GAATTC</u> ATGCCCCGCCCTGC	Cloning <i>dddP</i> of <i>Roseovarius</i>
	CCG	nubinhibens ISM into
		pET21a
SydDddPF	GGACCG/AACTCCGCTGGCGT	Amplification of conserved
	Т	regions of <i>dddP</i> in isolates of
		Aspergillus sydowii
SydDddPR	TCATAG/ACCCGTCTCCGTCAC	Amplification of conserved
		regions of <i>dddP</i> in isolates of
		Aspergillus sydowii
D295 For	CATCTCTTTCGcCACCGATCTC	Site directed mutagenesis of
	ATCGG	Roseovarius nubinhibens
		ISM dddP

M. Kirkwood	Chapter 8: Materials and methods	
D295 Rev	CCGATGAGATCGGTGgCGAAA	Site directed mutagenesis of
	GAGATG	Roseovarius nubinhibens
		ISM dddP
D297 For	CTTTCGACACCGcTCTCATCG	Site directed mutagenesis of
	GCAGC	Roseovarius nubinhibens
		ISM dddP
D297 Rev	GCTGCCGATGAGAgCGGTGTC	Site directed mutagenesis of
	GAAAG	Roseovarius nubinhibens
		ISM dddP
D307 For	GGCATCTGCGTCGcCATCTCG	Site directed mutagenesis of
	CGCAGC	Roseovarius nubinhibens
		ISM dddP
D307 Rev	GCTGCGCGAGATGgCGACGCA	Site directed mutagenesis of
	GATGCC	Roseovarius nubinhibens
		ISM dddP
H371 For	GGCTGCCTGATGGcTGGGGGTC	Site directed mutagenesis of
	GGGC	Roseovarius nubinhibens
		ISM dddP
H371 Rev	GCCCGACCCCAgCCATCAGGC	Site directed mutagenesis of
	AGCC	Roseovarius nubinhibens
		ISM dddP
E406 For	GCTCTGTGTCGcGGCGGCGGT	Site directed mutagenesis of
	CGGCG	Roseovarius nubinhibens
		ISM dddP
E406 Rev	CGCCGACCGCCGCCgCGACAC	Site directed mutagenesis of
	AGAGC	Roseovarius nubinhibens
		ISM dddP

M. Kirkwood	Chapter 8: Materials and metho	ods 2012
E421 For CCATCAAGCTCGcGGATCAGC		Site directed mutagenesis of
	TGC	Roseovarius nubinhibens
		ISM dddP
E421 Rev	GCACCTGATCCgCGAGCTTGA	Site directed mutagenesis of
	TGG	Roseovarius nubinhibens
		ISM dddP
Wprom 1	GC <u>GAATTC</u> CATCGTCAGCAGA	Cloning promoter of
	GTC	Ruegeria pomeroyi DSS-3
		dddW into pBIO1878
Wprom 2	GC <u>CTGCAG</u> CACCATGTCGCGCG	Cloning promoter of
	GCG	Ruegeria pomeroyi DSS-3
		dddW into pBIO1878
Wpet1	AACTGCAG <u>CATATG</u> ACCGCCAT	Amplification of <i>dddW</i> to
	GCT	create expression vector
	CGACAGTTTC	pBIO1948
Wpet2	AT <u>GGATCC</u> TCAGGCGCTGGCG	Amplification of <i>dddW</i> to
	GTGAACCG	create expression vector
		pBIO1948
RTdddPF	GCTGTGGAACACCCATAA	For qRT-PCR analysis of
		dddP
RTdddPR	GCCTCGGTCGAAATAGAACA	For qRT-PCR analysis of
		dddP
RTdddQF	AAACCTTCTGGCCGAGTTTC	For qRT-PCR analysis of
		dddQ
RTdddQR	ATAGGCTGTGGTCGTCAGGT	For qRT-PCR analysis of
		dddQ
RTdddWF	GTTTCGCAACCGATCTGACT	For qRT-PCR analysis of
		dddW

M. Kirkwood	Chapter 8: Materials and metho	ods 2012
RTdddWR	TCGAGGCCCAGATAGAACTC	For qRT-PCR analysis of <i>dddW</i>
SPO2904F	TAAGCTTTCCGGTCCTGATG	
SP02904F	TAGETTICCONCENTRATO	For qRT-PCR analysis,
SP02004P		control
SPO2904R	ACCGTCGATTTCAGCAACTT	For qRT-PCR analysis,
		control
Wmut1	CG <u>GGATCC</u> AGCCCGGCAACCT	Cloning internal <i>dddW</i>
	GCCG	fragment to create
		pBIO1949
Wmut2	CG <u>GGATCC</u> ATAGGCAAAGCG	Cloning internal <i>dddW</i>
	CAGACC	fragment to create
		pBIO1949
454P1	AT <u>CTGCAG</u> CAAACCGCGCTATT	Cloning R. pomeroyi DSS-3
	TGTGACT	SPO0454 and its promoter
		into pOT2
454P2	AT <u>GTCGAC</u> AGATCGGTTGCGAA	Cloning R. pomeroyi DSS-3
	ACTGTCG	SPO0454 and its promoter
		into pOT2
454prom1	GGCC <u>GAATTC</u> GGCGATGCCCAC	Cloning promoter of
		SPO0454 into pBIO1878
454prom2	AACTGCAGGCGCACCAGCGCG	Cloning promoter of
	CC	SPO0454 into pBIO1878
SPO1913F_Pst	GC <u>CTGCAG</u> GGGCCCGACGCGCT	For cloning fragment used in
	GCGG	pBIO1870
SPO1913R_Eco	GCGACATTCAGCCGAATTC	For cloning fragment used in
		pBIO1870
SPO1913/14_XbaF	GCG <u>TCTAGA</u> GGTCCTGACGCCG	For cloning fragment used
	GGTCGCAC	in pBIO2019

M. Kirkwood	Chapter 8: Materials and metho	ods 2012
SPO1913/14_BamR	CG <u>GGATCC</u> GGGCCTCTTGCCGC	For cloning fragment used in
	TCACTTC	pBIO2019 and pBIO2013
acuI+1_NsiR	ACCA <u>ATGCAT</u> TCGAACATCTG	For cloning fragment used in
		pBIO2084
Spo1914_NdeF	GGAATC <u>CATATG</u> TTCAATGCAT	For cloning fragment used in
	TGGTGG	pBIO2013
1912_EcoF	CGCG <u>GAATTC</u> CAGATCGACCCC	For cloning fragment used in
	AACAGC	pBIO2085
1912_PstR	CGCG <u>CTGCAG</u> TCGTATAGCGC	For cloning fragment used in
	ATCAGTTCG	pBIO2085
SPO2399_ecoR1	GC <u>GAATTC</u> CGGCTTTTGCAGG	For cloning fragment used in
for1	TGCC	pBIO2086
SPO2399_pst1	GC <u>CTGCAG</u> CGCGGCTTCGCGC	For cloning fragment used in
rev1	AGG	pBIO2086

Sequences of the oligonucleotide primers are shown, with cloning restriction sites underlined.

All primers were synthesised by MWG Biotech. Where required, primers were supplied with phosphate groups at their 5' ends.

# **Chapter 9**

Bibliography

Alavi, M., Miller, T., Erlandson, K., Schneider, R. and Belas, R. (2001) Bacterial community associated with *Pfiesteria*-like dinoflagellate cultures. *Environ Microbiol* **3**: 380–396

Alker, A.P. Smith, G.W. and Kim, K. (2001) Characterization of *Aspergillus sydowii* (Thom et Church), a fungal pathogen of Caribbean sea fan corals. *Hydrobiologia* **460**: 105-111

Andreae, M.O. (1990) Ocean-atmosphere interactions in the global biogeochemical sulfur cycle. *Mar Chem* **30:** 1-29

Ansede, J.H., Pellechia, P.J. and Yoch, D.C. (1999) Metabolism of acrylate to ßhydroxypropionate and its role in dimethylsulfoniopropionate lyase induction by a salt marsh sediment bacterium, *Alcaligenes faecalis*. *Appl Environ Microbiol* **65**: 5075-5081

Archer, S.D., Widdicombe, C.E., Tarran, G.A., Rees, A.P. and Burkill, P.H. (2001) Production and turnover of particulate dimethylsulphoniopropionate during a coccolithophore bloom in the northern North Sea. *Aquat Microb Ecol* **24**: 225–41

Auld D.S. (1995) Removal and replacement of metal ions in metallopeptidases. *Methods Enzymol* **248**: 228-242

Bacic, M.K., Newell, S.Y. and Yoch, D.C. (1998) Release of dimethylsulfide from dimethylsulfoniopropionate by plant-associated salt marsh Fungi. *Appl Env Microbiol* **64**: 1484-1489

Bacic, M.K. and Yoch, D.C. (1998) *In vivo* characterization of dimethylsulfoniopropionate lyase in the Fungus *Fusarium lateritium*. *Appl Environ Microbiol* **64:** 106-111

Bailey, T.L., Bodén, M., Buske, F.A., Frith, M., Grant, C.E., Clementi, L., Ren, J., Li, W.W. and Noble, W.S. (2009) MEME SUITE: tools for motif discovery and searching. *Nucleic Acids Res* **37**: W202-W208

Barber, R.D., Rott, M.A. and Donohue, T.J. (1996) Characterization of a glutathionedependent formaldehyde dehydrogenase from *Rhodobacter sphaeroides*. *J Bacteriol* **178**: 1386-1393 Baumann, P. and Baumann, L. (1981) The marine Gram-negative Eubacteria: genera *Photobacterium, Beneckea, Alteromonas, Pseudomonas*, and *Alcaligenes*. In: Starr, M.P., Stolp, H., Trüper, H.G., Balows, A. and Schlegel, H.G. (Eds.). The Prokaryotes, Springer-Verlag, Berlin, 1302–1331

Bazan, J. F., Weaver, L. H., Roderick, S. L., Huber, R. and Matthews, B.W. (1994) Sequence and structure comparison suggest that methionine aminopeptidase, prolidase aminopeptidase P and creatinase share a common fold. *Proc Nat Acad Sci USA* **91**: 2473-2477

Benz, R. and Bauer, K. (1988) Permeation of hydrophilic molecules through the outer membrane of Gram-negative bacteria. Review on bacterial porins. *Eur J Biochem* **176:** 1–19

Berg, I.A., Kockelkorn, D., Buckel, W. and Fuchs, G. (2007) A 3-hydroxypropionate/4hydroxybutyrate autotrophic carbon dioxide assimilation pathway in Archaea. *Science* **318**: 1782–1786

Beringer, J.E. (1974) R factor transfer in *Rhizobium leguminosarum. J Gen Microbiol* 84: 188-198

Beringer, J.E. and Hopwood, D.A. (1976) Chromosomal recombination and mapping in *Rhizobium leguminosarum. Nature* **264:** 291-293

Biers, E.J., Sun, S. and Howard, E.C. (2009) Prokaryotic genomes and diversity in surface ocean waters: interrogating the global ocean sampling metagenome. *Appl Environ Microbiol* **75:** 2221–29

Birnboim, H.C. and Doly, J. (1979) A rapid alkaline extraction procedure for screening recombinant plasmid DNA. *Nucleic Acids Res* **7:** 1513-1523

Bjellqvist, B., Hughes, G.J., Pasquali, C., Paquet, N., Ravier, F., Sanchez, J.C., Frutiger, S. and Hochstrasser, D.F. (1993) The focusing positions of polypeptides in immobilized pH gradients can be predicted from their amino acid sequences. *Electrophoresis* **14**: 1023-1031

Brinkmeyer, R., Knittel, K., Jürgens, J., Weyland, H., Amann, R. and Helmke, E. (2003) Diversity and structure of bacterial communities in Arctic versus Antarctic pack ice. *Appl Environ Microbiol* **69**: 6610–6619

Broadbent, A.D., Jones G.B. and Jones, R.J. (2002) DMSP in corals and Benthic algae from the Great Barrier Reef. *Estuar Coast Shelf S* **55**: 547-555

Brown, T.A. (2002) Transcriptomes and Proteomes. In: Genomes, John Wiley and Sons, Oxford, 69-93

Buchan, A., González, J.M. and Moran, M.A. (2005) Overview of the marine Roseobacter lineage. *Appl Environ Microbiol* **71:** 5665–5677

Bürgmann, H., Howard, E.C., Ye, W., Sun, F., Sun, S., Napierala, S. and Moran, M.A. (2007)
Transcriptional response of *Silicibacter pomeroyi* DSS-3 to dimethylsulfoniopropionate
(DMSP). *Environ Microbiol* 9: 2742–2755

Cairney, J., Booth, I.R. and Higgins, C.F. (1985) Osmoregulation of gene expression in *Salmonella typhimurium*: proU encodes an osmotically induced betaine transport system. *J Bacteriol* **164**: 1224–1232

Cantoni, G. L. and Anderson, D. G. (1956) Enzymatic cleavage of dimethylpropiothetin by *Polysiphonia lanosa. J Biol Chem* **222:** 171-177

Caspi, R., Foerster, H., Fulcher, C.A., Kaipa, P., Krummenacker, M., Latendresse, M., Paley, S., Rhee, S.Y., Shearer, A.G., Tissier, C., Walk, T.C., Zhang, P. and Karp, P.D. (2012) The MetaCyc database of metabolic pathways and enzymes and the BioCyc collection of pathway/genome databases. *Nucleic Acids Res* **36**: D623-631

Charlson, R.J., Lovelock, J.E., Andreae, M.O. and Warren, S.G. (1987) Oceanic phytoplankton, atmospheric sulphur, cloud albedo and climate. *Nature* **326**: 655-661

Choudhary, M., Zanhua, X, Fu, Y.X. and Kaplan, S. (2007) Genome analyses of three strains of *Rhodobacter sphaeroides*: evidence of rapid evolution of chromosome II. *J Bacteriol* **189**: 1914-1921

Chung, C.H. and Goldberg, A.L. (1981) The product of the *lon* (*capR*) gene in *Escherichia coli* is the ATP-dependent protease, protease La. *P Natl Acad Sci USA* **78:**4931-4935

Cleveland, W.S. (1979) Robust locally weighted regression and smoothing scatterplots. *J Am Stat Assoc* **74:** 829-836

Cole, J.L., Lary, J.W., Moody, T.P. and Laue, T.M. (2008) Analytical Ultracentrifugation: Sedimentation Velocity and Sedimentation Equilibrium. In: Correia, J.J. and Detrich, H.W. (Eds.) Methods in Cell Biology, Academic Press, 143-179

Cooney, R. P., Pantos, O., Tissier, M.D.A.L., Barer, M.R., O'Donnell, A.G. and Bythell. J.C. (2002) Characterization of the bacterial consortium associated with black band disease in coral using molecular microbiological techniques. *Environ Microbiol* **4**: 401–413

Cooper, R.M. and Rankin, B. (1978) Cell wall-degrading enzymes of vascular wilt fungi. II. Properties and modes of action of polysaccharidases of *Verticillium albo-atrum* and *Fusarium oxysporum* f.sp. *lycopersici*. *Physiol Plant Pathol* **13**: 101-134

Cosquer, A., Pichereau, V., Pocard, J.A., Minet, J., Cormier, M. and Bernard, T. (1999) Nanomolar levels of dimethylsulfoniopropionate, dimethylsulfonioacetate, and glycine betaine are sufficient to confer osmoprotection to *Escherichia coli*. *Appl Environ Microbiol* **65:** 3304–3311

Cowan, J.M., Urbanowski, M.L., Talmi, M. and Stauffer, G.V. (1993) Regulation of the *Salmonella typhimurium metF* gene by the MetR protein. *J Bacteriol* **175:** 5862-5866

Cunliffe, M. (2010) Correlating carbon monoxide oxidation with *cox* genes in the abundant marine Roseobacter clade. *ISME J* **5**: 685-691

Cunningham, G.B., Strauss, V. and Ryan, P.G. (2008) African penguins (*Spheniscus demersus*) can detect dimethyl sulfide, a prey-related odour. *J Exp Biol* **211**:3123-3127

Curson, A.R.J., Rogers, R., Todd, J.D., Brearley, C.A. and Johnston, A.W.B. (2008) Molecular genetic analysis of a dimethylsulfoniopropionate lyase that liberates the climatechanging gas dimethylsulfide in several marine α-proteobacteria and *Rhodobacter sphaeroides*. *Environ Microbiol* **10**: 757-767

Curson, A. R. J., Sullivan, M. J., Todd, J. D. and Johnston, A.W.B. (2010) Identification of genes for dimethyl sulfide production in bacteria in the gut of Atlantic herring (*Clupea harengus*). *ISME J* **4**: 144-146

Curson, A.R.J., Todd, J.D., Sullivan, M.J. and Johnston, A.W.B. (2011a) Catabolism of dimethylsulfoniopropionate: microorganisms, enzymes and genes. *Nat Rev Microbiol* **9:** 849-859

Curson, A.R.J., Fowler, E.K., Dickens, S., Johnston, A.W.B. and Todd, J.D. (2011b) Multiple DMSP lyases in the γ-proteobacterium *Oceanimonas doudoroffii*. *Biogeochem*. 1-11 DOI: 10.1007/s10533-011-9663-2

Curson, A.R.J., Sullivan, M.J., Todd, J.D. and Johnston, A.W.B. (2011c) DddY, a periplasmic dimethylsulfoniopropionate lyase found in taxonomically diverse species of Proteobacteria. *ISME J.* **5**: 1191-1200

Demeler, B. (2005) UltraScan – a comprehensive data analysis software package for analytical ultracentrifugation experiments. In: Scott, D.J., Harding, S.E. and Rowe, A.J. (Eds.). Analytical Ultracentrifugation: Techniques and Methods, Royal Society of Chemistry, Cambridge, 210–229

DeRisi, J.L., Iyer, V.R. and Brown, P.O. (1997) Exploring the metabolic and genetic control of gene expression on a genomic scale. *Science* **278**: 680–686

de Souza, M.P. and Yoch, D.C. (1995a) Purification and characterization of dimethylsulfonio-propionate lyase from an *Alcaligenes*-like dimethyl sulfide-producing marine isolate. *Appl Environ Microbiol* **61:** 21–26

de Souza, M.P. and Yoch, D.C. (1995b) Comparative physiology of dimethyl sulfide production by dimethylsulfoniopropionate lyase in *Pseudomonas doudoroffii* and *Alcaligenes* sp. strain M3A. *Appl Environ Microbiol* **61**: 3986–3991 de Souza M.P. and Yoch, D.C. (1996) N-terminal amino acid sequences and comparison of DMSP lyases from *Pseudomonas doudoroffii* and *Alcaligenes* strain M3A. In: Kiene, R.P., Visscher, P.T., Keller, M.D. and Kirst, G.O. (Eds.). Environmental and biological chemistry on dimethylsulfoniopropionate and related sulfonium compounds, Plenum Press, New York, 293–304

Dorrell, N., Ahmed, A.H. and Moss, S.H. (1993) Photoreactivation in a *phrB* mutant of *Escherichia coli* K-12: evidence for the role of a second protein in photorepair. *Photochem Photobiol* 58: 831-835

Dudoit, S. and Yang, Y.H. (2002) Bioconductor R packages for exploratory analysis and normalization of cDNA microarray data. In: Parmigiani, G., Garrett, E.S., Irizarry, R.A. and Zeger, S.L. (Eds.). The Analysis of Gene Expression Data: Methods and Software. Springer, New York

Dunwell, J.M., (1998) Cupins: a new superfamily of functionally diverse proteins that include germins and plant seed storage proteins. *Biotechnol Genet Eng* **15**: 1–32

Dunwell, J.M. and Gane, P.J. (1998) Microbial relatives of seed storage proteins: conservation of motifs in a functionally diverse superfamily of enzymes. *J Mol Evol* **46:** 147– 154

Dunwell, J.M., Khuri, S. and Gane, P.J. (2000) Microbial relatives of the seed storage proteins of higher plants: conservation of structure and diversification of function during the evolution of the cupin superfamily. *Microbiol Mol Biol R* **64**: 153–179

Dunwell, J.M., Culham, A., Carter, C.E., Sosa-Aguirre, C.R. and Goodenough, P.W. (2001) Evolution of functional diversity in the cupin superfamily. *Trends Biochem Sci* **26**: 740–745

Dunwell, J.M., Purvis, A. and Khuri, S. (2004) Cupins: the most functionally diverse protein superfamily? *Phytochem* **65:** 7-17

Edwards, D.M., Reed, R.H. and Stewart, W.D.P. (1988) Osmoacclimation in *Enteromorpha intestinalis*: long-term effects of osmotic stress on organic solute accumulation. *Mar Biol* **88**: 457-476

Eichler, K., Bourgis, F., Buchet, A., Kleber, H.P. and Mandrand-Berthelot, M.A. (1994) Molecular characterization of the *cai* operon necessary for carnitine metabolism in *Escherichia coli. Mol Microbiol* **13**: 775–786

Eilers, H., Pernthaler, J., Glöckner, F.O. and Amann, R. (2000) Culturability and *in situ* abundance of pelagic bacteria from the North Sea. *Appl Environ Microbiol* **66**: 3044–3051

Eitinger, T., Rodionov, D.A., Grote, M. and Schneider, E. (2011) Canonical and ECF-type ATP-binding cassette importers in prokaryotes: diversity in modular organization and cellular functions. *FEMS Microbiol Rev* **35**: 3–67

Elssner, T., Engemann, C., Baumgart, K. and Kleber, H.P. (2001) Involvement of coenzyme A esters and two new enzymes, an enoyl-CoA hydratase and a CoA-transferase, in the hydration of crotonobetaine to 1-carnitine by *Escherichia coli*. *Biochem* **40**: 11140–11148

Ferguson, S.J., Jackson, J.B. and McEwan, A. G. (1987) Anaerobic respiration in the Rhodospirillaceae: characterisation of pathways and evaluation of roles in redox balancing during photosynthesis. *FEMS Microbiol. Rev* **46**: 117–143

Figurski, D. H. and Helinski, D. R. (1979) Replication of an origin-containing derivative of plasmid RK2 dependent on a plasmid function provided *in trans. P Natl Acad. Sci USA* **76**: 1648–1652

Franklin, D.J., Steinke, M., Young, J., Probert, I. and Malin, G. (2010)Dimethylsulphoniopropionate (DMSP), DMSP-lyase activity (DLA) and dimethylsulphide(DMS) in 10 species of cocclithophore. *Mar Ecol Prog Ser* **410**: 13-23

Fuhrmann, S., Ferner, M., Jeffke, T., Henne, A., Gottschalk, G. and Meyer, O. (2003) Complete nucleotide sequence of the circular megaplasmid pHCG3 of *Oligotropha carboxidovorans*: function in the chemolithoautotrophic utilization of CO, H<sub>2</sub> and CO<sub>2</sub>.*Gene* **32**: 67–75

Fujita, Y. and Fujita, T. (1987) The gluconate operon *gnt* of *Bacillus subtilis* encodes its own transcriptional negative regulator. *P Natl Acad Sci USA* 84: 4524-4528

Gage, D.A., Rhodes, D., Nolte, K.D., Hicks, W.A., Leustek, T., Cooper, A.J.L. and Hanson, A.D. (1997) A new route for synthesis of dimethylsulphoniopropionate in marine algae. *Nature* **387**: 891-894

Galinski, E.A. (1995) Osmoadaptation in Bacteria. Adv Microb Physiol 37: 273-328

Galperin, M.Y., Gaidenko, T.A., Mulkidianian, A.Y., Nakano, M. and Price, C.W. (2001) MHYT, a new integral membrane sensor domain. *FEMS Microbiol Lett* **27:** 17-23

Galperin, M.Y. (2008) Telling bacteria: do not LytTR. Structure 16: 657-659

Gao, R., Mack, T.R. and Stock, A.M. (2007) Bacterial response regulators: versatile regulatory strategies from common domains. *Trends Biochem Sci* **32**: 225–234

Geiser, D.M., Taylor, J.W., Ritchie, K.B. and Smith GW (1998) Cause of sea fan death in the West Indies. *Nature* **394:** 137–138

Gill, S. C. and von Hippel, P. H. (1989) Calculation of protein extinction coefficients from amino acid sequence data. *Anal Biochem* **182**: 319–326

Gillespie, J.J., Wattam, A.R., Cammer, S.A., Gabbard, J., Shukla, M.P., Dalay, O., Driscoll, T., Hix, D., Mane, S.P., Mao, C., Nordberg, E.K., Scott, M., Schulman, J.R., Snyder, E.E., Sullivan, D.E., Wang, C., Warren, A., Williams, K.P., Xue, T., Yoo, H.S., Zhang, C., Zhang, Y., Will, R., Kenyon, R.W. and Sobral, B.W. (2011) PATRIC: The comprehensive bacterial bioinformatics resource with a focus on human pathogenic species. *Infect Immun* **79**: 4286-4298

Giovannoni, S.J., Britschqi, T.B., Moyer, C.L. and Field, K.G. (1990) Genetic diversity in Sargasso Sea bacterioplankton. *Nature* **345:** 60-63

Giovannoni, S.J., Tripp, H.J., Givan, S., Podar, M., Vergin, K.L., Baptista, D., Bibbs, L., Eads, J., Richardson, T.H., Noordewier, M., Rappe, M.S., Short, J.M., Carrington, J.C. and Mathur, E.J. (2005) Genome streamlining in a cosmopolitan oceanic bacterium. *Science* **309**: 1242-1245

González, J. M., Whitman, W. B., Hodson, R. E. and Moran, M. A. (1996) Identifying numerically abundant culturable bacteria from complex communities: an example from a lignin enrichment culture. *Appl Environ Microbiol* **62**: 4433–4440

González, J.M. and Moran, M.A. (1997) Numerical dominance of a group of marine bacteria in the alpha-subclass of the class Proteobacteria in coastal seawater. *Appl Environ Microbiol* **63**: 4237–4242

González, J.M., Mayer, F., Moran, M.A., Hodson, R.E. and Whitman, W.B. (1997) *Sagittula stellata* gen. nov., sp. nov., a lignin-transforming bacterium from a coastal environment. *Int J Syst Bacteriol* **47**: 773–780

González, J.M., Kiene, R.P. and Moran, M.A. (1999) Transformation of sulfur compounds by an abundant lineage of marine bacteria in the α-subclass of the class Proteobacteria. *Appl Environ Microbiol* **65**: 3810–3819

González, J.M., Simó,R., Massana, R., Covert, J.S., Casamayor, E.O., Pedrós-Alió, C. and Moran, M.A. (2000) Bacterial community structure associated with a dimethylsulfoniopropionate producing North Atlantic algal bloom. *Appl Environ Microbiol* **66:** 4237– 46

González, J. M., Covert, J. S., Whitman, W. B., Henriksen, J. B., Mayer, F., Scharf, B., Schmitt R., Buchan, A., Fuhrman, J. A., Kiene, R. P. and Moran M. A. (2003) *Silicibacter pomeroyi* sp. nov. and *Roseovarius nubinhibens* sp. nov., dimethylsulfoniopropionatedemethylating bacteria from marine environments. *Int J Syst Evol Microbiol* **53**: 1261–1269

Graham, S.C., Bond, C.S., Freeman, H.C. and Guss, J.M. (2005) Structural and functional implications of metal ion selection in aminopeptidase P, a metalloprotease with a dinuclear metal center. *Biochem* **44**: 13820–13836

Gruber, A., Vugrinec, S., Hempel, F., Gould, S.B., Maier, U-G. And Kroth, P.G. (2007) Protein targeting into complex diatom plastids: functional characterisation of a specific targeting motif. *Plant Mol Biol* **64**: 519-530 Haardt, M., Kempf, B., Faatz, E. and Bremer, E. (1995) The osmoprotectant proline betaine is a major substrate for the binding-protein-dependent transport system ProU of *Escherichia coli* K-12. *Mol Gen Genet* **246**: 783–786

Hanson, A.D., Rivoal, J., Paquet, L. and Gage, D.A. (1994a) Biosynthesis of 3dimethylsulfoniopropionate in *Wollastonia biflora* (L.) DC. (Evidence that *S*methylmethionine is an intermediate). *Plant Physiol* **105**: 103-110

Hanson, A.D., Rathinasabapathi, B., Rivoal, J., Burnet, M., Dillon, M.O. and Gage, D.A. (1994b) Osmoprotective compounds in the Plumbaginaceae: a natural experiment in metabolic engineering of stress tolerance. *P Natl Acad Sci USA* **91**:306–310

Hatakeyama, S., Okuda, M. and Akimoto, H. (1982) Formation of sulfur-dioxide and methanesulfonic-acid in the photo-oxidation of dimethylsulfide in the air. *Geophys Res Lett* 9: 583–586

Hernández, C.T., Montero, A.Z., González, A.B., Rodríguez, J.A., Sabat, A.M. and Bayman, P. (2008) Fungi in healthy and diseased sea fans (*Gorgonia ventalina*): is *Aspergillus sydowii* always the pathogen? *Coral Reefs* **27**: 707-714

Herrmann, G., Selmer, T., Jessen, H. J., Gokarn, R. R., Selifonova, O., Gort, S. J. and Buckel,
W. (2005) Two beta-alanyl-CoA: ammonia lyases in *Clostridium propionicum*. *FEBS J*272: 813–821

Hofstee, B.H.J. (1973) Hydrophobic affinity chromatography of proteins. *Anal Biochem* **52:** 430–448

Howard, E.C., Henriksen, J.R., Buchan, A., Reisch, C.R., Bürgmann, H., Welsh, R., Ye, W., González, J.M., Mace, K., Joye, S.B., Kiene, R.P., Whitman, W.B. and Moran, M.A. (2006) Bacterial taxa that limit sulfur flux from the ocean. *Science* **314**: 649-652

Howard, E.C., Sun, S., Biers, E.J. and Moran M.A. (2008) Abundant and diverse bacteria involved in DMSP degradation in marine surface waters. *Environ Microbiol* **10**: 2397–2410

Howard, E.C., Sun, S., Reisch, C.R., Del Valle, D.A., Bürgmann, H., Kiene, R.P. and Moran, M.A. (2011) Changes in dimethyl-sulfoniopropionate demethylase gene assemblages in response to an induced phytoplankton bloom. *Appl Environ Microbiol* **77:**524–31

Hügler, M., Krieger, R.S., Jahn, M. and Fuchs, G. (2003) Characterization of acetyl-CoA/propionyl-CoA carboxylase in *Metallosphaera sedula*. *European J Biochem* **270**: 736– 744

James, F., Paquet, L., Sparace, S.A., Gage, D.A. and Hanson, A.D. (1995) Evidence implicating dimethylsulfoniopropionaldehyde as an intermediate in dimethylsulfoniopropionate biosynthesis. *Plant Physiol* **108**: 1439-1448

Johnson, A., Moran, M. and Miller, W. (2007) Investigating carbon monoxide (CO) consumption in the marine bacteria *Silicibacter pomeroyi* with *coxL* gene expression. *Geophys Res Abs* **9:** 4535

Johnston, A.W.B., Beynon, J.L., Buchanan-Wollaston, A.V., Setchell, S.M., Hirsch, P.R. and Beringer, J.E. (1978) High frequency transfer of nodulating ability between strains and species of *Rhizobium*. *Nature* **276**: 634-636

Johnston, A.W.B., Todd, J.D., Sun, L., Nikolaidou-Katsaridou, M.N., Curson, A.R.J. and Rogers, R. (2008) Molecular diversity of bacterial production of the climate changing gas, dimethyl sulphide, a molecule that impinges on local and global symbioses. *J Exp Bot* **59**: 1059-1067

Jones, G.B. and Trevena, A.J. (2005) The influence of coral reefs on atmospheric dimethylsulphide over the Great Barrier Reef, Coral Sea, Gulf of Papua and Solomon and Bismarck Seas. *Mar Freshwater Res* **56**: 85-93

Karsten, U., Kirst, G.O. and Wiencke, C. (1992) Dimethylsulphoniopropionate (DMSP) accumulation in green macroalgae from polar to temperate regions: interactive effects of light versus salinity and light versus temperature. *Polar Biol* **12**: 603–607

Karsten,U., Kuck,K., Vogt,C. and Kirst, G.O. (1996) Dimethylsulphoniopropionate production in phototrophic organisms and its physiological function as a cryoprotectant. In:

Kersters, K., Devos, P., Gillis, M., Swings, J., Vandamme, P. and Stackebrandt, E. (2006) Introduction to the Proteobacteria. In Dworkin, M., Falkow, S., Rosenberg, E., Schleifer, K.H. and Stackebrandt, E. (Eds.). The Prokaryotes: A Handbook on the Biology of Bacteria, Springer, New York, 3-37

Kettle, A.J., Andreae, M.O., Amouroux, D., Andreae, T.W., Bates, T.S., Berresheim, H., Bingemer, H., Boniforti, R., Curran, M.A.J., DiTullio, G.R., Helas, G., Jones, G.B., Keller, M.D., Kiene, R.P., Leck, C., Levasseur, M., Malin, G., Maspero, M., Matrai, P., McTaggart, A.R., Mihalopoulos, N., Nguyen, B.C., Novo, A., Pustaud, J.P., Rapsomanikis, S., Roberts, G., Schebeske, G., Sharma, S., Simó, R., Staubes, R., Turner, S. and Uher, G. (1999) A global database of sea surface dimethylsulfide (DMS) measurements and a procedure to predict sea surface DMS as a function of latitude, longitude, and month. *Glob Biogeochem Cycles* **13**: 399–444

Kiene, R.P. and Taylor, B.F. (1988) Demethylation of dimethylsulfoniopropionate and production of thiols in anoxic marine sediments. *Appl Environ Microbiol* **54**: 2208-2212

Kiene, R. P. and Service, S. K. (1991) Decomposition of dissolved DMSP and DMS in estuarine waters: dependence on temperature and substrate concentration *Mar Ecol Prog Ser* 76: 1-11

Kiene, R. P. (1996) Production of methanethiol from dimethylsulfoniopropionate in marine surface waters. *Mar Chem* **54:** 69–83

Kiene, R.P., Linn, L.J., González, J., Moran, M.A. and Bruton, J.A. (1999) Dimethylsulfoniopropionate and methanethiol are important precursors of methionine and proteinsulfur in marine bacterioplankton. *Appl Environ Microbiol* **65:** 4549-4558

Kiene, R.P., Linn, L.J., and Bruton, J.A. (2000) New and important roles for DMSP in marine microbial communities. *J Sea Res* **43**: 209-224

King, J.M. (2003) Molecular and culture-based analyses of aerobic carbon monoxide oxidizer diversity. *Appl Environ Microbiol* **69(12):** 7257-7265

Kingsford, C.L., Ayanbule, K. and Salzberg, S.L. (2007) Rapid, accurate, computational discovery of Rho-independent transcription terminators illuminates their relationship to DNA uptake. *Genome Biol* **8**: R22

Kirkwood, M., Le Brun, N.E., Todd, J.D. and Johnston, A.W.B. (2010a) The *dddP* gene of *Roseovarius nubinhibens* encodes a novel lyase that cleaves dimethylsulfoniopropionate into acrylate plus dimethyl sulfide. *Microbiol* **156**: 1900-1906

Kirkwood, M., Todd, J.D., Rypien, K.L. and Johnston, A.W.B. (2010b) The opportunistic coral pathogen *Aspergillus sydowii* contains *dddP* and makes dimethyl sulphide from dimethylsulfoniopropionate. *ISME J* **4**: 147-150

Kirst, G.O., Thiel, C., Wolff, H., Nothnagel, J., Wanzek, M. and Ulmke, R. (1991)Dimethylsulfoniopropionate (DMSP) in ice algae and its possible biological role. *Mar Chem*35: 381-388

Kitaguchi, H., Uchida, A. and Ishida, Y. (1999) Purification and characterization of Lmethionine decarboxylase from *Crypthecodinium cohnii*. *Fisheries Sci* **65**: 613-617

Kocsis, M.G., Nolte, K.D., Rhodes, D., Shen, T-L., Gage, D.A. and Hanson, A.D. (1998) Dimethylsulfoniopropionate biosynthesis in *Spartina alterniflora*: evidence that *S*methylmethionine and dimethylsulfoniopropylamine are intermediates. *Plant Physiol* **117**: 273-281

Kocsis, M.G. and Hanson, A.D. (2000) Biochemical Evidence for Two Novel Enzymes in the Biosynthesis of 3-Dimethylsulfoniopropionate in *Spartina alterniflora*. *Plant Physiol* **123**: 1153-1162

Krell, T., Molina-Henares, A.J. and Ramos, J.L. (2006) The IclR family of transcriptional activators and repressors can be defined by a single profile. *Protein Sci* **15**: 1207-1213

Labrenz, M., Collins, M.D., Lawson, P.A., Tindall, B.J., Braker, G. and Hirsch, P. (1998) *Antarctobacter heliothermus* gen. nov., sp. nov., a budding bacterium from hypersaline and heliothermal Ekho Lake. *Int J Syst Bacteriol* **48**: 1363–1372

Lathe, G.H. and Ruthven, C.R. (1955) The separation of substances on the basis of their molecular weights, using columns of starch and water. *Biochem J* **60**: xxxiv

Levine, N. M., Varaljay, V. A., Toole, D. A., Dacey, J. W. H., Doney, S. C. and Moran, M. A. (2012) Environmental, biochemical and genetic drivers of DMSP degradation and DMS production in the Sargasso Sea. *Environ Microbiol* **14**: 1210–1223

Lyon, B.R., Lee, P.A., Bennett, J.M., DiTullio, G.R. and Janech, M.G. (2011) Proteomic analysis of a sea-ice Diatom: salinity acclimation provides new insight in the dimethylsulfoniopropionate production pathway. *Plant Physiol* **157**: 1926-1941

Maddocks S.E. and Oyston P.C.F. (2008) Structure and function of the LysR-type transcriptional regulator (LTTR) family proteins. *Microbiol* **154:** 3609–3623

Malmstrom, R.R., Kiene, R.P. and Kirchman, D.L. (2004a) Identification and enumeration of bacteria assimilating dimethylsulfoniopropionate (DMSP) in the North Atlantic and Gulf of Mexico. *Limnol Oceanogr* **49**: 597–606

Malmstrom, R.R., Kiene, R.P., Cottrell, M.T. and Kirchman D.L. (2004b) Contribution of SAR11 bacteria to dissolved dimethylsulfoniopropionate and amino acid uptake in the North Atlantic Ocean. *Appl Environ Microbiol* **70**: 4129-4135

Manukhov, I.V., Mamaeva, D.V., Rastorguev, S.M., Faleev, N.G., Morozova, E.A., Demidkina, T.V., and Zavilgelsky, G.B. (2005) A gene encoding 1-methionine γ-lyase is present in enterobacteriaceae family genomes: identification and characterization of *Citrobacter freundii* 1-methionine γ-lyase. *J Bacteriol* **187**: 3889–3893

Mao, F., Dam, P., Chou, J., Olman, V. and Xu, Y. (2008) DOOR: a database for prokaryotic operons. *Nucleic Acids Res* **37:** 459-463

Margaritis, T., Lijnzaad, P., van Leenen, D., Bouwmeester, D., Kemmeren, P., van Hooff, S.R. and Holstege, F.C.P. (2009) Adaptable gene-specific dye bias correction for two-channel DNA microarrays. *Mol Syst Biol* **5**: 266

Markowitz, V.M., Chen, I.A., Palaniappan, K., Chu, K., Szeto, E., Grechkin, Y., Ratner, A., Anderson, I., Lykidis, A., Mavromatis, K., Ivanova, N.N. and Kyrpides, N.C. (2010) The Integrated Microbial Genomes system: an expanding comparative analysis resource. *Nucleic Acids Res* **38**: Database Issue

Matrai, P.A. and Keller, M.D. (1994) Total organic sulfur and dimethylsulfoniopropionate (DMSP) in marine phytoplankton: intracellular variations. *Mar Biol* **119**: 61–68

Matthews, R.G., Sheppard, C. and Goulding, C. (1998) Methylenetetrahydrofolate reductase and methionine synthase: biochemistry and molecular biology. *Eur J Pediatr* **157**: S54–59

Matthews, R.G., Koutmoas, M. and Datta, S. (2008) Cobalamin-dependent and cobamidedependent methyltransferases. *Curr Opin Struct Biol* **18:** 658-666

McCarthy, A.A., Baker, H.M., Shewry, S.C., Patchett, M.L. and Baker, E.N. (2001) Crystal structure of methylmalonyl-coenzyme A epimerase from *P. shermanii*. *Structure* **9**: 637-646

McClung, C. R., Patriquin, D.G. and Davis, R.E. (1983) *Campylobacter nitrofigilis* sp. nov., a nitrogen-fixing bacterium associated with roots of *Spartina alterniflora* Loisel. *Int J Syst Bacteriol* **33:** 605-612

Merzouk. A., Levasseur, M., Scarratt, M., Michaud, S., Lizotte, M., Rivkin, R.B. and Kiene, R.P. (2008) Bacterial DMSP metabolism during the senescence of the spring diatom bloom in the Northwest Atlantic. *Mar Ecol Prog Ser* **369**: 1-11

Meyer, O., Frunzke, K., Gadkari, D., Jacobitz, S., Hugendieck, I. and Kraut, M. (1990) Utilization of carbon monoxide by aerobes: recent advances. *FEMS Microbiol Rev* 87: 253–260 Meyer, O., Frunzke, K. and Mörsdorf, G. (1993) Biochemistry of the aerobic utilization of carbon monoxide. In: Murrell, J.C. and Kelly, D.P. (eds.). Microbial Growth on C1 compounds. Andover, Mass.: Intercept, Ltd., 433–459

Mopper, K. and Taylor, B.F. (1986) Biogeochemical cycling of sulfur-thiols in coastal marine sediments. In: Sohn, M. (Ed.). Organic marine geochemistry, American Chemical Society, Washington, D.C. 324-339

Moran, M.A., González, J.M. and Kiene, R.P. (2003) Linking a bacterial taxon to sulfur cycling in the sea: studies of the marine Roseobacter group. *Geomicrobiol. J* **20:** 375–88

Moran, M.A., Buchan, A., González, J.M., Heidelberg, J.F., Whitman, W.B., Kiene, R.P., Henriksen, J.R., King, G.M., Belas, R., Fuqua, C., Brinkac, L., Lewis, M., Johri, S., Weaver, B., Pai, G., Eisen, J.A., Rahe, E., Sheldon, W.M., Ye, W., Miller, T.R., Carlton, J., Rasko, D.A., Paulsen, I.T., Ren, Q., Daugherty, S.C., Deboy, R.T., Dodson, R.J., Durkin, A.S., Madupu, R., Nelson, W.C., Sullivan, S.A., Rosovitz, M.J., Haft, D.H., Selengut, J. And Ward, N. (2004) Genome sequence of *Silicibacter pomeroyi* reveals adaptations to the marine environment. *Nature* **432:** 910–13

Moran, M.A. and Miller, W.L. (2007) Resourceful heterotrophs make the most of light in the coastal ocean. *Nat Rev Microbiol* **5**: 792–800

Morris, R.M., Rappé, M.S., Connon, S.A., Vergin, K.L., Siebold, W.A., Carlson, C.A. and Giovannoni, S.J. (2002) SAR11 clade dominates ocean surface bacterioplankton communities. *Nature* **420**: 806–10

Nei, M., Xu, P. and Glazko, G. (2001) Estimation of divergence times from multiprotein sequences for a few mammalian species and several distantly related organisms. *P Natl Acad Sci USA* **98**: 2497–2502

Nelson, D.M., Tréguer, P., Brzezinski, M.A., Leynaert, A. and Quéguiner, B. (1995) Production and dissolution of biogenic silica in the ocean. Revised global estimates, comparison with regional data and relationship to biogenic sedimentation. *Global Biogeochem Cycles* **9:** 359–372 Nevitt, G.A. and Bonadonna, F. (2005) Sensitivity to dimethyl sulphide suggests a mechanism for olfactory navigation by seabirds. *Bio Lett* **1:** 303–305

Nevitt, G.A. (2008) Sensory ecology on the high seas: the odor world of the procellariiform seabirds. *J Exp Biol* **211**: 1706-1713

Nevitt, G.A. (2011) The neuroecology of dimethyl sulfide: a global-climate regulator turned marine infochemical. *Integr Comp Biol* **51**: 819-25

Newton, R.J., Griffin, L.E., Bowles, K.M., Meile, C., Gifford, S., Givens, C.E., Howard,
E.C., King, E., Oakley, C.A., Reisch, C.R., Rinta-Kanto, J.M., Sharma, S., Sun, S., Varaljay,
V., Vila-Costa, M., Westrich, J.R. and Moran, M.A. (2010) Genome characteristics of a generalist marine bacterial lineage. *ISME J* 4:784–98

Nielsen, C.B., Friedman, B., Birren, B., Burge, C.B. and Galagan, J.E. (2004) Patterns of intron gain and loss in fungi. *PLoS Biol* **2:** 2234–2242

Noordkamp, D.J.B., Gieskes, W.W.C., Gottschal, J.C., Forney, L.J. and van Rijssel, M.
(2000) Acrylate in *Phaeocystis* colonies does not affect the surrounding bacteria. *J Sea Res*43: 287-296

Oh, J.I. and Kaplan, S. (2001) Generalized approach to the regulation and integration of gene expression. *Mol Microbiol* **39:** 1116-23

Otte, M.L. and Morris, J.T. (1994) Dimethylsulphoniopropionate (DMSP) in *Spartina alterniflora* Loisel. *Aquatic Bot* **48:** 239–259

Otte, M.L., Wilson, G., Morris, J.T. and Moran, B.M. (2004) Dimethylsulphoniopropionate (DMSP) and related compounds in higher plants. *J Exp Bot* 55: 1919-1925

Padmanabhan, B., Paehler, A. and Horikoshi, M. (2002) Structure of creatine amidinohydrolase from *Actinobacillus*. *Acta Crystallogr D Biol Crystallogr* **58:** 1322–1328

Patel, S.S. and Walt, D.R. (1987) Substrate specificity of acetyl coenzyme A synthetase. *J Biol Chem* **262**: 7132-7134

Pati, A., Gronow, S., Lapidus, A., Copeland, A., Del Rio, T.G., Nolan, M., Lucas, S., Tice,
H., Cheng, J., Han, C., Chertkov, O., Bruce, D., Tapia, R., Goodwin, L., Pitluck, S., Liolios,
K., Ivanova, N., Mavromatis, K., Chen, A., Palaniappan, K., Land, M., Hauser, L., Chang,
Y., Jeffries, C.D., Detter, J.C., Rohde, M., Göker, M., Bristow, J., Eisen, J.A., Markowitz, V.,
Hugenholtz, P., Klenk, H. and Kyrpides, N.C. (2010) Complete genome sequence of *Arcobacter nitrofigilis* type strain (CIT). *Stand Genomic Sci* 2: 300–308

Paul, L., Ferguson Jr., D.J. and Kryzycki, J.A. (2000) The trimethylamine methyltransferase gene and multiple dimethylamine methyltransferase gens of *Methanosarcina barkeri* contain in-frame and read-through amber codons. *J Bacteriol* **182**: 2520-2529

Pelzmann, A., Ferner, M., Gnida, M., Meyer-Klaucke, W., Maisel, T. and Meyer, O. (2009) The CoxD protein of *Oligotropha carboxidovorans* is a predicted AAA<sup>+</sup> ATPase chaperone involved in the biogenesis of the CO dehydrogenase [CuSMoO<sub>2</sub>] cluster. *J Biol Chem* **284**: 9578–9586

Persson, B., Hedlund, J. and Jörnvall, H. (2008) The MDR superfamily. *Cell Mol Life Sci* **65:** 3879-3894

Petersen, T.N., Brunak, S., von Heijne, G. and Nielsen, H. (2011) SignalP 4.0: discriminating signal peptides from transmembrane regions. *Nature Methods* **8:** 785-786

Prentki, P. and Krisch, H.M. (1984) *In vitro* insertional mutagenesis with a selectable DNA fragment. *Gene* **29:** 303-313

Quinn, P.K. and Bates, T.S. (2011) The case against climate regulation via oceanic phytoplankton sulphur emissions. *Nature* **480**: 51-6

Raina, J.B., Tapiolas, D., Willis, B.L. and Bourne, D.G. (2009) Coral-associated bacteria and their role in the biogeochemical cycling of sulfur. *Appl Environ Microbiol* **75**: 3492-3501

Raina J.B., Dinsdale, E. A., Willis, B. L. and Bourne, D. G. (2010) Do the organic sulfur compounds DMSP and DMS drive coral microbial associations? *Trends Microbiol* **18**: 101-108

Ramette, A., LiPuma, J. and Tiedje, J.M. (2005) Species abundance and diversity of *Burkholderia cepacia* complex in the environment. *Appl Environ Microbiol* **71**: 1193-1201

Rangarajan, E.S., Li, Y., Iannuzzi, P., Cygler, M. and Matte, A. (2005) Crystal structure of *Escherichia coli* crotonobetainyl-CoA: carnitine CoA-transferase (CaiB) and its complexes with CoA and carnitinyl-CoA. *Biochem* **44**: 5728–5738

Rawlings, N.D., Barrett, A.J. and Bateman, A. (2010) MEROPS: the peptidase database. *Nucleic Acids Res* **38**: D227-233

Reisch, C. R., Moran, M. A. and Whitman, W. B. (2008) Dimethylsulfoniopropionatedependent demethylase (DmdA) from *Pelagibacter ubique* and *Silicibacter pomeroyi*. *J Bacteriol* **190**: 8018-8024

Reisch, C.R., Noran, M.A. and Whitman, W.B. (2011a) Bacterial catabolism of dimethylsulfoniopropionate (DMSP). *Front Microbiol* **2:** 172 doi: 10.3389/fmicb.2011.00172

Reisch, C.R., Stoudemayer, M.J., Varaljay, V.A., Amster, I.J., Moran, M.A. and Whitman, W.B. (2011b) Novel pathway for assimilation of dimethylsulfoniopropionate widespread in marine bacteria *Nature* 473: 208–11

Rossen, L., Johnston, A.W.B. and Downie, J.A. (1985) DNA sequence of the *Rhizobium leguminosarum* nodulation genes *nodAB* and *C* required for root hair curling. *Nucleic Acids Res* 12: 9497-9508

Rozen, S., and Skaletsky, H. (2000) Primer3 on the WWW for general users and for biologist programmers. *Methods Mol Biol* **132**: 365-386

Rusch, D.B., Halpern, A.L., Sutton, G., Heidelberg, K.B., Williamson, S., Yooseph, S., Wu,
D., Eisen, J.A., Hoffman, J.M., Remington, K., Beeson, K., Tran, B., Smith, H., BadenTillson, H., Stewart, C., Thorpe, J., Freeman, J., Andrews-Pfannkoch, C., Venter, J.E., Li, K.,
Kravitz, S., Heidelberg, J.F., Utterback, T., Rogers, Y.H., Falcón, L.I., Souza, V., BonillaRosso, G., Eguiarte, L.E., Karl, D.M., Sathyendranath, S., Platt, T., Bermingham, E.,
Gallardo, V., Tamayo-Castillo, G., Ferrari, M.R., Strausberg, R.L., Nealson, K., Friedman,

R., Frazier, M. and Venter J.C. (2007) The *Sorcerer II* global ocean sampling expedition: Northwest Atlantic through Eastern Tropical Pacific. *PLoS Biol* **5**: 398–431

Rypien, K.L., Andras, J.P. and Harvell, C.D. (2008) Globally panmictic population structure in the opportunistic fungal pathogen *Aspergillus sydowii*. *Mol Ecol* **17**: 4068-4078

Saier, M.H.Jr., Yen, M.R., Noto, K., Tamang, D.G. and Elkan, C. (2009) The Transporter Classification Database: recent advances. *Nucleic Acids Res* **37**: D274-8

Sambrook, J., Fritsch, E.F. and Maniatis, T. (1989) Molecular cloning: A Laboratory Manual. Cold Spring Harbor, N.Y.; Cold Spring Harbor Laboratory

Santiago, B. and Meyer, O. (1997) Purification and molecular characterization of the H<sub>2</sub> uptake membrane-bound hydrogenase from the carboxidotrophic bacterium *Oligotropha carboxidovorans*. *J Bacteriol* **179**: 6053–6060

Santiago, B., Schübel, U., Egelseer, C. and Meyer, O. (1999) Sequence analysis, characterization and CO-specific transcription of the *cox* gene cluster on the megaplasmid pHCG3 of *Oligotropha carboxidovorans*. *Gene* **236**: 115-124

Schäfer, A., Tauch, A., Jäger, W., Jörn, K., Thierbach, G. and Pühler, A. (1994) Small mobilizable multi-purpose cloning vectors derived from the *Escherichia coli* plasmids pK18 and pK19: selection of defined deletions in the chromosome of *Corynebacterium glutamicum*. *Gene* **145**: 69-73

Schiffmann, R., Neugebauer, A., and Klein, C.D. (2006) Metal-mediated inhibition of *Escherichia coli* methionine aminopeptidase: structure-activity relationships and development of a novel scoring function for metal–ligand interactions. *J Med Chem* **49:** 511–522

Schneider, K., Asao, M., Carter, M.S. and Alber, B.E. (2012) *Rhodobacter sphaeroides* uses a reductive route *via* propionyl coenzyme A to assimilate 3-hydroxypropionate. *J Bacteriol* **194:** 225-232 Schübert, H.L., Blumenthal, R.M. and Cheng, X. (2003) Many paths to methyltransfer: a chronicle of convergence *Trends Biochem Sci* **28**: 329–335

Schuller, D.J., Reisch, C.R., Moran, M.A., Whitman, W.B. and Lanzilotta, W.N. (2012) Structures of dimethylsulfoniopropionate-dependent demethylase from the marine organism *Pelagabacter ubique. Prot Sci* **21**: 289-298

Shah, I. M. and Wolf Jr., R. E. (2006) Sequence requirements for Lon-dependent degradation of the *Escherichia coli* transcription activator SoxS: identification of the SoxS residues critical to proteolysis and specific inhibition of *in vitro* degradation by a peptide comprised of the N-terminal 21 amino acid residues. *J Mol Biol* **357**: 718-731

Shalon, D., Smith, S.J. and Brown P.O. (1996) A DNA microarray system for analyzing complex DNA samples using two-color fluorescent probe hybridization. *Genome Res* **6**: 639–645

Shiba, T., Simidu, U. and Taga, N. (1979) Distribution of aerobic-bacteria which contain bacteriochlorophyll-*a*. *Appl Environ Microbiol* **38**: 43–45

Shiba, T. (1991) *Roseobacter litoralis* gen. nov., sp. nov., and *Roseobacter denitrificans* sp. nov., aerobic pink-pigmented bacteria which contain bacteriochlorophyll-*a. Syst Appl Microbiol* **14**: 140–145

Smith, G.W., Ives, L.D., Nagelkerken, I.A. and Ritchie, K.B. (1996) Caribbean sea fan mortalities. *Nature* **383**: 487

Snijders, C.H.A. and Perkowski, J. (1990) Effects of head blight caused by *Fusarium culmorum* on toxin content and weight of wheat kernels. *Phytopathol* **80**: 566-570

Spaink, H.P., Okker, R.J.H., Wijffelman, C.A., Pees E., and Lugtenberg B.J.J. (1987) Promoters in the nodulation region of the *Rhizobium leguminosarum* Sym plasmid pRL1JI. *Plant Mol Biol* **9**: 27-39 Staskawicz, B., Dahlbeck, D., Keen, N. and Napoli, C. (1987) Molecular characterization of cloned avirulence genes from race 0 and race 1 of *Pseudomonas syringae* pv glycinea. *J Bacteriol* **169:** 5789–5794

Stauffer, L.T., Ghrist, A. and Stauffer, G.V. (1993) The *Escherichia coli gcvT* gene encoding the T-protein of the glycine cleavage enzyme system. *DNA Seq* **3:** 339-46

Stefels, M. J. (2000) Physiological aspects of the production and conversion of DMSP in marine algae and higher plants. *J Sea Res* **43**: 183–197

Stefels, J., Steinke, M., Turner, S., Malin, G. and Belviso, S. (2007) Environmental constraints on the production and removal of the climatically active gas dimethylsulphide (DMS) and implications for ecosystem modelling. *Biogeochem* **83**: 245–275

Steinke, M., Wolfe, G.V. and Kirst, G.O. (1998) Partial characterisation of dimethylsulfoniopropionate (DMSP) lyase isozymes in 6 strains of *Emiliania huxleyi*. *Mar Ecol Prog Ser* **175**: 215-225

Steinke, M., Malin, G. and Liss P.S. (2002) Trophic interactions in the sea: an ecological role for climate relevant volatiles. *J Phycol* **38**: 630-638

Steinke, M., Stefels, J. and Stamhuis, E. (2006) Dimethyl sulfide triggers search behavior in copepods. *Limnol Oceanogr* **51:** 1925-1930

Stines-Chaumeil, C., Talfournier, F. and Branlant, G. (2006) Mechanistic characterization of the MSDH (methylmalonate semialdehyde dehydrogenase) from *Bacillus subtilis*. *Biochem J* **395:** 107–115

Stoyanova, R., Querec, T.D. Brown, T.R. and Patriotis, C. (2004) Normalization of singlechannel DNA array data by principal component analysis. *Bioinformatics* **20**: 1772-1784

Studier, F. W. and Moffatt, B. A. (1986) Use of Bacteriophage T7 RNA polymerase to direct selective high-level expression of cloned genes. *J Mol Biol* **189**: 113-130

Sullivan, M.J., Curson, A.R.J., Shearer, N., Todd, J.D., Green, R.T. and Johnston, A.W.B. (2011) Unusual regulation of a leaderless operon involved in the catabolism of dimethylsulfonio-propionate in *Rhodobacter sphaeroides*. *PLoS One* **6**: e15972

Summers, P.S., Nolte, K.D., Cooper, A.J.L., Borgeas, H., Leustek, T., Rhodes, D. and Hanson, A.D. (1998) Identification and stereospecificity of the first three enzymes of 3dimethylsulfoniopropionate biosynthesis in a Chlorophyte Alga. *Plant Physiol* 116: 369-378

Sun, L., Curson, A.R.J., Todd, J.D. and Johnston, A.W.B. (2011) Diversity of DMSP transport in marine bacteria, revealed by genetic analyses. *Biogeochem* DOI 10.1007/s10533-011-9666-z

Sunda, W., Kieber, D.J., Kiene, R.P. and Huntsman, S. (2002) An antioxidant function for DMSP and DMS in marine algae. *Nature* **418**: 317–320

Sutton, J.C. (1982) Epidemiology of wheat head blight and maize ear rot caused by *Fusarium* graminearum. Canadian J Plant Pathol **4:** 195-209

Suylen, G.M.H., Large, P.J., van Dijken, J.P. and Kuenen, J.G. (1987) Methylmercaptan oxidase, a key enzyme in the metabolism of methylated sulphur compounds by *Hyphomicrobium* EG. *J Gen Microbiol* **133**: 2989–2997

Tang, T., François, N., Glatigny, A., Agier, N., Mucchielli, M.H., Aggerbeck, L. and Delacroix, H. (2007) Expression ratio evaluation in two-colour microarray experiments is significantly improved by correcting image misalignment. *Bioinformatics* **23**: 2686–2691

Tang, K., Huang, H., Jiao, N. and Wu, C.H. (2010) Phylogenomic analysis of marine *Roseobacters*. *PLoS One* **5**: e11604 1-9

Taylor, B. F., and Visscher, P.T. (1996) Metabolic pathways involved in DMSP degradation. In: Kiene, R.P., Visscher, P.T. Keller, M.D. and Kirst, G.O. (Eds.). Biological and environmental chemistry of DMSP and related sulfonium compounds, Plenum, New York, 265–276 Thrash, J.C., Boyd, A., Huggett, M.J., Grote, J., Carini, P., Yoder, R.J., Robbertse, B., Spatafora, J.W., Rappé, M.S. and Giovannoni, S.J. (2011) Phylogenomic evidence for a common ancestor of mitochondria and the SAR11 clade. *Scientific Reports* **1**: doi:10.1038/srep00013

Todd, J.D., Rogers, R., Li, Y.G., Wexler, M., Bond, P.L., Sun, L., Curson, A.R.J., Malin, G., Steinke, M. and Johnston, A. W. B. (2007) Structural and regulatory genes required to make the gas dimethyl sulfide in bacteria. *Science* **315**: 666–669

Todd, J.D., Curson, A.R.J., Dupont, C.L., Nicholson, P. and Johnston, A.W.B. (2009) The *dddP* gene, encoding a novel enzyme that converts dimethylsulfoniopropionate into dimethyl sulfide, is widespread in ocean metagenomes and marine bacteria and also occurs in some Ascomycete fungi. *Environ Microbiol* **11**: 1376-1385

Todd, J.D., Curson, A.R.J., Nikolaidou-Katsaridou, N., Brearley, C.A., Watmough, N.J., Chan, Y., Page, P.C.B., Sun, L. and Johnston, A.W.B. (2010) Molecular dissection of bacterial acrylate catabolism – unexpected links with dimethylsulfoniopropionate catabolism and dimethyl sulfide production. *Environ Microbiol* **12**: 327-343

Todd, J. D., Curson, A. R. J., Kirkwood, M., Sullivan, M. J., Green, R. T. and Johnston, A. W. B. (2011) DddQ, a novel, cupin-containing, dimethylsulfoniopropionate lyase in marine roseobacters and in uncultured marine bacteria *Environ Microbiol* **13**: 427–438

Todd, J.D., Kirkwood, M., Newton-Payne, S. and Johnston, A.W.B. (2012a) DddW, a third DMSP lyase in a model Roseobacter marine bacterium, *Ruegeria pomeroyi* DSS-3. *ISME J* 6: 223-226

Todd, J.D., Curson, A.R.J., Sullivan, M.J., Kirkwood, M. and Johnston, A.W.B. (2012b) The *Ruegeria pomeroyi acuI* gene has a role in DMSP catabolism and resembles *yhdH* of *E. coli* and other bacteria in conferring resistance to acrylate. *PLoS One* **7**: e35947

Tolli, J.D., Sievert, S.M. and Taylor, C.D. (2006) Unexpected diversity of bacteria capable of carbon monoxide oxidation in a coastal marine environment, and contribution of the *Roseobacter*-associated Clade to total CO oxidation. *Appl Environ Microbiol* **72:** 1966-1973

Trevena, A.J. and Jones, G.B. (2006) Dimethylsulfide and dimethylsulfoniopropionate in Antarctic sea ice and their release during sea ice melting. *Mar Chem* **98**: 210-222

Trinick, M.J. (1973) Symbiosis between *Rhizobium* and the non-legume *Trema aspera*. *Nature* **244:** 459–460

Tripp, H.J., Kitner, J.B., Schwalbach, M.S., Dacey, J.W.H., Wilhelm, L.J., Giovannoni, S.J. (2008) SAR11 marine bacteria require exogenous reduced sulphur for growth. *Nature* 452:741–744

Trossat, C., Rathinasabapathi, B., Weretilnyk, E.A., Shen, T-L., Huang, Z-H., Gage, D.A. and Hanson, A.D. (1998) Salinity promotes accumulation of 3-Dimethylsulfoniopropionate and its precursor *S*-methylmethionine in chloroplasts. *Plant Physiol* **116**: 165-171

Van Alstyne, K.L. and Houser, L.T. (2003a) Dimethylsulfide release during macroinvertebrate grazing and its role as an activated chemical defense. *Mar Ecol Prog Ser*250: 175-181

Van Alstyne, K.L., Schupp, P. And Slattery, M. (2006). The distribution of dimethylsulfoniopropionate in tropical Pacific coral reef invertebrates. *Coral Reefs* **25**: 321–327

Van Alstyne, K.L., Dominique III, V.J. and Muller-Parker, J. (2009) Is dimethylsulfoniopropionate (DMSP) produced by the symbionts or the host in an anemone– zooxanthella symbiosis? *Coral Reefs* **28**: 167-176

van der Palen, C.J., Slotboom, D.J., Jongejan, L., Reijnders, W.N., Harms, N., Duine, J.A. and van Spanning, R.J. (1995) Mutational analysis of *mau* genes involved in methylamine metabolism in *Paracoccus denitrificans*. *Eur J Biochem* **230**: 860-871

Van Duyl, F.C., Gieskes, W.W.C., Kop, A.J. and Lewis, W.E. (1998) Biological control of short-term variations in the concentration of DMSP and DMS during a *Phaeocystis* springbloom. *J Sea Res* **40**: 221–231

Varaljay, V.A., Howard, E.C., Sun, S. and Moran, M.A. (2010) Deep sequencing of a dimethylsulfoniopropionate degrading gene (*dmdA*) by using PCR primer pairs designed on the basis of marine metagenomic data. *Appl Environ Microbiol* **76:** 609-617

Venter, J.C., Remington, K., Heidelberg, J.F., Halpern, A.L., Rusch, D., Eisen, J.A., Wu, D., Paulsen, I., Nelson, K.E., Nelson, W., Fouts, D.E., Levy, S., Knap, A.H., Lomas, M.W., Nealson, K., White, O., Peterson, J., Hoffman, J., Parsons, R., Baden-Tillson, H., Pfannkoch, C., Rogers, Y.H. and Smith, H.O. (2004) Environmental genome shotgun sequencing of the Sargasso Sea. *Science* **304**: 66-74.

Vieira, J. and Messing, J. (1982) The pUC plasmids, an M13mp7-derived system for insertion mutagenesis and sequencing with synthetic universal primers. *Gene* **19**: 259–268

Vila-Costa, M., Simó, R., Harada, H., Gasol, J.M., Slezak, D. and Kiene, R.P. (2006) Dimethylsulfoniopropionate uptake by marine phytoplankton. *Science* **314**: 652–654

Visscher, P.T., Diaz, M.R. and Taylor, B.F. (1992) Enumeration of bacteria which cleave or demethylate dimethylsulfoniopropionate in the Caribbean Sea. *Mar Ecol Prog Ser* **89:** 293–26

Visscher, P.T., Taylor, B.F. (1994) Demethylation of dimethylsulfoniopropionate to 3mercaptopropionate by an aerobic marine bacterium. *Appl.Environ Microbiol* **60**: 4617–19

Wagner, J., Gruz, P., Kim, S.R., Yamada, M., Matsui, K., Fuchs, R.P. and Nohmi, T. (1999) The *dinB* gene encodes a novel *E. coli* DNA polymerase, DNA pol IV, involved in mutagenesis. *Mol Cell* **4:** 281-286

Wagner-Döbler, I. and Biebl, H. (2006) Environmental biology of the marine Roseobacter lineage. *Ann Rev Microbiol* **60:** 255–280

Wang, J., Nygaard, V., Smith-Sørensen, B., Hovig, E. and Myklebost, O. (2002) MArray: analyzing single, replicated or reversed microarray experiments. *Bioinformatics* **18**: 1139-1140

Wang, Y., Ma, X., Zhao, W., Jia, X., Kai, L. and Xu, X. (2006) Study on the creatinase from *Paracoccus* sp. strain WB1. *Process Biochem* **41**: 2072–2077

Watson, A.J. Fuller, L.J. Jeenes, D.J. and Archer, D.B (1999) Homologs of Aflatoxin biosynthesis genes and sequence of *aflR* in *Aspergillus oryzae* and *Aspergillus sojae*. *Appl Environ Microbiol* **65**: 307-310

Wexler, M., Yeoman, K.H., Stevens, J.B., de Luca, N.G., Sawers, G. and Johnston, A.W.B. (2001) The *Rhizobium leguminosarum tonB* gene is required for the uptake of siderophore and haem as sources of iron. *Mol Micro* **41**: 801-816

Williams, A. and Frasca, V. (2001) Ion-Exchange Chromatography. *Curr Protoc Protein Sci* 15: 8.2.1–8.2.30

Wolfe, G.V., Steinke, M. and Kirst, K.O. (1997) Grazing-activated chemical defence in a unicellular marine alga. *Nature* **387**: 894–897

Woo, E.J., Dunwell, J.M., Goodenough, P.W., Marvier, A.C. and Pickersgill, R.W. (2000) Germin is a manganese containing homohexamer with oxalate oxidase and superoxide dismutase activities. *Nat Struct Biol* **7:** 1036–1040

Wood, W. B. (1966) Host specificity of DNA produced by *Escherichia coli*; bacterial mutations affecting the restriction and modification of DNA. *J Mol Biol* **16**: 118-133

Yang, Y. H., Dudoit, S., Luu, P., and Speed, T. P. (2001) Normalization for cDNA microarray data. In: Bittner, M.L., Chen, Y., Dorsel, A.N. and Dougherty, E.R. (Eds.). Microarrays: Optical Technologies and Informatics. Proceedings of SPIE, International Society for Optical Engineering, Bellingham

Yanisch-Perron, C., Vieira, J., and Messing, J. (1985) Improved M13 phage cloning vectors and host strains: nucleotide sequences of the M13mp18 and pUC19 vectors. *Gene* **33**: 103–119

Yi, H., Lim, Y.W. and Chun, J. (2007) Taxonomic evaluation of the genera *Ruegeria* and *Silicibacter*: a proposal to transfer the genus *Silicibacter* Petursdottir and Kristjansson 1999 to the genus *Ruegeria* Uchino *et al.* 1999. *ISME J* **57**: 815-819

Yoch, D.C., Ansede, J.H. and Rabinowitz, K.S. (1997) Evidence for intracellular and extracellular dimethylsulfoniopropionate (DMSP) lyases and DMSP uptake sites in two species of marine bacteria. *Appl Environ Microbiol* **63**: 3182–3188

Yoch, D. C. (2002) Dimethylsulfoniopropionate: its sources, role in the marine food web, and biological degradation to dimethylsulfide. *Appl Environ Microbiol* **68**: 5804-5815

Young, J.P., Crossman, L.C., Johnston, A.W.B., Thomson, N.R., Ghazoui, Z.F., Hull, K.H.,
Wexler, M., Curson, A.R., Todd, J.D., Poole, P.S., Mauchline, T.H., East, A.K., Quail,
M.A., Churcher, C., Arrowsmith, C., Cherevach, I., Chillingworth, T., Clarke, K., Cronin,
A., Davis, P., Fraser, A., Hance, Z., Hauser, H., Jagels, K., Moule, S., Mungall,
K., Norbertczak, H., Rabbinowitsch, E., Sanders, M., Simmonds, M., Whitehead, S.
and Parkhill, J. (2006) The genome of *Rhizobium leguminosarum* has recognizable core and
accessory components. *Genome Biol* 7: R34

Zuo, Y. and Jones, R.D. (1995) Formation of carbon monoxide by photolysis of dissolved marine organic material and its significance in the carbon cycling of the oceans. *Naturwissenschaften* **82:** 472-474

#### Chapter 10

Appendix

#### **DNA** Deoxyribonucleic acid A1: Abbreviations **dNTP** deoxynucleotide triphosphate **ABC** ATP-binding cassete **DTT** dithiothreitol Acu Acrylate-utilising **EDTA** Ethylenediaminetetra-acetic acid **AMP** Adenosine monophosphate **EtBr** Ethidium bromide **Amp** Ampicillin FAD flavin adenine dinucleotide **Amp**<sup>**R**</sup> Ampicillin-resistant g gram **ATP** Adenosine triphosphate GC Gas chromatography **Bp** Base pair(s) Gent Gentamicin **BCCT** Betaine-Carnitine-Choline Gent<sup>R</sup> Gentamicin- resistant Transporter **GOS** Global Ocean Survey **BLAST** Basic Local Alignment Search **GTP** Guanosine triphosphate Tool **HGT** Horizontal gene transfer **CCN** Cloud condensation nuclei **3HP** 3-hydroxypropionate cDNA complementary DNA **HPLC** High pressure liquid **CO** Carbon monoxide chromatography CO<sub>2</sub> Carbon dioxide **IPTG** Isopropyl-β-D-CoA Coenzyme A thiogalactopyranoside **CODH** Carbon monoxide dehydrogenase k kilo **Da** Daltons Kan Kanamycin **Ddd** DMSP-dependent DMS Kan<sup>R</sup> Kanamycin-resistant **DEPC** diethyl pyrocarbonate **kbp** kilo base pair(s) dH2O Distilled H2O kDa kilo Daltons **DMS** Dimethyl sulfide **I** litre **DMSP** Dimethylsulfoniopropionate **LOWESS** Locally weighted scatterplot smoothing algorithm DMSHB D-4-dimethylsulfonio-2hydroxybutyrate m milli

transcriptase PCR

μ micro	<b>RBS</b> Ribosomal binding site
M Molar	Rif Rifampicin
MalSA Malonate semialdehyde	Rif <sup>R</sup> Rifampicin-resistant
Mb Megabase(s)	RNA Ribonucleic acid
MBM Marine basal medium	<b>rRNA</b> ribosomal RNA
MeSH methanethiol	<b>RT</b> Reverse transcriptase
MMPA methylmercaptopropionate	SDS Sodium dodecyl sulfate
MPA 3-mercaptopropionate	SMM S-methyl methionine
mol moles	sp. Species
mRNA messenger RNA	Spec Spectinomycin
MTHB D-4-methylthio-2-hydroxybutyrate	Spec <sup>R</sup> Spectionmycin-resistant
MTO Methanethiol oxidase	<b>spp.</b> Species (plural)
MTOB 4-methylthio-2-oxobutyrate	Str Streptomycin
<b>n</b> nano	Str <sup>R</sup> Streptomycin-resistant
NMR Nuclear magnetic resonance	Tet Tetracycline
<b>OD</b> Optical density	Tet <sup>R</sup> Tetracycline-resistant
<b>ONPG</b> <i>ortho</i> -nitrophenyl-β-D-	THF Tetrahydrofolate
galactopyranoside	Tris (hydroxymethyl) aminomethane
PAGE Polyacrylamide gel electrophoresis	UV Ultra-violet
PCR Polymerase chain reaction	X-gal 5-bromo-4-chloro-3-indolyl-β-D-
<b>p</b> pico	galactoside
q-RTPCR quantitative real-time reverse	YTSS Yeast-Tryptone-Sea Salts

#### **A2: Publications**

Todd, J.D., Curson, A.R.J., Sullivan, M.J., Kirkwood, M. and Johnston, A.W.B. (2012) The *Ruegeria pomeroyi acuI* gene has a role in DMSP catabolism and resembles *yhdH* of *E. coli* and other bacteria in conferring resistance to acrylate. *PLoS One* **7**: e35947

In this paper I was responsible for sensitivity tests, whereby YhdH was shown to facilitate growth in the presence of acrylate, and much of the work presented in chapter 5 directly precedes work presented in this journal article.

Todd, J.D., Curson, A.R.J., **Kirkwood, M**., Sullivan, M.J., Green, R.T., and Johnston, A.W.B. (2011) DddQ, a novel, cupin-containing, dimethylsulfoniopropionate lyase in marine Roseobacters and in uncultured marine bacteria. *Env Microbiol* **13**: 427-438

My work for this journal article involved alignments of the DddQ and DddL peptides, and several of the DMS assays showing activity of heterologous strains.

Todd, J.D., **Kirkwood, M.**, Newton-Payne, S. and Johnston, A.W.B. (2011) DddW, a third DMSP lyase in a model Roseobacter marine bacterium, *Ruegeria pomeroyi* DSS-3. *ISME J* 6: 223-226

Again, I was responsible for the alignment of the similar peptide regions. Also, work on heterologous expression of DddW, along with *lacZ* assays showing transcriptional regulation, were conducted by me, as shown in chapter 4 of this thesis.

**Kirkwood, M.**, Todd, J.D., Rypien, K.L. and Johnston, A.W.B. (2010) The opportunistic coral pathogen *Aspergillus sydowii* contains *dddP* and makes dimethyl sulphide from dimethylsulfoniopropionate. *ISME J* **4:** 147-150

**Kirkwood, M.**, Le Brun, N.E., Todd, J.D. and Johnston, A.W.B. (2010) The *dddP* gene of *Roseovarius nubinhibens* encodes a novel lyase that cleaves dimethylsulfoniopropionate into acrylate plus dimethyl sulphide. *Microbiol* **156**: 1900-1906

All work in both of these articles was carried out by me, and is presented in chapter 2.

## A3: Microarray data

# Table 10.1 Microarray data for the fold-change in gene expression in *Ruegeria pomeroyi* following exposure to

# DMSP, DMS or acrylate

SPO0029	SPO0028	SPO0027	SPO0026	SPO0025	SPO0024	SPO0023	SPO0022	SPO0021	SPO0020	SPO0019	SPO0018	SPO0017	SPO0016	SPO0015	SPO0014	SPO0013	SPO0012	SPO0011	SPO0010	SPO0009	SPO0008	SPO0007	SPO0006	SPO0005	SPO0004	SPO0003	SPO0002	SPO0001	Gene No.
rumA					hslO				ilvA		argG				msrA-1	rbsK	mae B	mutS	grpE	hrcA	rph				parB	parA	gidB	gidA	Name
23S rRNA (uracil-5-)-methyltransferase RumA	ABC transporter transmembrane ATP-binding protein	ABC transporter transmembrane ATP-binding protein	poly(A) polymerase	NUDIX family hydrolase	chaperonin, 33 kDa	NUDIX family hydrolase	response regulator	Hpt domain-containing protein	threonine dehydratase (EC:4.3.1.19)	hypothetical protein	argininosuccinate synthase (EC:6.3.4.5)	hypothetical protein	sterol carrier protein	hypothetical protein	methionine-S-sulfoxide reductase (EC:1.8.4)	ribokinase (EC:2.7.1.15)	malic enzyme (EC:1.1.1.40)	DNA mismatch repair protein MutS	co-chaperone GrpE	heat-inducible transcription repressor	ribonuclease PH (EC:2.7.7.56)	deoxyribonucleotide triphosphate pyrophosphatase	coproporphyrinogen III oxidase	hypothetical protein	chromosome partitioning protein parB	chromosome partitioning protein ParA	glucose-inhibited division protein B	tRNA uridine 5-carboxymethylaminomethyl modification protein GidA	Function
-1.25	-1.06	-1.27	-1.66	×	-1.69	-1.03	-1.76	1.51	1.06	-1.09	1.54	-3.41	-1.43	-1.47	1.30	1.52	1.48	x	1.81	1.51	-1.08	-1.87	-1.55	2.62	1.64	1.08	-1.17	-1.23	Dp1
0.09	0.74	0.75	0.20	х	0.00	0.83	0.01	0.00	0.67	0.23	0.04	0.04	0.04	0.00	0.15	0.11	0.01	х	0.00	0.00	0.74	0.00	0.40	0.00	0.16	0.97	x	0.04	Р
1.28	-1.12	1.56	-1.42	-1.36	-1.31	1.05	1.23	2.40	1.25	2.14	1.40	1.29	1.23	1.16	1.18	1.48	1.65	1.06	1.02	1.20	1.28	1.17	1.13	1.44	-1.08	0.98	1.17	-1.01	Dp2
0.01	0.57	0.10	0.33	0.01	0.03	0.66	0.05	0.07	0.01	0.00	0.06	0.20	0.53	0.58	0.22	0.03	0.01	0.59	0.99	0.01	0.08	0.39	0.42	0.45	0.60	0.84	0.33	0.92	Р
0.02	-1.09	0.15	-1.54	N/A	-1.50	0.01	-0.27	1.96	1.16	0.53	1.47	-1.06	-0.10	-0.16	1.24	1.50	1.57	N/A	1.42	1.36	0.10	-0.35	-0.21	2.03	0.28	1.03	0.00	-1.12	Mean
1.27	0.03	1.42	0.12	N/A	0.19	1.04	1.50	0.44	0.10	1.62	0.07	2.35	1.33	1.32	0.06	0.02	0.09	N/A	0.39	0.16	1.18	1.52	1.34	0.59	1.36	0.05	1.17	0.11	Error
2.03	-1.09	-1.19	1.85	3.31	2.19	1.86	-6.04	-7.49	1.33	-1.39	4.84	1.28	-1.46	1.19	-2.31	2.89	3.72	3.80	2.28	1.76	2.64	1.21	1.68	2.78	1.69	1.31	-0.89	-0.93	A1
0.00	0.92	0.87	0.19	0.00	0.00	0.00	0.00	0.00	0.06	0.08	0.00	0.47	0.04	0.01	0.00	0.01	0.00	0.00	0.00	0.00	0.02	0.07	0.15	0.02	0.03	0.40	0.03	0.04	Р
-1.59	1.03	-1.30	-1.66	-1.84	-1.33	-1.54	1.49	3.39	-1.41	1.07	-1.43	-1.42	-1.26	-1.17	1.09	-1.29	-1.26	-1.13	-1.37	-1.02	-1.42	-1.32	-1.21	-1.03	0.98	-1.28	-1.34	-1.46	A2
0.00	0.88	0.54	0.28	0.00	0.00	0.00	0.07	0.00	0.00	0.52	0.00	0.02	0.02	0.49	0.67	0.06	0.01	0.12	0.00	0.66	0.01		0.11		0.44	0.26	0.32	0.00	Р
0.22	-0.03	-1.25	0.10	0.74	0.43	0.16	-2.28	-2.05	-0.04	-0.16	1.71	-0.07	-1.36	0.01	-0.61	0.80	1.23	1.34	0.46	0.37	0.61	-0.06	0.24	0.88	1.33	0.02	-1.12	-1.19	Mean
1.81	1.06	0.06	1.76	2.58	1.76	1.70	3.77	5.44	1.37	1.23	3.14	1.35	0.10	1.18	1.70	2.09	2.49	2.47	1.83	1.39	2.03	1.27	1.45	1.91	0.36	1.30	0.22	0.27	Error
-1.08	-1.21	1.12	1.05	1.14	-1.12	-2.14	-1.54	-1.06	1.10	-1.02	-1.10	-1.26	1.09	1.15	-0.99	-1.01	-1.40	-1.26	-1.37	1.14	-1.37	-1.55	-1.37	-1.09	1.05	-1.16	-1.31	-1.64	Ds1
0.28	0.48	0.79	0.88	0.06	0.05	0.00	0.11	0.95			0.20	0.26	0.08	0.03	0.79	0.79	0.01	0.03	0.00	0.15	0.00	0.04	0.03		0.93	0.46	0.09	0.00	Р
-1.76	1.09	1.27	-1.02	-1.34	0.95	-2.21	-1.07	1.41	-1.00	-1.07	-1.46	-1.69	1.21	-1.03	1.58	-1.37	-1.93	-1.78	-1.38	-1.08	-1.82	-2.07	-1.64	-0.98	1.08	0.99	-1.48	-2.01	Ds2
0.00	0.59	0.41	1.00	0.00	0.18	0.00	0.75	0.37	0.75	0.36		0.00	0.27				0.00	_	0.00	0.29		-	0.07	0.86	0.88	0.80	0.29	0.00	Р
-1.42	-0.06	1.20	0.02	-0.10	-0.08	-2.18	-1.31	0.18	0.05	-1.05	-1.28	-1.48	1.15	0.06	0.29	-1.19	-1.67	-1.52	-1.38	0.03	-1.60	-1.81	-1.51	-1.04	1.07	-0.08	-1.40	-1.83	Mean
0.34	1.15	0.08	1.04	1.24	1.04	0.03	0.23	1.24	1.05	0.03	0.18	0.21	0.06	1.09	1.29	0.18	0.27	0.26	0.00	1.11	0.22	0.26	0.14	0.05	0.02	1.08	0.09	0.19	Error

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	nusA		infB	mutT	argJ		secA			radC						thiE	thiG	thiS		thiD	dnaJ	dnaK				cysQ	kdsB									
hypothetical protein peptide/opine/nickel uptake ABC transporter	transcription elongation factor NusA	hypothetical protein	translation initiation factor IF-2	mutator mutT protein (EC:3.6.1)	bifunctional ornithine acetyltransferase/N- acetylglutamate synthase (EC:2.3.1.1 2.3.1.35)	containing protein	preprotein translocase subunit SecA	hypothetical protein	OmpA domain-containing protein	DNA repair protein RadC	ABC transporter permease	ABC transporter ATP-binding protein	TENA/THI-4 family protein	ABC transporter substrate-binding protein	thiamine biosynthesis protein ThiF	(EC:2.5.1.3)	thiazole synthase	thiamine biosynthesis protein ThiS	thiamine biosynthesis oxidoreductase ThiO	phosphomethylpyrimidine kinase (EC:2.7.4.7)	molecular chaperone DnaJ	molecular chaperone Dna	alkylated DNA repair protein	hypothetical protein	ABC transporter permease	3'(2'),5'-bisphosphate nucleotidase (EC:3.1.3.7)	3-deoxy-manno-octulosonate cytidylyltransterase (EC:2.7.7.38)	hypothetical protein	hypothetical protein	core-2/I-branching enzyme family protein	hypothetical protein	LysR family transcriptional regulator	hypothetical protein	ErfK/YbiS/YcfS/YnhG family protein	voltage-gated sodium channel	(EC:2.1.1)
-1.72 -1.48	-1.53	-1.61	1.07	3.13	1.81	1.71	-1.06	-2.16	-1.17	1.13	Х	-1.63	-1.80	-1.23	Х	-1.06	Х	1.09	-1.26	-1.81	-1.34	0.97	1.13	1.33	1.19	-1.13	-1.09	-2.03	-1.56	-1.65	-1.14	-1.30	-1.49	1.17	1.80	
0.00 0.00	0.00	0.00	0.94	0.00	0.00	0.08	0.57	0.00	0.24	0.69	×	0.09	0.11	0.85	×	0.73	x	0.94	0.33	0.09	0.58	0.69	0.91	0.62	0.42	0.01	0.52	0.00	0.02	0.00	0.62	0.36	0.20	0.94	0.03	
1.05 1.05	0.97	-1.24	1.57	-1.12	-1.08	-1.35	-1.42	-1.08	1.04	1.19	-1.01	1.10	1.07	-1.06	-1.17	1.11	1.13	1.03	-1.11	-1.01	-1.19	-1.69	1.13	1.12	-1.18	1.26	-1.04	-1.19	-1.33	-1.02	-1.03	-1.10	-1.18	0.98	1.11	
0.75 0.73	0.34	0.04	0.02	0.06	0.32	0.17	0.00	0.46	0.89	0.28	0.97	0.35	0.48	0.81	0.26	0.28	0.14	0.90	0.57	1.00	0.28	0.01	0.26	0.53	0.26	0.30	0.67	0.04	0.15	0.91	0.87	0.19	0.31	0.84	0.48	
-0.34 -0.22	-0.28	-1.43	1.32	1.01	0.37	0.18	-1.24	-1.62	-0.06	1.16	N/A	-0.27	-0.37	-1.15	N/A	0.03	N/A	1.06	-1.19	-1.41	-1.27	-0.36	1.13	1.23	0.01	0.07	-1.07	-1.61	-1.45	-1.34	-1.09	-1.20	-1.34	1.07	1.46	
1.39 1.27	1.25	0.19	0.25	2.13	1.45	1.53	0.18	0.54	1.11	0.03	N/A	1.37	1.44	0.09	N/A	1.09	N/A	0.03	0.08	0.40	0.08	1.33	0.00	0.11	1.19	1.20	0.03	0.42	0.12	0.32	0.05	0.10	0.16	0.10	0.35	
1.34 -1.30	1.32	1.71	3.06	3.56	2.41	2.03	3.00	-2.96	2.11	-4.31	1.27	1.57	2.15	1.21	-0.97	1.45	1.48	1.41	1.38	1.15	2.22	2.62	1.41	0.86	-1.73	2.00	1.58	-1.97	-1.60	-1.42	-1.29	-1.10	-2.13	1.09	3.07	
0.02 0.01	0.08	0.00	0.00	0.00	0.00	0.04	0.00	0.00	0.00	0.00	0.01	0.09	0.06	0.83	0.01	0.00	0.00	0.70	0.32	0.22	0.37	0.05	0.64	0.54	0.00	0.00	0.00	0.00	0.08	0.04	0.49	0.25	0.01	0.94	0.00	
-1.58 1.16	-1.48	-1.30	-1.33	-1.27	-1.32	-1.30	-1.36	-1.15	1.02	1.44	-1.36	-1.65	-1.25	-1.30	-1.72	-1.73	-1.95	-1.21	-1.74	-2.14	-1.41	-1.63	-1.38	1.00	1.01	1.06	-1.06	-1.10	-1.06	-1.18	1.08	-1.46	-1.34	-1.23	-2.10	
0.00 0.18	0.00	0.01	0.11	0.02	0.01	0.09	0.01	0.50	0.94	0.01	0.10	0.00	0.08	0.19	0.03	0.00	0.02	0.55	0.00	0.02	0.20	0.00	0.14	0.71	0.74	0.81	0.51	0.21	0.71	0.29	0.85	0.00	0.14	0.34	0.00	
-0.12 -0.07	-0.08	0.21	0.87	1.15	0.55	0.37	0.82	-2.06	1.57	-1.44	-0.05	-0.04	0.45	-0.05	-1.34	-0.14	-0.24	0.10	-0.18	-0.50	0.41	0.50	0.02	0.93	-0.36	1.53	0.26	-1.54	-1.33	-1.30	-0.11	-1.28	-1.74	-0.07	0.49	
1.46 1.23	1.40	1.51	2.20	2.42	1.87	1.67	2.18	0.91	0.55	2.88	1.32	1.61	1.70	1.26	0.38	1.59	1.72	1.31	1.56	1.65	1.82	2.13	1.40	0.07	1.37	0.47	1.32	0.44	0.27	0.12	1.19	0.18	0.40	1.16	2.59	
-1.77 1.01	-1.61	-1.69	1.48	1.52	1.54	1.26	-1.30	-1.26	1.05	-1.05	1.08	1.09	-1.36	-1.42	-1.86	-1.38	-1.35	-1.00	-1.55	-0.97	1.18	1.21	1.26	1.65	1.23	-1.44	1.09	-1.75	-1.49	-1.15	-1.08	-1.12	-1.07	1.23	1.25	
0.00 0.95	0.00	0.00	0.04	0.00	0.00	0.31	0.04	0.17	0.71	0.98	0.35	0.82	0.01	0.07	0.00	0.03	0.17	0.97	0.01	0.82	0.58	0.32	0.10	0.00	0.01	0.00	0.75	0.00	0.07	0.34	0.64	0.29	0.79	0.53	0.16	
-2.65 -1.07	-2.11	-2.20	-1.32	1.30	1.04	1.72	0.97	-1.19	1.33	-1.22	-1.13	-0.98	-1.53	-1.68	-1.13	-2.03	х	1.28	-1.95	-1.26	1.32	1.49	1.11	1.61	1.35	-1.33	1.82	-1.18	-1.25	-1.68	-1.06	-1.42	-0.92	-1.09	-1.19	
0.00 0.49	0.00	0.00	0.21	0.03	0.41	0.05	0.20	0.89	0.12	x	0.74	0.77	0.08	0.12	0.97	0.01	×	0.25	0.00	0.63	0.55	0.28	0.40	0.01	0.02	0.06	0.02	0.24	0.30	0.00	0.75	0.10	0.74	0.66	0.29	
-2.21 -0.03	-1.86	-1.95	0.08	1.41	1.29	1.49	-0.16	-1.23	1.19	-1.14	-0.02	0.05	-1.45	-1.55	-1.50	-1.71	N/A	0.14	-1.75	-1.11	1.25	1.35	1.19	1.63	1.29	-1.39	1.46	-1.47	-1.37	-1.42	-1.07	-1.27	-1.00	0.07	0.03	
0.44 1.04	0.25	0.26	1.40	0.11	0.25	0.23	1.14	0.04	0.14	0.09	1.11	1.04	0.09	0.13	0.37	0.33	N/A	1.14	0.20	0.15	0.07	0.14	0.08	0.02	0.06	0.05	0.37	0.29	0.12	0.27	0.01	0.15	0.07	1.16	1.22	

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SP00059 SP00060 SP00062 SP00062 SP00063 SP00064 SP00065 SPO0058

SP00048 SP00050 SP00051 SP00052 SP00053 SP00054 SP00055 SP00055 SP00057

SP00038 SP00039 SP00040 SP00041 SP00042 SP00043 SP00043 SP00045 SP00046.1

SPO0036 SPO0037

SPO0030 SPO0031 SPO0032 SPO0033 SPO0034 SPO0035

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SPO0101	SPO0100	SPO0099	SPO0098	SPO0097	SPO0096	SPO0095	SPO0094	SPO0093	SPO0092	SPO0091	SPO0090	SPO0089	SPO0088	SPO0087	SPO0086	SPO0085	SPO0084	SPO0083	SPO0082	SPO0081	SPO0080	SPO0079	SPO0078	SPO0077	SPO0076	SPO0075	SPO0074	SPO0073	SPO0072	SPO0071	SPO0070	SPO0069	SPO0068	SPO0067	SPO0066	
						pncB		pncA									betB		kdsD					ptsN				hemH						ubiG	pip	
peptide/opine/nickel uptake ABC transporter	permease	permease permease pende/noine/nickel untake ABC transnorter	pepuer/Pener measuripease ABC transporter	aldehyde dehydrogenase	hypothetical protein	nicotinate phosphoribosyltransferase (EC:2.4.2.11)	sensory box sensor histidine kinase/response regulator (EC:2.7.3)	pyrazinamidase/nicotinamidase (EC:3.5.1.19)	molecular chaperone DnaK	rhodanese-like domain-containing protein	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	lipoprotein	hypothetical protein	betaine aldehyde dehydrogenase (EC:1.2.1.8)	exonuclease	arabinose 5-phosphate isomerase (EC:5.3.1.13)	hypothetical protein	hypothetical protein	ABC transporter ATP-binding protein	ribosomal subunit interface protein	r 1.5 utansporter subunit inte-fine introgen- regulatory protein PtsN	lipoprotein	ErfK/YbiS/YcfS/YnhG family protein	hypothetical protein	ferrochelatase (EC:4.99.1.1)	hypothetical protein	competence protein F	glutaredoxin	carbon-nitrogen family hydrolase	transcriptional regulator	3-ceme in y nordunnone-9-3-metri y nrans i erase (EC:2.1.1.64)	2 demethylubicuinore 0.3 methyltranefersor	substrate-binding protein
1.36	-1.86	-1.82	-1.44	-1.00	1.19	-1.38	-0.95	-1.35	-1.24	х	-1.23	1.95	-1.32	-2.00	1.13	-1.11	х	-1.20	1.33	1.06	1.12	-1.15	2.34	1.73	-1.18	-1.16	1.15	1.04	Х	0.97	0.88	1.39	-1.16	4.01	1.00	
0.55	0.10	0.02	0.00	0.94	0.12	0.12	0.61	0.00	×	×	0.79	0.00	0.16	0.04	0.02	0.72	х	0.02	0.08	0.43	0.84	0.03	0.04	0.07	0.01	0.37	0.64	0.99	х	0.15	0.70	0.01	0.56	0.00	0.72	
-1.03	-1.09	1.01	-1.01	1.40	1.09	-1.14	1.28	1.21	-1.33	1.25	-1.19	1.03	-1.38	-1.16	1.08	-1.68	-1.09	-1.40	-1.19	-1.51	-1.29	-1.56	0.98	1.04	2.61	-1.62	1.59	1.07	1.26	-1.21	-1.05	1.63	1.22	2.40	1.12	
0.72	0.50	0.95	0.99	0.08	0.52	0.54	0.28	0.00	0.32	0.00	0.58	0.66	0.16	0.77	0.50	0.09	0.53	0.03	0.20	0.08	0.03	0.00	0.70	0.87	0.00	0.00	0.00	0.68	0.08	0.36	0.62	0.01	0.06	0.00	0.28	_
0.17	-1.48	-0.41	-1.23	0.20	1.14	-1.26	0.17	-0.07	-1.29	N/A	-1.21	1.49	-1.35	-1.58	1.11	-1.40	N/A	-1.30	0.07	-0.23	-0.09	-1.36	1.66	1.39	0.72	-1.39	1.37	1.06	N/A	-0.12	-0.09	1.51	0.03	3.21	1.06	
1.20	0.39	1.42	0.21	1.20	0.05	0.12	1.11	1.28	0.05	N/A	0.02	0.46	0.03	0.42	0.02	0.29	N/A	0.10	1.26	1.29	1.21	0.21	0.68	0.35	1.90	0.23	0.22	0.02	N/A	1.09	0.96	0.12	1.19	0.80	0.06	
-1.83	-1.94	-1.58	-1.17	-1.15	1.18	1.46	-2.44	-2.14	4.45	3.59	-1.57	1.34	2.84	-2.18	-1.45	-2.58	1.86	1.88	3.70	2.60	1.93	1.77	1.25	1.21	1.10	1.00	1.29	1.27	1.37	-5.24	-1.86	1.51	1.44	3.21	1.19	
0.00	0.00	0.06	0.27	0.87	0.11	0.10	0.28	0.00	0.00	0.00	0.68	0.00	0.20	0.42	0.15	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.06	0.53	0.33	0.01	0.40	0.54	0.00	0.00	0.22	0.00	0.26	0.00	0.39	
1.90	1.50	1.86	1.34	1.84	1.15	1.01	x	1.38	-1.70	-1.92	-0.99	-1.35	1.04	-1.07	1.05	-1.53	-2.17	1.07	-1.36	-1.59	-1.54	-1.28	1.98	1.95	1.22	1.94	0.97	-1.44	-1.25	1.30	1.17	1.01	-1.28	-1.08	1.63	
0.01	0.01	0.03	0.05	0.03	0.61	0.93	×	0.01	0.07	0.00	0.95	0.04	0.83	0.77	х	0.06	0.00	0.71	0.00	0.00	0.00	0.00	0.00	0.00	0.04	0.00	0.02	0.00	0.04	0.10	0.26	0.91	0.01	0.25	0.00	
0.03	-0.22	0.14	0.09	0.35	1.17	1.24	N/A	-0.38	1.38	0.84	-1.28	-0.01	1.94	-1.63	-0.20	-2.06	-0.16	1.48	1.17	0.51	0.20	0.25	1.62	1.58	1.16	1.47	1.13	-0.09	0.06	-1.97	-0.35	1.26	0.08	1.07	1.41	
1.87	1.72	1.72	1.26	1.50	0.02	0.23	N/A	1.76	3.08	2.76	0.29	1.35	0.90	0.56	1.25	0.53	2.02	0.41	2.53	2.10	1.74	1.53	0.37	0.37	0.06	0.47	0.16	1.36	1.31	3.27	1.52	0.25	1.36	2.15	0.22	
1.36	-1.31	-1.36	1.26	1.26	1.37	1.11	-1.11	1.54	-2.08	-1.65	1.12	-1.33	-1.43	-1.10	-1.21	1.29	-1.50	-1.24	1.17	-1.17	1.42	-1.31	1.83	1.48	1.42	0.99	1.63	-1.28	-1.52	-1.29	1.05	-1.18	1.22	1.29	1.14	
0.09	0.09	0.06	0.01	0.30	0.03	0.76	0.80	0.03	0.02	0.00	0.80	0.44	0.29	0.82	0.35	0.13	0.00	0.04	0.27	0.15	0.00	0.01	0.00	0.01	0.01	0.34	0.02	0.15	0.00	0.34	0.92	0.07	0.09	0.02	0.34	
1.95	-1.08	-1.01	-1.16	1.19	-1.01	-1.14	×	-0.98	-1.85	-2.80	2.11	1.13	1.56	1.55	-0.98	2.66	-2.18	1.07	0.97	-1.21	1.15	-1.42	2.42	1.54	-1.32	1.21	1.28	-1.41	-2.30	2.01	1.03	-1.23	-1.02	-1.20	1.17	
0.02	0.84	0.92	0.61	0.27	0.88	0.58	x	0.19	0.08	0.00	0.10	0.34	0.15	0.05	0.64	0.00	0.00	0.72	0.15	0.01	0.14	0.00	0.00	0.01	0.02	0.17		0.01	0.00	0.00	0.90	0.11	0.83	0.20	0.02	
1.66	-1.20	-1.19	0.05	1.23	0.18	-0.01	N/A	0.28	-1.97	-2.23	-	-0.10	0.07	0.23	-1.09	1.98	-1.84	-0.09	1.07	-1.19		-1.37	2.13	1.51	0.05	1.10		-1.35			1.04	-1.21	0.10	0.05	1.16	
0.29	0.12	0.18	1.21	0.04	1.19	1.13	N/A	1.26	0.12	0.58	0.50	1.23	1.50	1.33	0.12	0.69	0.34	1.16	0.10	0.02	0.14	0.05	0.30	0.03	1.37	0.11		0.06	0.39	1.65	0.01	0.03	1.12	1.25	0.02	

SPO0140	SPO0139	SPO0138	SPO0137	SPO0136	SPO0135	SPO0134	SPO0133	SPO0132	SPO0131	SPO0130	SPO0129	SPO0128	SPO0127	SPO0126	SPO0125	SPO0124	SPO0123	SPO0122	SPO0120	SPO0119	SPO0118	SPO0116	SPO0115	SPO0114	SPO0113	SPO0112	SPO0111	SPO0110	SPO0109	SPO0108	SPO0107	SPO0106	SPO0105	SPO0104	SPO0103	SPO0102	
					pbpC							fabG				mepA						ackA			fabI-1						recQ			glpK	panC	panB	
nypotnetical protein	MoxR family ATPase	hypothetical protein	hypothetical protein	hypothetical protein	penicillin-binding protein 1C	hypothetical protein	PAN domain-containing protein	sensor histidine kinase/response regulator (EC:2.7.3)	isopentyl-diphosphate delta-isomerase (EC:5.3.3.2)	cytochrome c family protein	T4 family peptidase	3-ketoacyl-ACP reductase (EC:1.1.1.100)	peptidyl-tRNA hydrolase domain-containing protein	hypothetical protein	hypothetical protein	penicillin-insensitive murein endopeptidase	hypothetical protein	hypothetical protein	hypothetical protein	AsnC family transcriptional regulator	hypothetical protein	acetate kinase (EC:2.7.2.1)	orrunctional encyr-Cory nymatase/pirospirate acetyltransferase	hypothetical protein	enoyl-ACP reductase (EC:1.3.1.10)	poly(3-hydroxyalkanoate) polymerase	hypothetical protein	ABC transporter ATP-binding protein/permease	hypothetical protein	transposase, truncation	ATP-dependent DNA helicase RecQ (EC:3.6.1)	hypothetical protein	hypothetical protein	glycerol kinase (EC:2.7.1.30)	pantoatebeta-alanine ligase (EC:6.3.2.1)	3-methyl-2-oxobutanoate hydroxymethyltransferase (EC:2.1.2.11)	substrate-binding protein
-1.3/	0.93	-1.18	-1.48	-1.15	-1.21	-1.35	1.21	-1.10	3.38	5.48	-1.06	1.13	-2.84	-1.26	-2.14	-2.17	-1.59	1.39	-1.18	-1.22	1.68	1.51	2.35	-1.46	1.17	1.22	1.54	×	-1.03	х	-1.32	1.45	-1.06	-1.13	-1.11	1.44	
0.00	0.00	0.02	0.05	0.03	×	0.00	0.73	х	0.00	0.00	0.41	0.92	0.00	0.84	0.00	0.00	0.01	0.02	×	0.07	0.00	0.20	0.00	0.23	0.74	0.14	0.42	х	×	×	0.37	0.42	0.59	0.04	0.73	0.02	
-1.25	-1.35	-1.10	-1.12	-1.13	-1.01	1.10	1.14	1.42	1.83	1.76	1.02	-1.02	-1.32	1.31	-1.32	-1.42	-1.17	-1.85	-1.13	1.14	1.08	2.17	2.39	1.18	2.08	2.41	1.69	2.07	2.24	1.32	-1.06	1.09	1.01	-1.03	-1.26	-1.53	
0.02	0.03	0.04	0.06	0.02	0.97	0.29	0.20	0.14	0.00	0.01	0.72	0.81	0.10	0.24	0.06	0.04	0.00	0.00	0.49	0.05	0.32	0.03	0.00	0.29	0.03	0.00	0.04	0.00	0.00	0.00	0.76	0.48	0.96	0.77	0.10	0.06	
-1.30	-0.21	-1.14	-1.30	-1.14	-1.11	-0.13	1.18	0.16	2.61	3.62	-0.02	0.05	-2.08	0.03	-1.73	-1.80	-1.38	-0.23	-1.16	-0.04	1.38	1.84	2.37	-0.14	1.63	1.82	1.62	N/A	0.61	N/A	-1.19	1.27	-0.03	-1.08	-1.19	-0.05	
0.07	1.14	0.04	0.18	0.01	0.10	1.23	0.04	1.26	0.78	1.86	1.04	1.08	0.76	1.29	0.41	0.38	0.21	1.62	0.03	1.18	0.30	0.33	0.02	1.32	0.46	0.60	0.08	N/A	1.64	N/A	0.13	0.18	1.04	0.05	0.08	1.49	
C0.1	-1.86	-1.36	-1.72	-1.76	-1.22	-1.58	-1.47	-1.39	2.33	2.60	1.10	0.97	-3.60	2.35	1.08	1.19	-1.15	2.95	-1.49	-2.18	-1.48	2.98	9.42	1.25	5.37	4.63	8.13	21.40	12.70	2.40	0.91	2.19	-1.58	-1.14	1.88	5.27	
0.12	0.00	0.03	0.00	0.00	0.17	0.00	0.00	0.84	0.00	0.00	0.15	0.38	0.00	0.47	0.02	0.90	0.10	0.00	0.13	0.00	0.04	0.00	0.00	0.02	0.01	0.00	0.00	0.00	0.00	0.00	0.62	0.11	0.01	0.20	0.09	0.00	
1.01	1.52	1.72	1.40	1.49	1.46	1.28	1.57	-1.16	-1.27	1.57	-1.01	1.31	-1.01	-5.87	1.15	1.69	-1.25	-1.09	-1.11	-1.26	1.37	2.35	2.45	1.37	1.97	3.24	2.23	1.66	1.99	1.22	1.09	-1.10	1.04	-1.17	-2.37	-3.25	
0.99	0.00	0.00	0.00	0.00	0.02	0.03	0.01	x	0.02	0.01	0.76	0.08	0.58	0.00	0.49	0.00	0.01	0.07	x	0.06	0.01	0.02	0.00	0.05	0.00	0.00	0.01	0.00	0.01	0.02	0.82	0.18	0.86	0.01	0.00	0.00	
1.05	-0.17	0.18	-0.16	-0.14	0.12	-0.15	0.05	-1.28	0.53	2.09	0.05	1.14	-2.31	-1.76	1.12	1.44	-1.20	0.93	-1.30	-1.72	-0.05	2.67	5.94	1.31	3.67	3.94	5.18	11.53	7.35	1.81	1.00	0.55	-0.27	-1.16	-0.25	1.01	
0.02	1.69	1.54	1.56	1.63	1.34	1.43	1.52	0.12	1.80	0.52	1.06	0.17	1.30	4.11	0.03	0.25	0.05	2.02	0.19	0.46	1.43	0.32	3.49	0.06	1.70	0.70	2.95	9.87	5.36	0.59	0.09	1.65	1.31	0.02	2.13	4.26	
1.55	1.15	1.35	-1.00	1.25	1.49	1.65	1.62	1.13	1.19	1.26	1.04	1.13	-1.18	-2.18	1.52	1.64			1.09				-1.25	-1.60		-1.05	1.02		-1.34		1.36	1.01	1.18	-1.0	1.71	1.20	
5 U.UU		5 0.04	0 0.92	5 0.01	0.10	5 0.00	0.00	3 0.47	0.11	5 0.01	0.44	0.36	8 0.12	_	0.00	0.00	_	4 0.09	0.77				5 0.04	0.00	3 0.75	5 0.61	0.93		4 0.10	_	5 0.34	0.80	0.26	0.77	0.01	0.11	
) I. <u></u> 33		1.32	-1.03	1.19	) 1.32	) 1.29	) 1.66	7 X	1.22	1.66	4 -1.03	1.44	-1.32	3 -3.22	1.23	2.73		1.24	-1.33		-1.27	-	4 -1.41	) 1.22	-1.13		-1.06		-1.93	-	4 1.39	-1.28	5 1.46	-1.04	1.79	1.65	
0.06			ω				51			51	3	-	2	13	~	~	9	-	3	4	7	Ĩ			3	4	5	7	ω	5	1	8		4	Ĩ		

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1.41

1.47 1.64 N/A 0.01

0.07 0.16 1.04 0.20 0.02

1.29

1.25

1.21 1.46 0.00 0.10

2.19

-2.70

1.38

0.06

1.33

-0.19 -0.05 -1.15

-1.67 -1.51

-0.12

0.08 1.04 0.24 0.27 1.21 1.121 1.14 1.20 0.54 0.54

-1.16

-1.64

-2.01

 $\begin{array}{c} 0.23\\ 0.04\\ 0.02\\ 0.14\\ 1.15\\ 0.01\\ 0.16\\ 0.30\\ 0.21\\ 1.04\\ 1.08\\ 1.41\end{array}$ 

1.38 -0.14 1.32 -1.03

1.75 1.43

-0.02

1.39

1.34

-1.02 1.22

N/A 0.02 0.18 0.09 0.03 0.02 0.02 0.02

1.33

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SPO0180	SPO0179	SPO0178	SPO0177	SPO0176	SPO0175	SPO0174	SPO0173	SPO0172	SPO0171	SPO0170	SPO0169	SPO0168	SPO0167	SPO0166	SPO0165	SPO0164	SPO0163	SPO0162	SPO0161	SPO0160	SPO0157	SPO0156	SPO0155	SPO0154	SPO0153	SPO0152	SPO0151	SPO0150	SPO0149	SPO0148	SPO0147	SPO0146	SPO0145	SPO0144	SPO0143	SPO0142	SPO0141
fliE	fliQ	flgF	flgG	flgA	flgH			flhB	fliR	flhA												rtcR	gyrB				recF	dnaN	dnaA	rpsT		mutM	ubiE	ubiB		bktB	
flagellar hook-basal body protein FliE	flagellar biosynthetic protein FliQ	flagellar basal body rod protein FlgF	flagellar basal body rod protein FlgG	FigA	flagellar basal body L-ring protein flagellar basal body P-ring biosynthesis protein	hypothetical protein	hypothetical protein	flagellar biosynthesis protein FlhB	flagellar biosynthetic protein FliR	flagellar biosynthesis protein FlhA	Slt family transglycosylase	TetR family transcriptional regulator	trimethylamine methyltransferase	N-methylproline demethylase	pyrroline-5-carboxylate reductase	oxidoreductase, FMN-binding/pyridine nucleotide- disulfide oxidoreductase	sensory box histidine kinase/response regulator	hypothetical protein	LuxR family transcriptional regulator	hypothetical protein	TROVE domain-containing protein	transcriptional regulator RtcR	DNA gyrase subunit B (EC:5.99.1.3)	quinone family NAD(P)H dehydrogenase	TetR family transcriptional regulator	amino acid transporter LysE	recombination protein F	DNA polymerase III subunit beta (EC:2.7.7.7)	chromosome replication initiator DnaA	30S ribosomal protein S20	enoyl-CoA hydratase (EC:4.2.1.17)	EC:3.2.2.3)	methyltransferase UbiE	2-polyprenylphenol 6-hydroxylase (EC:1.14.13)	(Fe-S)-binding protein	beta-ketothiolase	DNA binding protein
2.05	1.71	2.25	2.54	2.13	2.13	2.04	Х	Х	-1.24	1.54	Х	Х	1.74	1.27	1.54	-1.13	-1.18	-1.02	-1.87	-1.00	-2.76	-1.57	1.31	-2.41	-1.36	-1.37	1.31	1.27	1.50	1.73	2.08	1.13	1.72	1.20	-2.39	6.45	-1.24
0.00	0.00	0.11	0.08	0.02	0.00	0.00	x	x	0.11	0.00	х	x	0.00	0.37	х	0.55	0.79	0.63	0.00	0.73	0.01	0.03	0.01	0.00	0.02	0.08	0.11	0.08	0.17	0.07	0.00	0.01	0.01	0.03	0.00	0.01	0.01
2.80	4.10	2.12	1.49	2.12	2.44	2.64	-1.06	-1.61	1.11	1.33	Х	-1.43	-1.31	-1.10	1.22	-1.17	1.43	2.35	-1.21	1.15	1.07	-1.17	-1.02	2.26	1.85	1.19	1.20	1.28	1.01	-1.32	1.16	-1.14	1.36	1.85	-1.33	2.25	-1.16
0.00	0.00	0.20	0.43	0.00	0.01	0.00	х	0.02	0.38	0.06	×	0.00	0.03	0.31	0.18	0.15	0.19	0.00	0.01	0.11	0.25	0.52	0.57	0.00	0.01	0.80	0.00	0.07	0.96	0.01	0.10	0.07	0.08	0.00	0.01	0.00	0.10
2.43	2.91	2.19	2.02	2.13	2.29	2.34	N/A	N/A	-0.06	1.44	N/A	N/A	0.22	0.09	1.38	-1.15	0.13	0.67	-1.54	0.08	-0.85	-1.37	0.15	-0.08	0.25	-0.09	1.26	1.28	1.26	0.21	1.62	-0.01	1.54	1.53	-1.86	4.35	-1.20
0.38	1.20	0.06	0.52	0.00	0.16	0.30	N/A	N/A	1.18	0.11	N/A	N/A	1.53	1.19	0.16	0.02	1.31	1.69	0.33	1.08	1.92	0.20	1.17	2.34	1.61	1.28	0.06	0.01	0.25	1.53	0.46	1.14	0.18	0.33	0.53	2.10	0.04
-4.92	-13.70	-2.68	-3.19	-5.39	-4.21	-3.81	-2.21	-1.06	-1.62	-1.08	Х	1.80	-1.56	-1.35	1.54	-1.54	-1.40	-1.76	1.10	1.36	1.18	-1.49	1.42	1.09	1.93	1.66	2.27	1.66	1.32	5.84	3.56	1.37	2.50	-1.37	-2.97	1.67	1.88
0.00	0.00	0.40	0.04	0.00	0.00	0.00	0.00	x	0.03	0.32	Х	0.00	0.00	0.07	0.01	0.34	0.72	0.05	0.18	0.00	0.27	0.72	0.01	0.04	0.00	0.23	0.00	0.00	0.34	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
4.39	7.01	3.18	1.86	3.74	2.44	5.14	Х	-1.52	-1.07	1.18	Х	-1.10	1.27	-1.13	-1.01	-1.16	1.25	3.21	1.09	1.04	1.20	-1.16	-1.45	-1.94	-1.08	-1.11	-1.29	-1.13	1.07	-1.37	1.10	1.31	-1.11	-1.47	1.85	1.96	1.14
0.00	0.00	0.07	0.01	0.00	0.18	0.09	x	0.18	x	0.44	x	0.46	0.17	0.15	0.88	0.23	0.33	0.00	0.29	0.59	0.07	0.61	0.00	0.00	0.32	0.81	0.03	0.09	0.94	0.01	0.55	0.01	0.25	0.00	0.00	0.00	0.20
-0.27	-3.35	0.25	-0.67	-0.83	-0.89	0.67	N/A	-1.29	-1.35	0.05	N/A	0.35	-0.15	-1.24	0.27	-1.35	-0.08	0.73	1.10	1.20	1.19	-1.33	-0.02	-0.43	0.43	0.28	0.49	0.27	1.20	2.24	2.33	1.34	0.70	-1.42	-0.56	1.82	1.51
4.66	10.36	2.93	2.53	4.57	3.33	4.48	N/A	0.23	0.27	1.13	N/A	1.45	1.42	0.11	1.28	0.19	1.33	2.49	0.01	0.16	0.01	0.16	1.44	1.52	1.51	1.39	1.78	1.40	0.13	3.61	1.23	0.03	1.81	0.05	2.41	0.15	0.37
-1.63	-1.60	-1.05	-1.14	-1.55	-1.18	-1.18	-1.28	-1.77	1.41	-1.07	Х	-1.46	-1.46	-1.18	1.17	-1.28	1.52	1.17	-1.10	1.14	-1.77	-1.09	1.01	1.21	1.25	1.28	1.17	1.17	-1.14	-1.98	1.14	-1.11	-1.84	1.19	-1.23	1.74	-1.42
0.00	0.01	0.98	0.69	0.00	0.70	0.11	0.29	0.03	0.01	0.89	x	0.00	0.02	0.14	0.15	0.06	0.12	0.15	0.02	0.13	0.00	-	0.76	0.02	0.01	0.50	0.01	0.19	0.52	0.00	0.01	0.07	0.00	0.01	0.01	0.00	
-1.25	x	2.00	1.52	×	-1.65	x	х	-1.02	Х	x	Х	-1.76	-1.18	-1.21	-1.08	-0.98	1.30	х	1.19	-0.96	-2.50	-1.15	-1.27	4.53	1.19	1.08	-1.34	-1.14	-1.18	-2.87	-1.26	-1.13	-2.09	-1.60	-1.08	2.31	1.34
x	x	0.11	x	×		×	x	0.90	x	x	x	0.00	0.29	0.63	×	0.35	0.22	x	0.07	0.15	0.00		0.01	0.00	0.17	0.86	-	0.30	0.48	0.00	0.10	0.26	0.00	0.00	0.84		0.02
-1.44	N/A	0.48	0.19	N/A	-1.42	N/A	N/A	-1.40	N/A	N/A	N/A	-1.61	-1.32	-1.20	0.04	-1.13	1.41	N/A		0.09						1.18			-1.16			-1.12	-1.97				
0.19	N/A	1.53	1.33	N/A	0.23	N/A	N/A	0.38	N/A	N/A	N/A	0.15	0.14	0.02	1.13	0.15	0.11	N/A	1.15	1.05	0.37	0.03	1.14	1.66	0.03	0.10	1.26	1.16	0.02	0.45	1.20	0.01	0.13	1.40	0.08	0.29	1.38

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SPO0218	SPO0217	SPO0216	SPO0215	SPO0214	SPO0213	SPO0212	SPO0211	SPO0210	SPO0209	SPO0207	SPO0206	SPO0205	SPO0204	SPO0203	SPO0202	SPO0201	SPO0200	SPO0199	SPO0198	SPO0197	SPO0196	SPO0195	SPO0194	SPO0193	SPO0192	SPO0191	SPO0190	SPO0189	SPO0188	SPO0187	SPO0186	SPO0185	SPO0184	SPO0183	SPO0182	SPO0181
		leuC-1	leuD					leuB						motA			fliL	fliF		fiiN	fliP	flgI		flgK		motB								fiil	flgB	flgC
mechanosensitive ion channel protein MscS	hypothetical protein	isopropylmalate isomerase large subunit (EC:4.2.1.33)	sopropy imatate isomerase small subunit (EC:4.2.1.33)	hypothetical protein	hypothetical protein	hypothetical protein	LysR family transcriptional regulator	3-isopropylmalate dehydrogenase (EC:1.1.1.85)	Na/Pi-cotransporter family protein	glycine cleavage system protein T	LysR family transcriptional regulator	MmgE/PrpD family protein	hypothetical protein	flagellar motor protein MotA	hypothetical protein	hypothetical protein	flagellar basal body protein FliL	flagellar MS-ring protein	ABC transporter ATP-binding protein, flagellar	flagellar motor switch protein FliN	flagellar biosynthesis protein FliP	надения разая осод и типд отозунинсять расски. FlgA	flagellar basal body P_ring biosynthesis motein	flagellar hook-associated protein Flg	flagellar hook protein FlgE	chemotaxis protein MotB	GMC family oxidoreductase	hypothetical protein	sensor histidine kinase (EC:2.7.3)	DNA-binding response regulator	hypothetical protein	hypothetical protein	hypothetical protein	ri+-transporting two-sector ATPase, magenum- specific (EC:3.6.3.14)	flagellar basal-body rod protein FlgB	flagellar basal body rod protein FlgC
-1.61	Х	1.55	1.63	-1.75	-1.26	-2.33	Х	2.41	1.46	29.60	1.33	43.00	2.21	2.42	2.58	2.40	1.17	3.40	1.90	2.74	1.21	х	2.10	2.43	1.88	-1.03	2.33	1.07	1.15	-1.53	1.01	2.32	2.69	1.27	2.03	2.25
0.00	х	0.18	0.00	0.01	0.01	0.00	х	0.00	0.04	0.00	0.01	0.00	0.00	0.00	0.00	0.00	0.68	0.00	0.12	0.00	0.77	×	0.00	0.00	0.01	0.47	0.00	0.86	0.44	0.00	0.02	0.12	0.00	0.65	0.00	0.00
1.19	-1.37	-1.24	1.02	1.01	-1.06	-1.53	-1.13	1.48	1.09	-1.18	1.14	-1.34	1.54	2.94	1.50	2.39	-1.22	1.43	2.14	3.19	-1.16	1.50	1.37	1.78	2.06	1.37	1.45	1.02	1.40	-1.21	-1.77	-1.48	-1.34	1.29	1.03	4.52
0.11	0.50	0.15	0.99	0.98	0.62	0.37	0.31	0.04	0.11	0.20	0.04	0.07	0.00	0.00	0.09	0.00	0.48	0.24	0.01	0.01	0.68	0.09	0.11	0.01	0.01	0.40	0.12	0.84	0.01	0.18	0.00	0.02	0.19	0.35	0.58	0.00
-0.21	N/A	0.16	1.33	-0.37	-1.16	-1.93	N/A	1.95	1.28	14.21	1.24	20.83	1.88	2.68	2.04	2.40	-0.03	2.42	2.02	2.97	0.03	N/A	1.74	2.11	1.97	0.17	1.89	1.05	1.28	-1.37	-0.38	0.42	0.68	1.28	1.53	3.39
1.40	N/A	1.40	0.31	1.38	0.10	0.40	N/A	0.47	0.19	15.39	0.10	22.17	0.34	0.26	0.54	0.00	1.20	0.98	0.12	0.23	1.19	N/A	0.37	0.33	0.09	1.20	0.44	0.03	0.13	0.16	1.39	1.90	2.02	0.01	0.50	1.14
-1.59	-2.85	3.46	3.80	-1.72	1.01	-6.48	-1.50	3.43	1.23	-1.26	-0.98	-4.63	-2.23	-3.46	-5.84	-4.81	-5.67	-7.27	-5.06	-4.14	1.46	x	-1.11	-1.24	-4.65	-3.10	-1.34	1.40	-1.06	-2.30	-3.57	1.76	1.64	-1.13	-5.68	-13.00
0.01	0.07	0.01	0.00	0.07	0.91	0.00	0.30	0.00	0.05	0.41	0.05	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.58	х	0.30	0.83	0.00	0.00	0.37	0.22	0.40	0.00	0.00	0.38	0.00	0.85	0.00	0.00
1.44	-1.11	-1.39	-1.52	-1.71	-1.23	-1.26	-1.42	-1.07	1.37	1.08	1.06	х	1.10	4.86	2.15	2.28	1.13	3.64	2.06	4.19	1.21	×	1.72	2.14	2.66	1.21	1.48	1.25	1.11	-1.04	-1.02	-1.21	-1.30	1.11	1.35	5.51
0.00	0.82	0.01	0.01	0.03	0.21	0.53	0.20	0.08	0.00	0.31	0.48	x	×	0.00	0.00	0.00	0.77	0.00	0.07	0.02	0.69	×	0.02	0.04	0.01	0.57	0.04	0.11	0.33	0.75	0.61	0.27	0.43	0.63	0.16	0.00
-0.08	-1.98	1.04	1.14	-1.72	-0.11	-3.87	-1.46	1.18	1.30	-0.09	0.04	N/A	-0.57	0.70	-1.85	-1.27	-2.27	-1.82	-1.50	0.03	1.34	N/A	0.31	0.45	-1.00	-0.95	0.07	1.33	0.03	-1.67	-2.30	0.28	0.17	-0.01	-2.17	-3.75
1.52	0.87	2.43	2.66	0.01	1.12	2.61	0.04	2.25	0.07	1.17	1.02	N/A	1.67	4.16	4.00	3.55	3.40	5.46	3.56	4.17	0.13	N/A	1.42	1.69	3.66	2.16	1.41	0.08	1.09	0.63	1.28	1.49	1.47	1.12	3.52	9.26
1.55	-1.10	1.00	-1.30	-1.18	-0.98	-1.06	-1.05	-1.05	1.37	1.17	1.31	-1.27	1.12	-1.07	-1.59	1.14	-1.33	-1.25	-1.08	-1.07	-1.35	-1.18	-1.31	-1.18	-1.28	-1.14	1.42	-1.40	1.10	-1.13	-3.34	-2.37	-1.73	-0.96	-1.05	-1.66
0.00	0.83	0.65	0.16	0.36	0.58	0.95	0.71	0.42	0.02	0.02	0.01	0.13	0.11	0.89	0.05	0.09	0.54	0.36	0.81	x	0.25	0.29	0.22	0.52	0.42	0.65	0.16	0.34	0.27	0.41	0.00	0.00	0.23	0.78	0.89	0.06
1.16	х	-1.32	-1.71	-1.25	-1.05	1.44	-1.02	-1.32	-1.02	-1.38	-1.16	x	x	Х	-0.95	x	1.65	-0.93	1.58	x	1.57	x	-1.04	-1.04	-0.96	1.20	2.64	-1.11	-1.22	1.34	-3.84	-5.01	-1.68	1.13	1.47	Х
0.10	х	0.05	0.01	0.27	0.62	x	0.56	0.00	0.31	0.32	0.94	x	x	Х	0.57	x	0.36	0.39	0.14	x	0.28	x	0.83	0.78	0.42	0.45	0.02	0.76	0.22	0.03	0.00	0.00	0.30	0.53	0.02	×
1.36	N/A	-0.16	-1.51	-1.22	-1.01	0.19	-1.04	-1.19	0.18	-0.11	0.08	N/A	N/A	N/A	-1.27	N/A	0.16	-1.09	0.25	N/A	0.11	N/A	-1.18	-1.11	-1.12		2.03	-1.26	-0.06	0.11	-3.59	-3.69	-1.71	0.09	0.21	N/A
0.20	N/A	1.16	0.21	0.04	0.04	1.25	0.02	0.14	1.20	1.28	1.24	N/A	N/A	N/A	0.32	N/A	1.49	0.16	1.33	N/A	1.46	N/A	0.14	0.07	0.16	1.17	0.61	0.15	1.16	1.24	0.25	1.32	0.03	1.04	1.26	N/A

## Chapter 10: Appendix

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SPO0259	SPO0258	SPO0257	SPO0256	SPO0255	SPO0254	SPO0253	SPO0252	SPO0251	SPO0250	SPO0249	SPO0248	SPO0247	SPO0241	SPO0240	SPO0239	SPO0238	SPO0237	SPO0236	SPO0235	SPO0234	SPO0233	SPO0232	SPO0231	SPO0230	SPO0229	SPO0228	SPO0227	SPO0226	SPO0225	SPO0224	SPO0223	SPO0222	SPO0221	SPO0220	SPO0219
					ohr	ilvE	petP	petR		xseB	ispA	dxs		ugpB	ugpA	ugpE	ugpC								rpsU							ald			
hypothetical protein	AraC family transcriptional regulator	hypothetical protein	plastocyanin/azurin family copper binding protein	MarR family transcriptional regulator	organic hydroperoxide resistance protein	branched-chain amino acid aminotransierase (EC:2.6.1.42)	transcriptional regulator PetP	DNA-binding response regulator PetR	histone deacetylase	(EC:3.1.11.6)	farnesyl diphosphate synthase (EC:2.5.1.10) exodeoxyribonuclease VII small subunit	I-deoxy-D-xy1ulose-5-pnospnate synthase (EC:2.2.1.7)	LysR family transcriptional regulator	SN-glycerol-3-phophate ABC transporter substrate- binding protein	SN-glycerol-3-phosphate ABC transporter	permease	glycerol-3-phosphate transporter ATP-bunding subunit (EC:3.6.3.20) SN-olycerol-3-phosphate ABC transporter	glycerophosphoryl diester phosphodiesterase	aldehyde dehydrogenase	hypothetical protein	AsnC family transcriptional regulator	quinone oxidoreductase	alcohol dehydrogenase	hypothetical protein	30S ribosomal protein S21	hypothetical protein	PaxA	cytochrome P450 family protein	transcriptional regulator	polyphosphate kinase	AsnC family transcriptional regulator	alanine dehydrogenase (EC:1.4.1.1)	hypothetical protein	rRNA large subunit methyltransferase	iojap family protein
-1.19	-1.33	-0.97	-1.04	-1.31	-1.54	1.11	-1.62	-2.15	-1.30	0.99	1.40	1.35	-1.31	-1.36	1.14	-1.13	1.56	-2.31	-3.94	-4.02	1.29	1.13	2.19	1.38	-1.39	4.77	-2.63	×	-1.48	1.03	-1.01	15.40	-1.22	-1.02	-1.45
0.04	0.44	0.73	0.94	0.08	0.00	0.20	0.00	0.08	0.43	0.53	0.74	0.09	0.60	0.02	0.33	0.82	0.01	0.17	0.00	0.00	0.03	0.59	0.00	0.02	0.00	0.00	0.00	x	0.01	0.85	0.21	0.00	0.87	0.80	0.00
-1.27	-1.37	-1.60	-1.42	1.47	1.47	-1.59	-1.28	-2.12	-1.82	1.06	1.12	1.23	-1.18	-1.53	-1.21	-1.07	1.03	-1.29	-1.56	-1.00	-1.03	-1.29	1.02	-1.10	1.16	1.78	-2.69	-1.06	-1.11	-1.40	-1.47	-1.21	-1.62	-1.51	-1.67
0.05	0.18	0.05	0.18	0.05	0.00	0.01	0.05	0.00	0.05	0.76	0.77	0.02	0.76	0.02	0.42	0.65	0.61	0.20	0.07	0.99	0.88	0.07	0.93	0.27	0.59	0.00	0.00	0.57	0.55	0.17	0.00	0.16	0.04	0.03	0.00
-1.23	-1.35	-1.28	-1.23	0.08	-0.04	-0.24	-1.45	-2.14	-1.56	1.03	1.26	1.29	-1.25	-1.45	-0.04	-1.10	1.30	-1.80	-2.75	-2.51	0.13	-0.08	1.61	0.14	-0.12	3.28	-2.66	N/A	-1.30	-0.19	-1.24	7.10	-1.42	-1.27	-1.56
0.04	0.02	0.32	0.19	1.39	1.51	1.35	0.17	0.01	0.26	0.04	0.14	0.06	0.07	0.09	1.18	0.03	0.27	0.51	1.19	1.51	1.16	1.21	0.59	1.24	1.28	1.50	0.03	N/A	0.19	1.22	0.23	8.31	0.20	0.24	0.11
1.48	0.87	-1.03	1.28	1.99	11.20	1.98	-1.89	-2.44	-1.65	1.13	1.70	1.79	-1.24	-1.23	1.21	-0.99	1.70	-2.39	-16.50	-25.60	-1.50	-1.33	2.23	1.81	5.36	-1.60	3.16	1.44	-1.10	1.40	-1.55	8.29	1.95	1.40	0.92
0.00	0.66	0.64	0.00	0.00	0.00	0.00	0.00	0.00	0.20	0.41	0.52	0.00	0.97	0.04	0.15	0.65	0.00	0.01	0.00	0.00	0.02	0.00	0.00	0.00	0.00	0.06	0.00	0.00	0.26	0.20	0.00	0.00	0.61	0.07	0.00
-1.13	1.23	1.12	-1.01	1.27	1.04	0.98	-1.02	-1.14	-1.12	1.03	1.02	-1.15	-1.24	-1.04	-1.16	-1.24	-1.51	-1.59	-1.21	-1.23	1.22	1.52	-1.38	-1.17	-1.39	1.25	-1.37	-1.16	-1.08	-1.09	-1.20	-2.47	-1.13	-1.16	1.11
0.28	0.48	0.67	0.99	0.12	0.91	0.56	0.48	0.08	0.38	0.85	0.95	0.29	0.70	0.42	0.18	0.30	0.05	0.08	0.36	0.56	0.00	0.00	0.01	0.02	0.00	0.04	0.00	0.31	0.48	0.63	0.01	0.00	0.36	0.01	0.63
0.18	1.05	0.05	0.14	1.63	6.12	1.48	-1.46	-1.79	-1.39	1.08	1.36	0.32	-1.24	-1.14	0.03	-1.12	0.10	-1.99	-8.86	-13.42	-0.14	0.10	0.43	0.32	1.99	-0.18	0.90	0.14	-1.09	0.16	-1.38	2.91	0.41	0.12	1.02
1.31	0.18	1.08	1.15	0.36	5.08	0.50	0.43	0.65	0.27	0.05	0.34	1.47	0.00	0.10	1.19	0.12	1.61	0.40	7.65	12.19	1.36	1.43	1.81	1.49	3.38	1.43	2.27	1.30	0.01	1.25	0.18	5.38	1.54	1.28	0.09
-1.15	-1.00	1.03	-1.01	1.05	-1.86	-2.19	-1.28	-2.00	-1.86	-1.28	-1.07	-1.19	-1.17	-1.27	-1.10	-1.32	-1.15	-1.33	-1.27	-1.18	-1.46	1.44	1.70	-1.03	2.15	1.21	3.28	1.50	-1.31	1.44	-1.29	1.55	1.23	1.60	2.04
0.20	0.93	0.86	0.98	0.81	0.00	0.00	0.01	0.00	0.03	0.00	0.79	0.12	0.79	0.01	0.47	0.11	0.32	0.06	0.16	0.58	0.00	0.01	0.00	0.16	0.00	0.22	0.00	0.01	0.01	0.18	0.00	0.00	0.27	0.00	0.00
X	1.01	2.01	1.60	-1.04	-2.02	-4.68	-1.57	-2.48	-2.74	-2.34	-1.29	-1.60	1.12	1.23	-1.12	-1.18	-1.35	1.35	1.76	1.37	-1.46	1.90	1.32	-1.21	-1.52	1.67	1.87	-1.60	-1.82	1.65	-1.55	1.79	0.99	1.17	1.76
X	0.99	0.08	0.14	0.98	0.00	0.00	0.00	0.00	0.00	0.00	0.55	0.01	0.77	0.09	0.94	0.86	0.46	0.05	0.00	0.24	0.02	0.04	0.01	0.00	0.00	0.03	0.00	x	0.00	0.12	0.00	0.00	0.93	0.50	0.01
N/A	0.01	1.52	0.30	0.01	-1.94	-3.44	-1.43	-2.24	-2.30	-1.81	-1.18	-1.40	-0.02	-0.02	-1.11	-1.25	-1.25	0.01	0.25	0.10	-1.46	1.67	1.51	-1.12	0.32	1.44	2.58	-0.05	-1.57	1.55	-1.42	1.67	1.11	1.39	1.90
N/A	1.01	0.49	1.31	1.05	0.08	1.25	0.14	0.24	0.44	0.53	0.11	0.21	1.15	1.25	0.01	0.07	0.10	1.34	1.52	1.28	0.00	0.23	0.19	0.09	1.84	0.23	0.70	1.55	0.26	0.11	0.13	0.12	0.12	0.22	0.14

SPO0269 SPO0270

SPO0266 SPO0267 SPO0268

SPO0271

SPO0272

SPO0262 SPO0263 SPO0264 SPO0265

SPO0260 SPO0261

## Chapter 10: Appendix

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										udgA	pyrB								dapF					petC	petB	petA			ribH-1	aroC					cobO		
acyl-CoA dehydrogenase	Sua5/YciO/YrdC family protein	hypothetical protein	hypothetical protein	NUDIX family hydrolase	AcrB/AcrD/AcrF family transporter	RND family efflux transporter MFP subunit	molybdenum cofactor biosynthesis protein B	transmembrane amino acid efflux protein	Ser/Thr protein phosphatase	uracil-DNA glycosylase (EC:3.2.2)	aspartate carbamoyltransferase (EC:2.1.3.2)	hypothetical protein	hypothetical protein	dihydroorotase (EC:3.5.2.3)	hypothetical protein	hypothetical protein	hypothetical protein	transposase, truncation	diaminopimelate epimerase (EC:5.1.1.7)	MiaB-like tRNA modifying enzyme	LuxR family transcriptional regulator	cytochrome b	hypothetical protein	ионуцинот-су ослионие с технисказе, суностнотие ст (EC:1.10.2.2)	(EC:1.10.2.2) historical contracts of the second contr	subunit (EC:1.10.2.2) ubituinol-evtrochrome c reductase, evtochrome B	glutathione S-transferase	negative transcriptional regulator	riboflavin synthase subunit beta (EC:2.5.1.9)	chorismate synthase (EC:4.2.3.5)	HD domain-containing protein	auxin efflux carrier protein	CaiB/BaiF family protein	hypothetical protein	cob(I)alamin adenosyltransferase (EC:2.5.1.17)	hypothetical protein	alkaline phosphatase
2.75	-1.28	-1.24	1.17	-1.10	-1.53	-1.21	1.43	1.23	1.55	1.02	1.66	2.29	1.20	1.41	1.04	-1.24	1.36	Х	-1.30	-1.15	-2.30	1.17	Х	5.01	6.19	4.47	1.40	-1.14	-1.84	-1.52	-2.51	-1.46	1.16	Х	1.62	х	-1.18
0.03	0.02	0.54	0.04	0.64	0.00	0.32	0.02	0.54	0.78	0.97	0.00	0.00	0.03	0.00	0.16	0.06	0.09	х	0.00	0.13	0.01	0.41	x	0.00	0.00	0.00	0.00	0.69	0.00	0.00	0.01	0.38	0.11	x	0.00	х	0.13
2.95	-1.10	1.09	-1.22	-1.11	-1.17	1.37	-1.07	1.06	1.31	1.00	1.33	1.36	1.29	1.30	1.27	1.34	-1.23	-0.99	-1.25	-1.37	-1.08	1.10	-1.08	1.80	1.34	1.17	1.01	1.08	1.00	1.03	1.03	1.36	1.53	-1.20	1.27	1.21	1.03
0.00	0.05	0.69	0.02	0.43	0.31	0.03	0.13	0.60	0.01	1.00	0.02	0.00	0.05	0.03	0.27	0.05	0.15	0.46	0.03	0.05	0.71	0.45	0.83	0.01	0.10	0.51	0.95	0.47	1.00	0.84	0.86	0.60	0.00	0.42	0.11	0.00	0.86
2.85	-1.19	-0.08	-0.03	-1.11	-1.35	0.08	0.18	1.15	1.43	1.01	1.50	1.83	1.25	1.36	1.16	0.05	0.07	N/A	-1.28	-1.26	-1.69	1.14	N/A	3.41	3.77	2.82	1.21	-0.03	-0.42	-0.25	-0.74	-0.05	1.35	N/A	1.45	N/A	-0.08
0.10	0.09	1.17	1.20	0.01	0.18	1.29	1.25	0.09	0.12	0.01	0.16	0.47	0.05	0.05	0.12	1.29	1.30	N/A	0.03	0.11	0.61	0.03	N/A	1.61	2.43	1.65	0.19	1.11	1.42	1.28	1.77	1.41	0.19	N/A	0.18	N/A	1.11
-1.26	-2.96	-1.24	1.21	-1.09	-1.82	1.24	1.53	1.45	1.76	2.34	3.00	5.23	1.71	1.81	-1.41	1.51	-1.91	2.74	1.68	1.94	-2.25	1.90	1.37	6.11	5.69	3.89	1.27	1.10	1.06	1.49	-2.74	-1.35	1.55	-1.10	4.05	4.32	-1.07
0.56	0.00	0.00	0.02	0.42	0.00	0.76	0.00	0.01	0.63	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.00	0.00	0.00	0.00	0.00	0.03	0.25	0.56	0.00	0.00	0.77	0.00	0.16	0.00	0.00	0.13
1.25	1.04	1.11	-1.10	-1.12	-1.13	1.19	1.51	1.17	1.41	1.38	-1.43	-1.46	-1.08	-1.20	-1.20	-1.09	-1.19	-1.20	-1.49	-1.36	-1.02	1.23	-1.18	-1.46	1.04	1.27	1.00	-1.04	1.26	-1.31	-1.52	-1.24	1.05	-1.02	1.06	-1.26	1.38
0.23	0.92	0.68	0.04	0.09	0.37	0.13	0.01	0.05	0.03	0.00	0.00	0.00	0.16	0.01	0.08	0.29	0.04	0.02	0.00	0.01	0.97	0.06	0.23	0.00	0.59	0.23	0.66	0.55	0.23	0.00	0.04	0.71	0.73	0.94	0.82	0.11	0.02
-0.01	-0.96	-0.06	0.05	-1.11	-1.48	1.22	1.52	1.31	1.59	1.86	0.79	1.89	0.32	0.31	-1.31	0.21	-1.55	0.77	0.10	0.29	-1.64	1.57	0.10	2.33	3.37	2.58	1.13	0.03	1.16	0.09	-2.13	-1.30	1.30	-1.06	2.56	1.53	0.16
1.26	2.00	1.18	1.16	0.02	0.35	0.03	0.01	0.14	0.17	0.48	2.22	3.35	1.40	1.51	0.11	1.30	0.36	1.97	1.59	1.65	0.62	0.34	1.28	3.79	2.33	1.31	0.14	1.07	0.10	1.40	0.61	0.06	0.25	0.04	1.50	2.79	1.23
1.43	-1.63	2.20	1.53	1.30	-1.11	1.77	-1.16	1.12	1.71	1.49	-1.38	-1.41	-1.28	-1.13	1.10	-1.19	-1.07	-1.01	-1.06	-1.46	1.07	1.54	-1.05	1.11	1.15	-1.32	1.10	-1.37	-1.65	-1.91	-1.12	-1.18	1.14	-1.04	1.22	-1.02	1.51
0.07	0.00	0.00	0.00	0.15	0.65	0.02	0.00	0.41	0.00	0.00	0.00	0.00	0.02	0.12	0.68	0.10	0.30	0.95	0.36	0.00	0.59	0.00	0.96	0.67	0.29	0.02	0.31	0.04	0.04	0.00	0.54	0.79	0.14	0.98	0.01	0.78	0.00
2.97	-1.30	2.07	1.39	1.56	1.77	1.69	-1.05	1.03	1.46	1.05	-1.81	-2.21	-1.27	-1.14	1.74	-1.52	-1.24	-1.39	-1.32	-1.62	1.43	1.36	1.09	-1.42	1.25	0.93	-1.21	-1.42	-1.39	-1.61	1.09	-1.26	-1.22	-0.90	-1.15	-1.63	×
0.00	0.11	0.01	0.00	0.06	0.02	0.02	0.38	0.46	0.05	0.46	0.00	0.00	0.05	0.30	0.03	0.03	0.05	0.20	0.01	0.00	0.11	0.01	x	0.00	0.67	0.18	0.04	0.06	0.24	0.02	0.53	0.70	0.16	0.56	0.01	0.01	×
2.20	-1.47	2.14	1.46	1.43	0.33	1.73	-1.11	1.08	1.59	1.27	-1.60	-1.81	-1.28	-1.14	1.42	-1.36	-1.16	-1.20	-1.19	-1.54	1.25	1.45	0.02	-0.16	1.20	-0.20	-0.05	-1.40	-1.52	-1.76	-0.02	-1.22	-0.04	-0.97	0.04	-1.33	N/A
0.77	0.17	0.07	0.07	0.13	1.44	0.04	0.05	0.05	0.13	0.22	0.21	0.40	0.01	0.01	0.32	0.16	0.09	0.19	0.13	0.08	0.18	0.09	1.07	1.27	0.05	1.13	1.16	0.02	0.13	0.15	1.11	0.04	1.18	0.07	1.19	0.31	N/A

SPO0296 SPO0297 SPO0298

SPO0294 SPO0295

SPO0288 SPO0289 SPO0290 SPO0291 SPO0292 SPO0293

SPO0286 SPO0287

SPO0281 SPO0282 SPO0283 SPO0284 SPO0285

SPO0280

SPO0273 SPO0274 SPO0275 SPO0276 SPO0277 SPO0278

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SPO0336	SPO0335	SPO0334	SPO0333	SPO0332	SPO0331	SPO0330	SPO0329	SPO0328	SPO0327	SPO0326	SPO0325	SPO0324	SPO0323	SPO0322	SPO0321	SPO0320	SPO0319	SPO0318	SPO0317	SPO0316	SPO0315	SPO0314	SPO0313	SPO0312	SPO0311	SPO0310	SPO0309	SPO0308	SPO0307	SPO0306	SPO0305	SPO0304	SPO0303	SPO0302	SPO0301	SPO0300	SPO0299
		lysA		argH		ccpA				phbA	phbB						ddsA	ispE				soxR		greA		moeA		mobB	mobA	fdhD							
hypothetical protein	hypothetical protein	diaminopimelate decarboxylase (EC:4.1.1.20)	lipoprotein	argininosuccinate lyase (EC:4.3.2.1)	thiol:disulfide interchange protein	cytochrome c peroxidase (EC:1.11.1.5)	hypothetical protein	hypothetical protein	diguanylate phosphodiesterase	acetyl-CoA acetyltransferase (EC:2.3.1.9)	acetoacetyl-CoA reductase (EC:1.1.1.36)	DNA-binding transcriptional activator GcvA	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	decaprenyl diphosphate synthase (EC:2.5.1.31)	+-шриоsриосуцаут-2-С-шенцут-2-етуштнот клиаse (ЕС:2.7.1.148)	A disheer heartidal 2 C mathal D cartherital Linese	electrotransfer ubiquinone oxidoreductase	hypothetical protein	redox-sensitive transcriptional activator SoxR	glyoxalase	transcription elongation factor GreA	hypothetical protein	molybdopterin biosynthesis protein MoeA	hypothetical protein	molybdopterin-guanine dinucleotide biosynthesis protein MobB	molybdoptern-guanne dinucleotide biosynthesis protein MobA	formate dehydrogenase accessory protein FdhD	AzIC family protein	lipoprotein	cytochrome c oxidase subunit IV	hypothetical protein	glycerophosphoryl diester phosphodiesterase	hypothetical protein	metallo-beta-lactamase
-1.25	1.02	-1.18	1.87	-2.20	1.11	-1.07	1.00	1.48	×	3.58	1.43	-1.12	-1.36	1.03	-1.37	-1.77	-1.15	-1.41	-1.33	1.48	-2.05	-1.46	-1.57	-1.42	1.25	-1.72	-1.08	1.18	1.14	1.04	-1.58	3.37	4.12	1.02	0.95	1.01	2.41
0.02	0.68	0.13	0.02	0.00	0.26	0.24	0.53	0.00	×	0.01	0.62	0.22	0.21	0.20	0.00	0.12	0.01	0.00	0.00	0.08	0.00	0.22	0.01	0.14	0.52	0.02	0.78	0.01	0.48	0.51	0.05	0.05	0.00	0.99	0.87	0.86	0.00
1.03	-1.19	1.13	1.46	1.49	-1.09	1.02	-1.67	-1.18	1.20	1.37	1.66	1.25	1.86	-1.16	-1.10	1.04	1.22	-1.22	-1.05	1.16	-1.42	-1.11	-1.38	1.00	1.31	-1.34	1.09	1.11	-1.21	-1.09	1.21	1.75	1.75	1.05	1.06	1.47	3.44
0.77	0.12	0.10	0.06	0.03	0.45	0.89	0.00	0.04	0.16	0.07	0.04	0.00	0.01	0.11	0.46	0.74	0.09	0.04	0.54	0.54	0.01	0.73	0.01	0.78	0.29	0.14	0.76	0.11	0.47	0.43	0.01	0.06	0.00	0.83	0.85	0.01	0.00
-0.11	-0.09	-0.03	1.67	-0.36	0.01	-0.03	-0.34	0.15	N/A	2.48	1.55	0.06	0.25	-0.06	-1.24	-0.37	0.04	-1.32	-1.19	1.32	-1.74	-1.29	-1.48	-0.21	1.28	-1.53	0.01	1.15	-0.04	-0.03	-0.19	2.56	2.94	1.04	1.00	1.24	2.93
1.14	1.11	1.16	0.21	1.85	1.10	1.05	1.34	1.33	N/A	1.11	0.12	1.19	1.61	1.10	0.14	1.41	1.19	0.10	0.14	0.16	0.32	0.17	0.10	1.21	0.03	0.19	1.09	0.03	1.18	1.07	1.40	0.81	1.19	0.02	0.06	0.23	0.52
-1.63	-2.41	4.21	2.06	-1.30	1.72	-1.85	-3.20	-2.23	х	2.90	0.87	1.20	-1.57	-1.62	1.68	-1.46	2.53	1.68	1.40	1.54	-3.08	-1.37	1.59	0.99	2.50	-1.81	1.02	1.34	5.00	1.21	1.16	4.35	3.56	-1.22	-1.23	-1.13	-1.29
0.00	0.00	0.00	0.00	0.02	0.00	0.00	0.00	0.00	×	0.00	0.02	0.00	0.55	0.00	0.00	0.16	0.00	0.00	0.01	0.08	0.00	0.65	0.01	0.06	0.04	0.00	0.77	0.00	0.00	0.00	0.09	0.00	0.00	0.51	0.77	0.74	0.61
-1.36	1.65	-1.64	-1.16	-1.23	-1.05	1.08	1.36	1.29	х	1.77	2.35	1.05	-1.46	2.18	1.28	-1.27	-1.35	1.15	1.22	1.00	1.81	-1.45	1.01	1.03	-1.30	1.04	1.02	-1.03	-1.19	-1.11	-1.25	1.28	1.35	1.24	1.16	-1.05	-1.00
0.01	0.00	0.00	0.27	0.09	0.52	0.55	0.01	0.06	×	0.00	0.00	0.40	0.14	0.00	0.07	0.07	0.01	0.11	0.06	0.68	0.00	0.15	0.96	0.51	0.30	0.95	0.98	0.78	0.44	0.08	0.03	0.46	0.00	0.29	0.43	0.19	0.82
-1.50	-0.38	1.29	0.45	-1.27	0.34	-0.39	-0.92	-0.47	N/A	2.34	1.61	1.13	-1.52	0.28	1.48	-1.37	0.59	1.42	1.31	1.27	-0.64	-1.41	1.30	1.01	0.60	-0.39	1.02	0.16	1.91	0.05	-0.05	2.82	2.46	0.01	-0.04	-1.09	-1.14
0.14	2.03	2.93	1.61	0.04	1.39	1.47	2.28	1.76	N/A	0.57	0.74	0.08	0.06	1.90	0.20	0.10	1.94	0.27	0.09	0.27	2.45	0.04	0.29	0.02	1.90	1.43	0.00	1.19	3.10	1.16	1.21	1.54	1.11	1.23	1.20	0.04	0.15
1.22	1.25	-1.92	1.46	1.52	-1.03	1.37	-1.12	1.38	-1.28	1.20	0.98	-1.68	1.28	-1.11	-1.41	1.04	-1.29	1.81	1.85	-1.24	1.41	1.25	-1.24	-1.68	-1.11	1.02	1.29	1.10	1.06	1.21	-1.15	1.12	1.07	1.02	1.41	1.51	1.20
0.01	0.04	0.00	0.03	0.00	0.80	0.00	0.11	0.01	0.28	0.03	0.33	0.00	0.09	0.42	0.04	0.50	0.00	0.00	0.00	0.22	0.01	0.18	0.03	0.00	0.58	0.99	0.32	0.06	0.83	0.00	0.31	0.80	0.77	0.92	0.11	0.01	0.06
-1.01	1.56	-2.48	1.18	1.09	-1.05	1.30	2.27	2.46	x	1.65	1.18	-2.06	-1.20	1.66	-1.36	-1.23	-1.59	1.32	1.59	1.24	1.76	1.32	-1.58	-1.34	-1.55	2.15	1.06	-1.24	1.16	1.05	-1.29	0.98	0.95	-1.17	1.62	1.30	1.65
0.74	0.00	0.00	0.28	0.32	0.92	0.15	0.00	0.00	x	0.00	0.76	x	0.73	0.04	0.13	0.33	0.05	0.01	0.01	0.59	0.01	0.14	0.00	0.00	0.14	0.02	0.74	0.10	0.55	0.69	0.04	0.68	0.07	0.59	0.07	0.01	0.00
0.11	1.41	-2.20	1.32	1.31	-1.04	1.34	0.58	1.92	N/A	1.43		-1.87	0.04	0.28		-0.10		1.57	1.72	0.00	1.59	1.29	-1.41	-1.51	-1.33	1.59	1.18	-0.07	1.11	1.13	-1.22	1.05	1.01	-0.08	1.52	1.41	1.43
1.12	0.16			0.21	0.01	0.04	1.70	0.54	N/A	0.23		0.19	1.24	1.39	0.02	1.14	0.15	0.25	0.13	1.24	0.17				0.22	0.57	0.12	1.17	0.05	0.08	0.07	0.07	_	1.10			0.23

## Chapter 10: Appendix

SPO0374	SPO0373	SPO0372	SPO0371	SPO0370	SPO0369	SPO0368	SPO0367	SPO0366	SPO0365	SPO0364	SPO0363	SPO0362	SPO0361	SPO0360	SPO0359	SPO0358	SPO0357	SPO0356	SPO0355	SPO0354	SPO0353	SPO0352	SPO0351	SPO0350	SPO0349	SPO0348	SPO0347	SPO0346	SPO0345	SPO0344	SPO0343	SPO0342	SPO0341	SPO0340	SPO0339	SPO0338	SPO0337
			luxR-1	ccrA				deoD					sdhB	sdhA	sdhD	sdhC						citE			mdh		sucC	sucD		sucA	sucB			lpdA			fisE
hypothetical protein	helicase, ATP-dependent	autoinducer synthesis protein	LuxR family transcriptional regulator	crotonyl-CoA reductase	acyltransferase	methylmalonyl-CoA mutase (EC:5.4.99.2)	H-NS family DNA-binding protein	purine nucleoside phosphorylase (EC:2.4.2.1)	hypothetical protein	decarboxylase, pyridoxal-dependent	hypothetical protein	type I secretion target repeat-containing protein	Succritate denydrogenase fron-sultur subunit (EC:1.3.99.1)	(EC:1.3.5.1)	anchor protein (EC:1.3.99.1)	succinate dehydrogenase, cytochrome b556 subunit (EC:1.3.99.1)	von Willebrand factor A	hypothetical protein	hypothetical protein	hypothetical protein	NnrU family protein	citrate lyase subunit beta (EC:4.1.3.34 4.1.3.6)	hypothetical protein	hypothetical protein	malate dehydrogenase (EC:1.1.1.37)	hypothetical protein	succinyl-CoA synthetase subunit beta (EC:6.2.1.5)	succinyl-CoA synthetase subunit alpha (EC:6.2.1.5)	hypothetical protein	2-oxoglutarate dehydrogenase E1 (EC:1.2.4.2)	dihydrolipoamide succinyltransferase (EC:2.3.1.61)	hypothetical protein	hypothetical protein	2-oxoglutarate dehydrogenase E3 (EC:1.8.1.4)	acyltransferase	cell division permease FtsX	cell division ATP-binding protein FtsE
1.06	-1.70	-1.72	-1.80	-3.94	-1.36	-1.59	-1.48	-1.36	3.24	-0.96	3.14	-1.35	3.05	3.22	3.31	3.07	-1.38	1.06	1.66	1.45	1.11	-1.74	1.42	2.01	2.67	-1.88	1.72	1.37	1.29	1.37	1.54	-2.62	-1.10	1.18	1.21	-1.33	1.08
0.98	0.00	0.00	0.26	0.00	0.55	0.09	0.00	0.19	0.02	0.45	0.00	0.00	0.06	0.00	0.00	0.00	0.34	0.92	0.00	0.43	0.90	0.00	0.00	0.00	0.00	0.00	0.03	0.02	0.68	0.01	0.50	0.00	0.70	0.80	0.09	0.01	0.11
1.13	-1.17	2.47	3.01	1.58	-1.10	1.29	-1.12	1.31	1.59	-1.02	3.76	1.47	2.07	1.24	1.85	1.40	1.03	1.20	1.65	1.21	1.15	1.39	1.18	1.73	1.52	-1.15	1.44	2.04	2.04	1.45	1.79	1.39	1.64	1.53	1.41	1.26	1.21
0.68	0.46	0.01	0.01	0.04	0.90	0.09	0.42	0.05	0.03	1.00	0.00	0.05	0.01	0.34	0.00	0.01	0.88	0.27	0.01	0.19	0.10	0.04	0.34	0.00	0.03	0.31	0.05	0.00	0.00	0.01	0.00	0.05	0.06	0.00	0.04	0.06	0.02
1.10	-1.44	0.38	0.61	-1.18	-1.23	-0.15	-1.30	-0.03	2.42	-0.99	3.45	0.06	2.56	2.23	2.58	2.24	-0.18	1.13	1.66	1.33	1.13	-0.18	1.30	1.87	2.10	-1.52	1.58	1.71	1.67	1.41	1.67	-0.62	0.27	1.36	1.31	-0.04	1.15
0.03	0.27	2.10	2.41	2.76	0.13	1.44	0.18	1.34	0.83	0.03	0.31	1.41	0.49	0.99	0.73	0.84	1.21	0.07	0.01	0.12	0.02	1.57	0.12	0.14	0.58	0.37	0.14	0.33	0.38	0.04	0.13	2.01	1.37	0.18	0.10	1.30	0.06
1.43	2.52	-3.54	-5.51	-1.51	-1.01	-1.36	1.74	1.94	1.35	-1.20	12.90	1.11	5.68	4.82	4.69	4.46	-2.23	1.74	1.47	1.39	1.86	2.03	-1.40	1.28	5.12	1.26	7.46	6.69	5.93	4.70	6.11	1.10	1.64	6.21	2.55	1.51	2.47
0.37	0.05	0.00	0.01	0.02	0.72	0.12	0.00	0.06	0.16	0.75	0.00	0.88	0.00	0.00	0.00	0.00	0.00	0.05	0.00	0.54	0.00	0.00	0.74	0.00	0.00	0.01	0.00	0.00	0.00	0.00	0.02	0.07	0.01	0.00	0.00	0.00	0.00
1.03	1.00	4.09	5.46	1.46	1.05	1.25	-1.09	-1.16	-1.07	-1.45	4.58	1.61	2.29	2.43	2.38	1.94	1.39	1.15	-1.19	1.18	0.99	1.04	2.69	-0.97	1.07	-1.06	4.59	3.45	4.38	3.44	3.66	3.15	2.01	3.31	-1.25	-1.33	-1.29
0.98	0.86	0.00	0.00	0.02	0.95	0.23	0.06	0.11	0.84	0.32	0.00	0.00	0.01	0.00	0.00	0.00	0.15	0.24	0.02	0.13	0.38	0.78	0.00	0.60	0.89	0.27	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.00	0.05	0.29	0.01
1.23	1.76	0.28	-0.02	-0.03	0.02	-0.06	0.33	0.39	0.14	-1.33	8.74	1.36	3.99	3.63	3.54	3.20	-0.42	1.45	0.14	1.29	1.43	1.54	0.65	0.15	3.10	0.10	6.03	5.07	5.16	4.07	4.89	2.13	1.83	4.76	0.65	0.09	0.59
0.20	0.76	3.82	5.49	1.49	1.03	1.31	1.42	1.55	1.21	0.13	4.16	0.25	1.70	1.20	1.16	1.26	1.81	0.30	1.33	0.11	0.43	0.50	2.05	1.13	2.03	1.16	1.44	1.62	0.78	0.63	1.23	1.03	0.19	1.45	1.90	1.42	1.88
1.41	-1.03	-1.14	1.13	1.04	-1.03	-1.12	0.99	1.60	1.19	-1.80	-1.65	-1.51	-1.12	-1.20	-1.18	-1.04	-1.10	1.26	-1.01	-1.30	-1.36	-1.36	-0.98	-0.99	1.24	1.37	-1.42	1.19	-1.10	-1.59	-1.04	-1.25	-1.35	-1.25	1.52	-1.06	-1.14
0.17	0.82	0.56	0.63	0.94	0.99	0.35	0.19	0.01	0.25	0.03	0.02	0.01	0.57	0.02	0.00	0.43	0.65	0.05	0.98	0.00	0.03	0.02	0.55	0.73	0.12	0.00	0.00	0.08	0.03	0.00	0.33	0.09	0.04	0.06	0.01	0.61	0.11
1.05	-1.17	-0.89	1.73	1.23	2.41	-1.41	1.05	1.10	-1.44	-1.96	-1.09	-1.45	-1.31	0.94	-1.20	1.09	-1.20	-1.06	-1.49	-1.19	-1.42	-1.52	-1.03	-1.60	0.91	-1.15	0.89	1.20	1.04	-1.55	-1.15	1.77	1.69	0.90	-1.00	-1.15	-1.66
0.93	0.33	0.59	0.10	0.33	0.11	0.04	0.53	0.84	0.54	0.03	0.45	0.03	0.09	0.03	0.04	0.69	0.43	0.44	0.02	0.02	0.00	0.00	0.81	0.06	0.02	0.23	0.01	0.63	0.53	0.00	0.08	0.05	x	0.05	0.98	0.76	0.00
1.23	-1.10	-1.01	1.43	1.14	0.69	-1.27	1.02	1.35	-0.13	-1.88	-1.37	-1.48	-1.22	-0.13	-1.19	0.03	-1.15	0.10	-1.25	-1.25	-1.39	-1.44	-1.00	-1.30	1.07	0.11	-0.27	1.20	-0.03	-1.57	-1.10	0.26	0.17	-0.18	0.26	-1.11	-1.40
0.18	0.07	0.13	0.30	0.10	1.72	0.14	0.03	0.25	1.32	0.08	0.28	0.03	0.10	1.07	0.01	1.07	0.05	1.16	0.24	0.06	0.03	0.08	0.03	0.30	0.17	1.26	1.16	0.01	1.07	0.02	0.05	1.51	1.52	1.08	1.26	0.04	0.26

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## Chapter 10: Appendix

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	dut	coaBC		rpoH2				comM	gshB				glnD	mviN				trpS	bphC3																
molvhdonterin hiosynthesis protein MoeB	скоху или не 5 - и приозрнате пистеот получнотазе (EC:3.6.1.23)	ortunctional puospinopantothenate synthase decarboxylase/phosphopantothenate synthase (EC:4.1.1.36 6.3.2.5)	hypothetical protein	RNA polymerase factor sigma-32	guanyltransferase	phosphoglycerate mutase	hypothetical protein	competence protein ComM	glutathione synthetase (EC:6.3.2.3)	hypothetical protein	tetrapyrrole methylase	hypothetical protein	PII uridylyl-transferase (EC:2.7.7.59)	integral membrane protein MviN	cyclic nucleotide-binding protein	hypothetical protein	rhomboid family protein	tryptophanyl-tRNA synthetase (EC:6.1.1.2)	bipnenyi-2,3-dioi 1,2-dioxygenase III (EC:1.13.11.39)	dehydrogenase	AsnC family transcriptional regulator	aminotransferase	hypothetical protein	universal stress family protein	2-hydroxychromene-2-carboxylate isomerase	hypothetical protein	ACP phosphodieterase	NifU domain-containing protein	protease	ribosomal-protein-alanine acetyltransferase	sugar ABC transporter substrate-binding protein	sugar ABC transporter ATP-binding protein	sugar ABC transporter permease	sugar ABC transporter permease	hypothetical protein
-1 61	1.06	-1.21	1.20	1.08	-1.02	-1.07	2.43	1.47	1.62	-1.00	1.06	-1.32	-1.24	1.10	-1.28	1.97	-1.03	1.18	1.45	-1.15	-1.74	1.09	-1.89	1.39	1.57	-2.48	-1.45	0.92	1.01	-1.72	-1.34	-1.37	-1.45	-1.01	-1.02
96.0	1.00	0.43	0.92	0.16	0.89	0.25	0.00	0.30	0.13	0.43	0.37	0.24	0.07	0.46	0.49	0.00	0.13	0.78	0.05	0.76	0.16	0.98	0.00	0.10	0.00	0.00	0.23	0.08	0.95	0.00	0.00	0.41	0.00	0.86	0.87
1.12	1.38	1.08	-1.07	-2.19	1.07	1.39	2.90	2.36	1.13	1.63	1.75	1.06	1.17	1.18	1.09	-1.10	1.03	1.09	1.48	1.36	-1.36	-1.54	-1.39	1.12	1.11	-1.22	-1.25	1.29	1.03	-1.19	-1.63	-1.18	-1.11	1.09	-1.16
045	0.08	0.68	0.67	0.00	0.56	0.01	0.00	0.00	0.35	0.00	0.00	0.84	0.24	0.53	0.72	0.41	0.96	0.67	0.01	0.43	0.03	0.02	0.22	0.32	0.52	0.30	0.56	0.08	0.90	0.42	0.00	0.25	0.14	0.52	0.37
-0.25	1.22	-0.06	0.06	-0.56	0.03	0.16	2.67	1.92	1.38	0.32	1.41	-0.13	-0.04	1.14	-0.10	0.44	0.00	1.14	1.47	0.11	-1.55	-0.23	-1.64	1.26	1.34	-1.85	-1.35	1.11	1.02	-1.46	-1.49	-1.28	-1.28	0.04	-1.09
1 37	0.16	1.15	1.14	1.64	1.05	1.23	0.24	0.44	0.25	1.31	0.35	1.19	1.21	0.04	1.19	1.54	1.03	0.04	0.02	1.26	0.19	1.32	0.25	0.14	0.23	0.63	0.10	0.18	0.01	0.26	0.15	0.10	0.17	1.05	0.07
-1.33	1.65	1.12	1.26	0.90	1.44	4.29	4.44	1.23	2.81	1.31	1.34	1.51	1.26	1.52	1.15	-1.29	1.43	1.29	1.19	0.84	-2.40	-1.35	-2.07	1.32	1.55	-1.08	-2.19	0.84	1.33	-1.86	-1.55	-1.48	-1.35	1.03	1.58
0 69 0	0.47	0.91	0.85	0.02	0.00	0.00	0.00	0.78	0.00	0.01	0.01	0.02	0.04	0.04	0.63	0.11	0.00	0.59	0.09	0.62	0.05	0.00	0.00	0.10	0.00	0.87	0.00	0.00	0.66	0.00	0.00	0.00	0.01	0.74	0.28
-1.34	1.04	1.19	1.34	1.02	1.16	1.06	1.13	1.97	-1.23	1.38	-1.02	1.24	1.07	-1.02	-0.99	-1.10	-1.27	-1.52	-1.14	-1.61	-1.07	1.02	-1.11	1.19	1.12	1.30	-1.24	1.03	-1.07	1.03	-1.47	-1.49	-1.63	-1.71	-1.03
0 03	0.85	0.39	0.18	0.96	0.10	0.38	0.11	0.00	0.01	0.14	0.73	0.34	0.71	0.81	0.99	0.45	0.04	0.01	0.36	0.19	0.65	0.77	0.67	0.14	0.20	0.06	0.54	0.66	0.85	0.88	0.00	0.01	0.00	0.01	0.77
-1 34	1.35	1.16	1.30	0.96	1.30	2.68	2.79	1.60	0.79	1.35	0.16	1.38	1.17	0.25	0.08	-1.20	0.08	-0.12	0.03	-0.39	-1.74	-0.17	-1.59	1.26	1.34	0.11	-1.72	0.94	0.13	-0.42	-1.51	-1.49	-1.49	-0.34	0.28
0.01	0.31	0.03	0.04	0.06	0.14	1.62	1.66	0.37	2.02	0.03	1.18	0.14	0.10	1.27	1.07	0.10	1.35	1.41	1.17	1.22	0.67	1.19	0.48	0.07	0.22	1.19	0.48	0.09	1.20	1.45	0.04	0.01	0.14	1.37	1.31
1 08	1.45	1.14	1.37	1.21	-1.49	-1.17	1.51	-1.02	1.13	-1.08	1.30	1.19	1.10	1.40	1.41	-1.07	1.29	-1.63	-1.29	1.24	-1.40	-1.05	-1.21	1.27	1.25	-1.28	1.34	1.30	-1.01	-1.44	-1.02	-1.09	-1.07	1.15	-1.04
031	0.01	0.34	0.13	0.02	0.00	0.02	0.00	0.96	0.34	0.48	0.00	0.36	0.43	0.00	0.02	0.81	0.00	0.02	0.02	0.41	0.01	0.50	0.14	0.04	0.01	0.06	0.39	0.00	0.97	0.03	0.43	0.31	0.10	0.27	0.75
-1 48	1.03	1.22	1.15	-1.02	-1.84	-1.73	-0.94	1.18	-1.21	-1.09	-0.94	1.17	-1.11	-0.93	1.12	2.43	1.25	-2.24	-1.53	1.56	1.13	-1.57	-1.06	1.31	-1.14	-1.30	2.63	1.10	1.35	1.60	1.20	1.26	1.01	1.01	-1.05
0.02	0.72	0.28	0.40	0.85	0.00	0.00	0.19	0.58	0.05	0.92	0.22	0.68	0.66	0.13	0.70	0.00	0.34	0.00	0.05	0.30	0.22	0.00	0.98	0.03	0.11	0.07	0.02	0.75	0.45	0.01	0.23	0.19	0.99	0.92	0.75
-0.20	1.24	1.18	1.26	0.10	-1.67	-1.45	0.29	0.08	-0.04	-1.09	0.18	1.18	-0.01	0.24	1.27	0.68	1.27	-1.94	-1.41	1.40	-0.14	-1.31	-1.14	1.29	0.06	-1.29	1.99	1.20	0.17	0.08	0.09	0.09	-0.03	1.08	-1.05
1.78	0.21	0.04	0.11	1.12	0.18	0.28	1.22	1.10	1.17	0.01	1.12	0.01	1.11	1.16	0.14	1.75	0.02	0.31	0.12	0.16	1.27	0.26	0.08	0.02	1.20	0.01	0.65	0.10	1.18	1.52	1.11	1.18	1.04	0.07	0.01

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SPO0390

SPO0409 SPO0410 SPO0408

SPO0405 SPO0406 SPO0407

SPO0449	SPO0448	SPO0447	SPO0446	SPO0445	SPO0444	SPO0443	SPO0442	SPO0441	SPO0440	SPO0439	SPO0438	SPO0437	SPO0436	SPO0435	SPO0434	SPO0433	SPO0432	SPO0431	SPO0430	SPO0429	SPO0428	SPO0427	SPO0426	SPO0425	SPO0424	SPO0423	SPO0422	SPO0421	SPO0420	SPO0419	SPO0418	SPO0417	SPO0416	SPO0415	SPO0414	SPO0413	SPO0412	SPO0411
nadD																			gltX				nadE				leuA		mreB	mre C		pbpA	mreD					
nicotinic acid mononucleotide adenylyltransferase	MOSC domain-containing protein	hypothetical protein	ABC transporter ATP-binding protein	hypothetical protein	hypothetical protein	hypothetical protein	thioredoxin	hypothetical protein	thioesterase	hypothetical protein	ErfK/YbiS/YcfS/YnhG family protein	cold shock family protein	hypothetical protein	acyl-CoA thioesterase	ABC transporter ATP-binding protein	ABC transporter permease	rrf2 family protein	hypothetical protein	glutamyl-tRNA synthetase (EC:6.1.1.17)	metallo-beta-lactamase	amino acid deaminase	hypothetical protein	NAD synthetase (EC:6.3.1.5)	MORN repeat-containing protein	hypothetical protein	DNA-binding protein	2-isopropylmalate synthase (EC:2.3.3.13)	hypothetical protein	rod shape-determining protein MreB	rod shape-determining protein MreC	hypothetical protein	penicillin-binding protein 2	rod shape-determining protein MreD	D-isomer specific 2-hydroxyacid dehydrogenase	peptidyl-dipeptidase	Cystoret metaconsm rlr-dependent enzyme family protein	hypothetical protein	hypothetical protein
-1.67	1.32	х	1.62	1.09	1.02	-1.03	1.08	1.22	-1.27	1.15	1.17	-1.46	1.10	1.21	-1.05	-0.97	1.07	1.14	-1.02	1.46	-1.26	1.88	1.85	0.94	-2.05	-1.71	1.54	-1.63	-1.40	-1.07	-1.10	-1.15	1.36	-1.04	1.44	1.29	-1.00	-1.80
0.48	0.92	×	0.00	0.51	0.51	0.89	0.23	0.39	0.01	0.37	0.23	0.00	0.38	0.67	0.08	0.88	0.46	0.33	х	0.01	0.33	0.00	0.00	0.00	0.02	0.04	0.00	0.01	0.00	0.21	0.80	0.01	0.15	0.80	0.42	0.02	0.62	0.01
1.15	1.15	-1.31	0.98	1.12	-1.12	-1.16	-1.23	-1.02	1.50	-1.09	-1.05	1.27	1.23	1.07	-1.42	-1.21	1.36	1.42	1.18	-1.01	1.65	2.21	1.29	1.02	1.03	1.10	1.12	1.23	-1.26	1.18	1.51	1.04	1.19	-1.20	1.09	-1.66	1.08	-1.04
0.78	0.74	0.03	0.87	0.67	0.52	0.66	0.17	0.89	0.02	0.40	0.04	0.10	0.00	0.44	0.00	0.36	0.03	0.46	0.33	0.99	0.00	0.00	0.17	1.00	0.80	0.70	0.24	0.22	0.02	0.14	0.00	0.77	0.31	0.49	0.66	0.00	0.21	0.55
-0.26	1.24	N/A	1.30	1.11	-0.05	-1.10	-0.08	0.10	0.12	0.03	0.06	-0.10	1.17	1.14	-1.24	-1.09	1.22	1.28	0.08	0.23	0.20	2.05	1.57	0.98	-0.51	-0.31	1.33	-0.20	-1.33	0.05	0.21	-0.05	1.28	-1.12	1.27	-0.19	0.04	-1.42
1.41	0.09	N/A	0.32	0.02	1.07	0.06	1.16	1.12	1.39	1.12	1.11	1.37	0.06	0.07	0.19	0.12	0.15	0.14	1.10	1.24	1.46	0.17	0.28	0.04	1.54	1.41	0.21	1.43	0.07	1.13	1.31	1.10	0.09	0.08	0.18	1.48	1.04	0.38
1.84	1.42	-1.00	2.22	-2.03	-1.88	-1.13	-1.22	1.15	-1.20	-1.61	-2.54	1.65	-1.09	1.36	1.35	1.53	7.87	1.93	5.68	-1.10	-1.10	-1.74	3.15	0.94	-3.83	1.10	3.69	-1.28	3.51	1.28	1.17	1.14	1.72	1.12	1.77	-2.02	3.60	-1.63
0.39	0.86	0.01	0.00	0.05	0.00	0.69	0.58	0.50	0.46	0.03	0.00	0.00	0.96	0.01	0.03	0.18	0.00	0.17	0.00	0.61	0.86	0.00	0.00	0.00	0.00	0.73	0.00	0.29	0.00	0.02	0.10	0.01	0.04	0.63	0.13	0.00	0.00	0.00
0.99	-1.14	-1.30	-1.33	-1.02	1.03	-1.30	-1.52	-1.34	-1.29	-1.14	1.07	-1.17	-1.03	1.10	-1.64	-1.60	1.05	1.09	-1.31	2.99	-1.11	2.37	-1.03	1.23	-1.27	-1.39	-1.09	1.04	-1.27	-1.14	-1.02	-1.08	-1.11	1.02	-1.11	2.13	1.43	1.02
0.93	0.78	0.05	0.07	0.95	0.99	0.34	0.01	0.22	0.36	0.19	0.83	0.17	0.71	0.20	0.00	0.05	0.63	0.87	0.01	0.00	0.08	0.00	0.51	0.15	0.20	0.23	0.03	0.76	0.00	0.18	0.48	0.14	0.33	0.86	0.10	0.00	0.00	0.84
3 1.41	0.14	-1.15	0.45	-1.53	-0.43	-1.22	-1.37	-0.10		-1.38	-0.74		-1.06	) 1.23	0.15	-0.04	3 4.46	1.51	2.19	0.95	-1.11	0.32	1.06	5 1.09		-0.15		-0.12	) 1.12	3 0.07	3 0.08	4 0.03	3 0.31	5 1.07	0.33	0.05		-0.31
0.43	4 1.28	5 0.15	5 1.78	3 0.51	3 1.46	2 0.09	7 0.15	0 1.25	5 0.05	8 0.24	4 1.81	4 1.41	6 0.03	3 0.13	5 1.50	4 1.57	5 3.41	0.42	€ 3.50	5 2.05	0.01	2 2.06	5 2.09	€ 0.14	5 1.28	5 1.25	) 2.39	2 1.16	2 2.39	7 1.21	3 1.10	3 1.11	1 1.42	0.05	3 1.44	5 2.08		1 1.33
43 1.	-																																					
11 0.	1.14 0.	1.46 0.	1.40 0.	1.17 0.	-1.10 0.	1.22 0.	1.13 0.	1.12 0.	1.18 0.	1.84 0.	0.99 0.	1.42 0.	1.32 0.	1.41 0.	1.53 0.	0.98 0.	1.31 0.	1.32 0.	-1.42 0.	1.60 0.	1.10 0.	-1.02 0.	-1.14 0.	1.10 0.	1.32 0.	-1.29 0.	-1.10 0.	-1.26 0.	-1.54 0.	1.08 0.	1.19 0.	1.01 0.	-1.08 0.	-1.47 0.	1.92 0.		1.79 0.	
0.83 -]	0.76 1	0.00	0.01 -1	0.64	0.41 -1	0.51 1	0.06 -(	0.15 1	0.07 -1	0.00 -1	0.67 1	0.11 1	0.01 1	0.00 1	0.00 1	0.69 -1		0.61 -1	0.00 -1	0.01 1	0.37 -1	0.68 -1	0.38 -]	0.17 1	0.07 -(		0.33 -]	0.07 -1	0.00 - ]	0.19 -1	0.02 -1	0.50 - ]	0.55 -1	0.04 -1	0.00 1	0.00 1		0.41 1
1.27	1.59	Х	-1.50	Х	1.17	1.88	-0.92	1.29	-1.10	-1.55	1.15	1.20	1.12	1.15	1.32	-1.00	-1.36 (	-1.52	-1.61	1.21	-1.05		_			-0.91 (				-1.17 (		-1.33	-1.43	-1.25	1.58 (	1.27	1.22	1.29
0.56 -	0.44	×	0.01 -	x	0.56 -	0.16	0.11	0.13	0.92	0.06	0.28	0.78	0.11	0.05	0.02	0.92 -		0.51 -	0.01 -	0.38	0.76 -	0.03 -	0.00 -	0.71	0.57 -		0.00 -	0.08 -	0.00 -	0.12 -	0.01 -	0.00 -	0.04 -	0.16 -	0.00	0.07	0.00	0.01
-0.08	1.37	N/A	-1.45	N/A	-1.14	1.55	0.11	0.09	0.04	-1.70	1.07	1.31	1.22	1.28	1.43	-0.99	-0.03	-1.42	-1.52	1.41	-1.08	-1.17	-1.32	1.09	-1.16	-1.10	-1.25	-1.32	-1.51	-0.04	-0.06	-1.17	-1.26	-1.36	1.75	1.25	1.51	1.18
1.19	0.23	N/A	0.05	N/A	0.03	0.33	1.02	1.21	1.14	0.15	0.08	0.11	0.10	0.13	0.11	0.01	1.34	0.10	0.10	0.20	0.03	0.14	0.18	0.02	0.16	0.19	0.15	0.06	0.03	1.13	1.25	0.16	0.18	0.11	0.17	0.02	0.29	0.12

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# Chapter 10: Appendix

#### 2012

## M. Kirkwood

## Chapter 10: Appendix

rpsC	rplV	rpsS	rplB							phnM		phnN	phnL	phnK	phnJ	phnI	phnH	phnG													lysK		dddW		dacB		
30S ribosomal protein S3	50S ribosomal protein L22	30S ribosomal protein S19	50S ribosomal protein L2	microcystin dependent protein	NADH-ubiquinone oxidoreductase	acetyltransferase	ISSpo8, transposase	hypothetical protein	hypothetical protein	alkylphosphonate utilization protein PhnM	hypothetical protein	alkylphosphonate utilization protein PhnN	alkylphosphonate utilization protein PhnL	phosphonate C-P lyase system protein Phn	alkylphosphonate utilization protein PhnJ	alkylphosphonate utilization protein PhnI	carbon-phosphorus lyase complex subunit	alkylphosphonate utilization protein PhnG	GntR family transcriptional regulator	formate/nitrate transporter	pyridoxamine 5'-phosphate oxidase	hypothetical protein	universal stress family protein	antibiotic biosynthesis monooxygenase domain- containing protein	ParA family protein	hypothetical protein	lysyl-tRNA synthetase (EC:6.1.1.6)	LysR family transcriptional regulator	dimethylsulfoniopropionate lyase	hypothetical protein	alanine-endopeptidase (EC:3.4.16.4)	hypothetical protein D aland D alania carbox mentidace/D alany! D	(EC:2.7.7.18)				
-1.96	-1.42	-1.49	-1.61	-1.30	1.33	Х	1.08	1.92	1.05	-1.03	-1.25	1.19	-0.99	1.49	1.36	1.21	1.03	1.32	1.19	-1.10	Х	1.79	1.49	0.99	2.60	-1.06	1.28	-1.82	1.51	1.71	1.51	3.30	36.60	4.61	-1.49	1.17	
0.00	0.00	0.00	0.00	0.00	0.74	×	0.11	0.00	0.07	0.98	0.11	0.91	0.92	0.15	0.31	0.22	0.99	0.90	0.28	0.05	×	0.00	0.01	0.00	0.05	0.20	0.03	0.12	0.00	0.00	0.47	0.00	0.00	0.00	0.12	0.27	
-1.15	-1.17	-1.65	-1.48	-1.83	1.23	1.26	1.99	-1.24	1.26	1.04	1.13	1.20	1.17	1.14	1.12	-1.03	-1.12	-1.01	-1.06	-1.35	1.13	1.18	1.32	1.10	2.78	1.28	1.32	-1.00	1.01	1.12	1.34	2.09	45.60	1.98	1.11	1.46	
0.26	0.15	0.00	0.00	0.00	0.04	0.12	0.00	0.15	0.10	0.86	0.52	0.34	0.12	0.09	0.59	0.85	0.24	0.98	0.90	0.15	0.10	0.10	0.03	0.22	0.00	0.46	0.05	0.94	0.97	0.53	0.30	0.00	0.00	0.01	0.45	0.17	
-1.56	-1.30	-1.57	-1.55	-1.57	1.28	N/A	1.54	0.34	1.16	0.01	-0.06	1.20	0.09	1.32	1.24	0.09	-0.05	0.16	0.06	-1.23	N/A	1.49	1.41	1.05	2.69	0.11	1.30	-1.41	1.26	1.42	1.43	2.70	41.10	3.30	-0.19	1.32	
0.41	0.13	0.08	0.07	0.27	0.05	N/A	0.46	1.58	0.11	1.04	1.19	0.01	1.08	0.18	0.12	1.12	1.08	1.17	1.13	0.13	N/A	0.31	0.09	0.05	0.09	1.17	0.02	0.41	0.25	0.29	0.09	0.61	4.50	1.32	1.30	0.15	_
11.20	13.20	11.80	11.70	-3.98	2.28	4.51	-1.15	-1.27	-3.09	-1.32	1.51	0.99	-1.66	-1.41	-1.61	-2.15	-2.40	1.27	1.36	1.69	х	1.46	3.25	-3.75	5.50	-7.99	-1.08	-1.84	-1.78	-1.44	3.38	-0.95	2.66	2.57	-1.98	1.84	
0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.40	0.07	0.00	0.80	0.00	0.98	0.16	0.46	0.14	0.01	0.00	0.88	0.65	0.02	x	0.00	0.00	0.00	0.00	0.00	0.73	0.53	0.00	0.00	0.07	0.27	0.01	0.01	0.07	0.09	
-1.98	-1.78	-1.70	-1.62	1.12	0.99	-1.28	×	-1.74	1.24	-1.29	-1.55	-0.99	-1.13	-1.11	-1.22	1.08	-1.15	1.03	-1.13	1.04	1.28	1.24	1.02	1.19	-1.46	1.24	3.15	2.07	1.69	1.39	-1.68	-1.22	-1.12	1.24	1.21	-1.40	
0.00	0.00	0.00	0.01	0.45	0.18	0.04	×	0.00	0.05	0.30	0.09	0.93	0.47	0.26	0.52	0.49	0.49	0.96	0.76	0.87	x	0.11	0.95	0.03	0.00	х	0.00	0.03	0.01	0.00	0.01	0.14	0.34	0.38	0.14	0.05	
4.61	5.71	5.05	5.04	-1.43	1.63	1.62	N/A	-1.51	-0.93	-1.31	-0.02	0.00	-1.40	-1.26	-1.42	-0.54	-1.78	1.15	0.12	1.37	N/A	1.35	2.14	-1.28	2.02	-3.38	1.04	0.12	-0.05	-0.03	0.85	-1.08	0.77	1.91	-0.39	0.22	
6.59	7.49	6.75	6.66	2.55	0.65	2.90	N/A	0.24	2.17	0.02	1.53	0.99	0.27	0.15	0.20	1.62	0.63	0.12	1.25	0.33	N/A	0.11	1.12	2.47	3.48	4.62	2.12	1.96	1.74	1.42	2.53	0.14	1.89	0.67	1.60	1.62	
-1.09	-1.17	-2.02	-2.07	1.62	1.19	1.13	-1.07	-1.37	-2.25	-0.97	-1.32	1.07	-0.97	-1.10	1.08	-1.59	-1.46	-1.08	-1.21	-1.08	1.14	1.12	1.51	1.15	1.40	-1.46	-1.16	-1.46	-1.74	-1.23	-1.39	1.04	-1.72	1.24	1.18	-1.33	
0.00	0.02	0.00	0.00	0.00	0.25	0.08	0.78	0.00	0.00	0.81	0.27	0.62	0.69	0.87	0.76	0.00	0.05	0.88	0.56	0.65	0.06	0.18	0.01	0.14	0.03	0.07	0.28	0.23		0.00	0.02	0.76	0.00	0.10	0.06	0.03	
-2.07	-2.35	-3.71	-3.55	3.22	1.02	-1.48	-1.60	-1.41	-1.54	-1.00	-1.49	1.13	-1.54	-1.39	-1.11	-1.76	-1.18	1.61	-1.19	1.31	-1.17	1.34	1.13	1.15	1.21	х	-2.14	1.17	-1.10	-1.17	-1.66	1.08	х	1.07	1.29	-1.81	
0.00	0.00	0.00	0.00	0.00	0.15	0.01	×	0.09	0.03	0.71	0.11	0.72	0.58	0.03	0.96	0.00	0.76	0.42	0.86	0.26	×	0.17	0.17	0.06	0.01	x	0.00	0.12	0.51	0.01	0.00	0.51	x	0.94	0.01	0.00	
-1.58	-1.76	-2.87	-2.81	2.42	1.11	-0.18	-1.34	-1.39	-1.90	-0.98	-1.41	1.10	-1.26	-1.25	-0.02	-1.68	-1.32	0.27	-1.20	0.12	-0.02	1.23	1.32	1.15	1.31	N/A	-1.65	-0.15	-1.42	-1.20	-1.53	1.06	N/A	1.16	1.24	-1.57	
0.49	0.59	0.85	0.74	0.80	0.09	1.31	0.27	0.02	0.36	0.02	0.09	0.03	0.28	0.14	1.10	0.09	0.14	1.35	0.01	1.20	1.16	0.11	0.19	0.00	0.10	N/A	0.49	1.32	0.32	0.03	0.14	0.02	N/A	0.09	0.06	0.24	

SPO0484 SPO0485 SPO0486 SPO0487

SPO0483 SPO0482 SPO0481

SPO0475 SPO0476 SPO0477 SPO0478 SPO0479

SPO0474 SPO0473

SP00468 SP00469 SP00470 SP00471 SP00472

SPO0467 SPO0466 SPO0465

SPO0462 SPO0463 SPO0464

SPO0460 SPO0461

SPO0458 SPO0457

SPO0459

SP00452 SP00453 SP00454 SP00455 SP00456

SPO0451 SPO0450

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								rluC				rplQ	rpoA	rpsK	rpsM		adk	secY	rplO		rpmD	rpsE	rplR	rplF	rpsH	rpsN	rplE	rplX	rplN	rpsQ	rpmC					rplP
hypothetical protein	phosphoglycerate mutase	glutamate/glutamine/aspartate/aspartagine ABC transporter ATP-binding protein	transporter permease	dutamate/dutaming/accountate/accountaing ABC	transporter substrate-binding protein glutamate/glutamine/aspartate/asparagine ABC	hypothetical protein glutamate/glutamine/aspartate/asparagine ABC	HAD family hydrolase	(EC:4.2.1.70)	recombination factor protein RarA	serine protease	hypothetical protein	50S ribosomal protein L17	EC:2.7.7.6)	30S ribosomal protein S11	30S ribosomal protein S13	hypothetical protein	adenylate kinase (EC:2.7.4.3)	preprotein translocase subunit SecY	50S ribosomal protein L15	hypothetical protein	50S ribosomal protein L30	30S ribosomal protein S5	50S ribosomal protein L18	50S ribosomal protein L6	30S ribosomal protein S8	30S ribosomal protein S14	50S ribosomal protein L5	50S ribosomal protein L24	50S ribosomal protein L14	30S ribosomal protein S17	50S ribosomal protein L29	cyclic nucleotide-binding protein	hypothetical protein	calcium-binding domain-containing protein	ISSpo2, transposase	50S ribosomal protein L16
3.55	-1.12	-1.63	-1.42	-1.60	-2.19	-1.32	-1.21	1.15	-1.08	1.91	-3.43	-1.17	-1.22	-1.40	-1.22	-1.53	-1.21	1.74	1.08	Х	1.74	1.12	-1.05	0.93	1.04	1.38	1.05	1.05	-1.08	-1.04	1.26	-1.36	-0.95	1.44	-1.00	-2.08
0.00	0.04	0.07	0.27	0.49	0.00	0.63	0.76	0.24	0.79	0.00	0.00	0.81	0.28	0.00	0.27	0.05	0.11	0.00	0.80	×	0.02	0.62	0.06	0.01	0.69	0.52	0.95	0.63	0.56	0.15	0.02	0.01	0.06	0.30	0.85	0.03
-1.01	1.32	-2.28	-2.44	-1.95	-2.27	-1.05	-1.11	1.00	1.02	1.38	-1.41	1.56	-1.20	-1.14	-1.69	1.13	1.00	-1.17	1.06	-1.20	1.25	0.97	1.05	-1.20	-1.18	0.99	-1.27	1.25	-1.07	-1.26	-1.34	1.11	1.08	1.61	-1.04	2.24
0.94	0.00	0.02	0.00	0.00	0.00	0.76	0.46	0.97	0.72	0.00	0.05	0.12	0.11	0.14	0.03	0.38	0.78	0.22	0.91	0.50	0.11	0.52	0.96	0.08	0.00	0.82	0.10	0.25	0.32	0.08	0.20	0.42	0.60	0.30	0.94	0.00
1.27	0.10	-1.96	-1.93	-1.78	-2.23	-1.19	-1.16	1.08	-0.03	1.65	-2.42	0.20	-1.21	-1.27	-1.46	-0.20	-0.11	0.29	1.07	N/A	1.50	1.04	0.00	-0.14	-0.07	1.19	-0.11	1.15	-1.08	-1.15	-0.04	-0.13	0.07	1.53	-1.02	0.08
2.28	1.22	0.33	0.51	0.18	0.04	0.14	0.05	0.08	1.05	0.27	1.01	1.37	0.01	0.13	0.23	1.33	1.10	1.46	0.01	N/A	0.24	0.08	1.05	1.06	1.11	0.19	1.16	0.10	0.01	0.11	1.30	1.24	1.01	0.09	0.02	2.16
2.79	-1.25	0.89	-1.28	-1.25	-2.32	2.94	3.02	4.30	1.17	1.96	-2.31	11.70	5.35	5.74	2.39	1.62	1.04	3.67	7.38	1.27	5.33	7.63	9.72	7.63	10.30	11.00	9.38	10.60	11.30	7.67	8.43	-1.33	6.88	-4.41	-1.52	1.79
0.00	0.05	0.48	0.36	0.05	0.00	0.04	0.02	0.00	0.00	0.00	0.00	0.05	0.00	0.00	0.01	0.00	0.35	0.00	0.00	х	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.00	0.00	0.56	0.01
1.21	1.47	1.01	-1.22	-1.16	-1.30	-1.55	-1.33	-1.40	-1.13	-1.42	-1.12	-1.28	-1.39	-1.71	-1.15	-1.40	-1.30	-1.43	-1.84	-1.59	-2.52	-1.48	-1.21	-1.51	-1.78	-1.80	-1.64	-1.08	-1.44	-1.57	-1.73	1.05	-1.03	2.15	1.14	-2.03
0.22	0.00	0.89	0.02	0.10	0.00	0.12	0.04	0.02	0.17	0.01	0.52	0.25	0.00	0.00	0.01	0.00	0.03	0.00	0.00	0.24	0.00	0.00	0.02	0.00	0.00	0.00	0.00	0.04	0.00	0.00	0.00	0.65	0.51	0.06	0.36	0.00
2.00	0.11	0.95	-1.25	-1.21	-1.81	0.70	0.85	1.45	0.02	0.27	-1.72	5.21	1.98	2.02	0.62	0.11	-0.13	1.12	2.77	-0.16	1.41	3.08	4.26	3.06	4.26	4.60	3.87	4.76	4.93	3.05	3.35	-0.14	2.93	-1.13	-0.19	-0.12
0.79	1.36	0.06	0.03	0.05	0.51	2.25	2.18	2.85	1.15	1.69	0.60	6.49	3.37	3.73	1.77	1.51	1.17	2.55	4.61	1.43	3.93	4.56	5.47	4.57	6.04	6.40	5.51	5.84	6.37	4.62	5.08	1.19	3.96	3.28	1.33	1.91
-1.29	1.43	1.01	-1.14	1.13	1.25	-1.42	-1.57	-1.37	1.51	1.17	1.13	1.06	-1.64	-1.30	-1.68	-1.42	-1.41	-1.50	-1.18	-1.24	1.28	1.21	-1.22	-1.26	-1.38	-1.33	-1.34	1.28	-1.56	-2.24	-2.65	1.41	-1.59	-1.09	-1.24	1.97
0.00	0.00	0.93	0.06	0.68	0.22	0.08	0.00	0.03	0.01	0.36	0.40	0.90	0.00	0.01	0.00	0.08	0.02	0.00	0.04	0.06	0.06	0.14	0.05	0.00	0.00	0.00	0.01	0.03	0.01	0.00	0.00	0.00	0.00	0.81	0.26	0.00
-1.22	1.18	1.02	1.06	1.12	1.42	-2.37	-2.17	-2.27	-1.15	-1.13	1.17	-2.00	-1.75	-1.77	-1.95	-2.02	-1.77	-1.55	-2.07	1.24	-2.53	-1.63	-1.83	-2.24	-2.99	-2.99	-2.81	-1.64	-2.61	-4.81	-4.38	1.17	-2.51	1.44	-1.20	1.01
0.20	0.01	0.78	0.65	0.96	0.05	0.01	0.00	0.00	0.41	0.21	0.36	0.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	х	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.00	0.38	0.81	0.19
-1.26	1.31	1.02	-0.04	1.13	1.34	-1.90	-1.87	-1.82	0.18	0.02	1.15	-0.47	-1.70	-1.54	-1.82	-1.72	-1.59	-1.53	-1.63	0.00	-0.63	-0.21	-1.53	-1.75	-2.19	-2.16	-2.08	-0.18	-2.09	-3.53	-3.52	1.29	-2.05	0.18	-1.22	1.49
0.04	0.13	0.01	1.10	0.00	0.09	0.48	0.30	0.45	1.33	1.15	0.02	1.53	0.06	0.24	0.14	0.30	0.18	0.03	0.44	1.24	1.91	1.42	0.31	0.49	0.81	0.83	0.74	1.46	0.53	1.29	0.87	0.12	0.46	1.27	0.02	0.48

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															uvrB						rpmH								engB				argB	
hypothetical protein glutathione S-transferase glycine cleavage system protein T	oligopeptide ABC transporter permease oligopeptide ABC transporter substrate-binding protein	oligopeptide ABC transporter permease	mechanosensitive ion channel protein MscS	UDP-glucose/GDP-mannose dehydrogenase	hypothetical protein	glycoside hydrolase	group 1 family glycosyltransferase	phosphoglycerate mutase	hypothetical protein	excinuclease ABC subunit B	glycine cleavage system protein T	hypothetical protein	sensor histidine kinase (EC:2.7.3)	mercuric reductase	hypothetical protein	50S ribosomal protein L34	ribonuclease P protein component (EC:3.1.26.5)	hypothetical protein	K+-dependent Na+/Ca+ exchanger-like protein	C32 tRNA thiolase	diguanylate cyclase	тны полотик росси наполосос сопролеть YidC	MOSC domain-containing protein	ribosome biogenesis GTP-binding protein YsxC	hypothetical protein	hypothetical protein	oxidoreductase	acetylglutamate kinase (EC:2.7.2.8)	sterol desaturase-like protein					
-1.17 3.06 1.16	1.50 -1.63	1.24	-1.06	-1.71	1.24	1.16	1.07	1.65	1.10	1.86	1.39	-2.04	-1.32	-1.15	-1.69	3.68	2.71	-1.02	1.13	-2.77	1.02	Х	1.29	Х	-1.26	1.28	-1.12	-1.06	1.28	1.20	1.13	1.72	1.10	-1.19
0.70 <mark>0.00</mark> 0.18	0.70 0.08	0.01	0.65	0.00	0.02	0.06	0.41	0.11	0.94	0.01	0.69	0.09	0.69	0.50	0.15	0.00	0.00	0.96	0.28	0.04	0.64	x	0.44	×	0.06	0.01	0.30	0.92	0.22	0.96	0.71	0.00	0.03	0.03
-1.07 1.35 -1.25	-1.15 1.03	1.05	1.03	-1.02	1.23	1.35	1.53	1.26	-1.34	1.18	1.26	-1.05	1.10	1.02	-1.26	1.68	1.19	1.14	-1.06	1.10	1.45	1.73	-1.07	-1.36	1.37	1.63	1.32	1.27	1.50	2.27	3.45	1.55	1.77	1.16
0.59 <mark>0.02</mark> 0.16	0.40 0.89	0.64	0.96	0.77	0.09	0.20	0.00	0.15	0.14	0.51	0.01	0.45	0.68	0.80	0.15	0.00	0.15	0.36	0.66	0.29	0.08	0.00	0.76	0.00	0.07	0.01	0.15	0.14	0.01	0.00	0.00	0.00	0.00	0.28
-1.12 2.21 -0.05	0.18	1.15	-0.02	-1.37	1.24	1.26	1.30	1.46	-0.12	1.52	1.33	-1.55	-0.11	-0.06	-1.48	2.68	1.95	0.06	0.03	-0.84	1.24	N/A	0.11	N/A	0.06	1.46	0.10	0.11	1.39	1.74	2.29	1.64	1.44	-0.02
0.05 0.86 1.21	1.33 1.33	0.10	1.05	0.35	0.01	0.10	0.23	0.19	1.22	0.34	0.06	0.50	1.21	1.09	0.21	1.00	0.76	1.08	1.10	1.94	0.22	N/A	1.18	N/A	1.32	0.17	1.22	1.17	0.11	0.54	1.16	0.09	0.34	1.18
1.84 1.37 -1.69	1.59 -2.39	1.03	1.16	-1.46	1.12	1.17	1.42	-1.01	2.70	-1.73	1.99	-1.18	-1.81	1.38	1.10	-1.19	-0.89	1.65	2.00	-3.20	5.47	6.91	2.92	2.78	1.81	-2.79	4.19	4.08	1.85	1.45	2.11	1.80	2.05	1.70
0.00 0.01 0.01	0.63	0.53	0.82	0.01	0.08	0.06	0.01	0.21	0.00	0.00	0.46	0.10	0.63	0.12	0.40	0.45	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.02	0.00	0.00	0.00	0.11	0.00	0.00	0.00	0.00
-1.55 -1.12 -1.64	1.30 1.26	1.53	1.19	-1.01	-1.55	-1.36	-1.06	-1.27	-1.27	-1.08	-1.05	-1.04	1.30	1.26	-1.12	1.40	-1.01	-1.14	1.11	-1.11	-1.57	-1.79	1.13	-2.02	-1.10	1.71	-1.09	-1.07	-1.21	1.68	1.31	1.09	1.14	1.13
0.00 0.13 0.00	0.48 0.16	0.01	0.64	0.57	0.00	0.03	0.16	0.02	0.34	0.34	0.58	0.54	0.31	0.07	0.08	0.01	0.95	0.37	0.39	0.64	0.00	0.00	0.62	0.00	0.45	0.00	0.49	0.47	0.04	0.00	0.02	0.21	0.27	0.12
0.15 0.13 -1.67	1.45 -0.57	1.28	1.18	-1.24	-0.22	-0.10	0.18	-1.14	0.72	-1.41	0.47	-1.11	-0.26	1.32	-0.01	0.11	-0.95	0.26	1.56	-2.16	1.95	2.56	2.03	0.38	0.36	-0.54	1.55	1.51	0.32	1.57	1.71	1.45	1.60	1.42
1.70 1.25 0.03	0.15 1.83	0.25	0.02	0.23	1.34	1.27	1.24	0.13	1.99	0.33	1.52	0.07	1.56	0.06	1.11	1.30	0.06	1.40	0.44	1.05	3.52	4.35	0.90	2.40	1.46	2.25	2.64	2.58	1.53	0.12	0.40	0.36	0.46	0.28
-1.20 1.36 1.34	-1.10 1.21	1.18	1.23	-1.54	-1.05	-1.01	-1.34	-1.10	-1.09	1.04	-1.14	1.61	-1.18	-1.10	1.14	-1.10	-1.23	1.15	1.11	-1.27	-1.09	-1.55	1.36	1.54	-1.01	-1.19	-1.52	-1.69	-1.65	-1.15	-1.34	1.20	-1.41	-1.05
0.05 0.00 0.07	0.70 0.01	0.00	0.47	0.00	0.53	0.98	0.01	0.63	0.61	0.70	0.09	0.01	0.84	0.26	0.13	0.39	0.01	0.03	0.47	0.16	0.14	0.00	0.09	0.00	0.79	0.14	0.04	0.00	0.01	0.01	0.02	0.03	0.03	0.33
-1.43 1.40 1.22	-1.03 1.06	1.15	1.68	-1.64	-1.40	-1.48	-1.68	-1.38	2.25	1.45	-1.07	-1.01	1.47	-1.09	1.28	х	-0.98	-1.25	1.17	1.20	-1.59	-3.75	1.43	1.32	1.25	-0.97	-1.59	-2.50	-2.17	3.01	3.35	-1.29	-1.63	-1.43
0.20 <mark>0.00</mark> 0.06	0.87 0.34	0.07	0.26	0.00	0.04	0.06	0.00	0.08	0.05	0.19	0.61	0.84	x	0.95	0.03	×	0.68	0.31	0.25	0.17	0.00	0.00	0.26	0.00	0.55	0.28	0.04	0.00	0.00	0.00	0.00	0.01	0.04	0.01
-1.32 1.38 1.28	-1.07 1.14	1.17	1.46	-1.59	-1.23	-1.25	-1.51	-1.24	0.58	1.25	-1.11	0.30	0.15	-1.10	1.21	N/A	-1.11	-0.05	1.14	-0.04	-1.34	-2.65	1.40	1.43	0.12	-1.08	-1.56	-2.10	-1.91	0.93	1.01	-0.05	-1.52	-1.24
0.12 0.02 0.06	0.04 0.08	0.02	0.22	0.05	0.17	0.23	0.17	0.14	1.67	0.20	0.03	1.31	1.33	0.01	0.07	N/A	0.12	1.20	0.03	1.24	0.25	1.10	0.03	0.11	1.13	0.11	0.04	0.41	0.26	2.08	2.35	1.25	0.11	0.19

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	nspC	speB-I	slcD	hpsQ	hpsP	hpsO	hpsN	hpsM	hpsL	hpsK	hpsR						aspC-1		acdA-2							acdA-1										
saccharopine dehydrogenase (EC:1.5.1.7)	carboxynorspermidine decarboxylase	agmatinase (EC:3.5.3.11)	membrane-bound sulfolactate dehydrogenase	UspA family stress protein	s or n-uniyoroxypropanesurionate-2- dehydrogenase	K of S-uniyuroxypropanesurionate-2- dehydrogenase (EC:1.1.1.6) 6 or D dibudesenesesson	EC:1.1.23) Der S. d.L. 1.23	dihydroxypropanesulfonateTRAP transporter dihydroxypropanesulfonate-3-dehydrogenase	dihydroxypropanesulfonateTRAP transporter	dihydroxypropanesulfonateTRAP transporter	LacI family transcriptional regulator	oxitoreductase, INAL-binding/iron-sultur cluster- binding protein	LysR family transcriptional regulator	hypothetical protein	acyl carrier protein	dehydrogenase/transketolase	aspartate aminotransferase (EC:2.6.1.1)	LysR family transcriptional regulator	acyl-CoA dehydrogenase (EC:1.3.99)	hypothetical protein	2-nitropropane dioxygenase	acetyl-CoA carboxylase, biotin carboxylase	acetyl-CoA carboxylase carboxyltransferase	hypothetical protein	TetR family transcriptional regulator	acyl-CoA dehydrogenase (EC:1.3.99)	protein	A BC transporter transmembrane A TP hinding	guanylate cyclase	PKD domain-containing protein	hypothetical protein	AsnC family transcriptional regulator	2-oxoacid ferredoxin oxidoreductase	oxidoreductase, FAD-binding	TetR family transcriptional regulator	trimethylamine methyltransferase
1.38	-1.14	1.22	1.53	-2.57	х	х	Х	x	1.17	-5.75	х	-1.57	1.03	1.16	-1.38	1.15	1.53	-2.42	4.18	-1.05	-1.83	1.60	0.98	1.31	-1.55	1.10	-1.28	-3.04	-1.49	1.05	2.26	3.97	2.54	1.36	1.21	-1.35
0.66	0.31	0.02	0.01	0.01	×	х	x	х	0.73	0.00	×	0.00	0.20	0.28	0.01	0.62	0.02	0.03	0.05	0.21	0.34	0.05	0.00	0.01	0.01	0.51	0.00	0.01	0.02	0.96	0.02	0.00	0.00	0.00	0.00	0.25
1.06	1.11	1.32	-1.33	-1.30	-1.49	1.05	-1.28	Х	-1.26	-1.68	1.39	-1.32	-1.54	1.12	1.20	1.26	-1.07	-1.07	1.41	-1.15	-1.17	1.01	1.22	1.12	-1.39	-1.10	1.10	-1.31	1.04	1.08	1.24	1.04	-1.47	1.10	-1.16	-1.11
0.87	0.38	0.03	0.17	0.15	0.03	0.60	0.19	х	0.05	0.08	0.00	0.67	0.19	0.31	0.07	0.10	0.92	0.68	0.06	0.53	0.53	0.96	0.03	0.18	0.01	0.80	0.57	0.12	0.85	0.53	0.27	0.82	0.41	0.09	0.14	0.56
1.22	-0.01	1.27	0.10	-1.94	N/A	N/A	N/A	N/A	-0.05	-3.72	N/A	-1.45	-0.26	1.14	-0.09	1.21	0.23	-1.75	2.80	-1.10	-1.50	1.31	1.10	1.22	-1.47	0.00	-0.09	-2.18	-0.23	1.07	1.75	2.51	0.54	1.23	0.03	-1.23
0.16	1.13	0.05	1.43	0.63	N/A	N/A	N/A	N/A	1.22	2.04	N/A	0.13	1.29	0.02	1.29	0.06	1.30	0.68	1.39	0.05	0.33	0.30	0.12	0.10	0.08	1.10	1.19	0.87	1.27	0.02	0.51	1.47	2.01	0.13	1.19	0.12
1.46	0.98	1.30	2.68	-4.91	-1.75	-1.44	-2.23	-2.32	1.76	-6.29	-1.21	-1.10	-0.98	2.62	1.01	1.33	-1.97	-2.64	-2.07	-2.70	-3.09	0.99	-4.96	-2.34	-2.28	-1.27	-2.34	-5.98	-1.80	2.45	1.12	-1.79	-8.32	1.59	2.69	-1.46
0.58	0.88	0.00	0.00	0.00	0.45	0.58	0.00	0.00	0.01	0.00	0.69	0.49	0.89	0.00	1.00	0.24	0.17	0.00	0.22	0.00	0.00	0.97	0.00	0.00	0.00	0.05	0.00	0.00	0.00	0.12	0.22	0.03	0.00	0.00	0.00	0.46
1.50	1.17	-1.21	-1.19	-1.02	1.18	Х	-1.12	Х	-1.34	1.28	-1.57	1.22	-1.25	-1.12	-1.24	-1.25	-1.02	-1.06	1.88	-1.16	-1.39	-1.14	-1.05	-0.97	-1.49	-1.20	1.24	-1.16	-1.02	1.19	1.20	-1.81	-1.48	1.07	1.03	-1.27
0.02	0.14	0.04	0.29	0.99	0.13	х	x	x	0.11	0.12	0.01	0.52	0.53	0.49	0.03	0.09	0.95	0.85	0.02	0.59	0.33	0.75	0.96	0.71	0.00	0.42	0.26	0.46	1.00	0.41	0.28	0.01	0.66	0.37	0.54	0.05
1.48	1.08	0.05	0.75	-2.97	-0.29	N/A	-1.68	N/A	0.21	-2.51	-1.39	0.06	-1.12	0.75	-0.12	0.04	-1.50	-1.85	-0.10	-1.93	-2.24	-0.08	-3.01	-1.66	-1.89	-1.24	-0.55	-3.57	-1.41	1.82	1.16	-1.80	-4.90	1.33	1.86	-1.37
0.02	0.09	1.26	1.94	1.95	1.47	N/A	0.56	N/A	1.55	3.79	0.18	1.16	0.13	1.87	1.13	1.29	0.48	0.79	1.98	0.77	0.85	1.07	1.96	0.68	0.40	0.04	1.79	2.41	0.39	0.63	0.04	0.01	3.42	0.26	0.83	0.10
-1.33	-2.10	-1.52	-1.24	-1.20	-1.54	-1.02	-0.99	-1.11	-1.22	1.08	1.07	-1.36	-1.54	-1.03	-1.18	-1.15	-1.31	-1.12	1.40	1.21	1.12	1.30	1.34	1.12	-1.29	1.13	1.06	1.64	-1.10	-1.13	-0.98	-1.43	-1.45	1.40	-1.02	1.23
0.02	0.00	0.00	0.24	0.11	0.01	0.74	0.55	0.68	0.44	0.35	0.33	0.51	0.16	0.87	0.03	0.32	0.24	0.63	0.07	0.12	0.79	0.29	0.00	0.23	0.01	0.29	0.53	0.01	0.38	0.62	0.67	0.00	0.36	0.02	0.83	0.26
-1.12	-2.19	-1.96	1.76	1.24	х	x	Х	Х	1.40	-1.16	-1.01	1.44	-1.37	1.10	-1.01	-1.45	1.05	1.13	3.86	1.96	1.84	1.45	Х	-0.89	-1.25	-1.22	1.07	2.16	1.22	1.31	-0.91	-1.24	-1.11	-1.21	-1.09	1.29
0.30	0.00	0.00	0.11	0.06	×	х	х	Х	0.02	0.98	х	0.37	0.58	0.23	0.31	0.11	x	0.19	0.00	0.04	0.22	0.12	×	0.39	0.23	0.77	0.61	0.00	0.01	0.35	0.33	х	х	0.34	0.87	0.02
-1.23	-2.15	-1.74	0.26	0.02	N/A	N/A	N/A	N/A	0.09	-0.04	0.03	0.04	-1.46	0.04	-1.10	-1.30	-0.13	0.00	2.63	1.59	1.48	1.38	N/A	0.12	-1.27	-0.05	1.07	1.90	0.06	0.09	-0.95	-1.34	-1.28	0.10	-1.06	1.26
0.11	0.04	0.22	1.50	1.22	N/A	N/A	N/A	N/A	1.31	1.12	1.04	1.40	0.09	1.07	0.09	0.15	1.18	1.13	1.23	0.38	0.36	0.08	N/A	1.00	0.02	1.18	0.01	0.26	1.16	1.22	0.03	0.10	0.17	1.31	0.04	0.03

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SPO0645	SP00644	SPO0643	SP00642	SP00641	SP00640	SPO0639	SPO0638	SPO0637	SPO0636	SPO0635	SPO0634	SPO0633 ggt	SPO0632	SPO0631	SPO0629	SPO0628	SPO0624	SPO0623	SPO0622	SPO0621	SPO0620	SPO0619	SPO0618	SPO0616	SPO0615	SP00614	SPO0613	SPO0612	SPO0611	SPO0610	SPO0609	SPO0608	SPO0607	SPO0606	SPO0605	SPO0604	SPO0603	SPO0602 speA
multidrug resistance efflux pump	hypothetical protein	aldo/keto reductase	oxidoreductase, FAD-binding	hypothetical protein	alkylhydroperoxidase	LysR family transcriptional regulator	hypothetical protein	hypothetical protein	EF hand domain-containing protein	glycine cleavage system protein T	oxidoreductase, FAD-binding	gamma-glutamyltranspeptidase (EC:2.3.2.2)	2-hydroxyacid dehydrogenase	hypothetical protein	ISSpo3, transposase	transposase, degenerate	hypothetical protein	cytochrome c oxidase domain-containing protein	ISSpo1, transposase	ankyrin repeat-containing protein	hypothetical protein	sterol desaturase	AraC family transcriptional regulator	zinc-binding dehydrogenase oxidoreductase	hypothetical protein	transcriptional regulator	hypothetical protein	sugar ABC transporter permease	sugar ABC transporter permease	sugar ABC transporter ATP-binding protein	sugar ABC transporter ATP-binding protein	sugar ABC transporter substrate-binding protein	proline racemase	hypothetical protein	MarR family transcriptional regulator	hypothetical protein	hypothetical protein	arginine decarboxylase (EC:4.1.1.19)

1.20	0.97	-1.60	-1.00	1.58	Х	-2.54	2.61	2.08	х	-1.55	-1.24	-1.59	2.08	0.92	0.90	-2.41	-1.04	-1.22	-2.30	-1.60	-0.99	1.09	-2.48	-1.18	-1.31	-1.36	-1.11	-1.42	-1.85	-1.62	-1.83	-3.13	0.95	1.04	-2.15	-1.45	-1.18	-1.05
0.80	0.08	0.00	0.98	0.00	x	x	0.00	0.00	×	0.03	0.00	0.00	0.00	0.00	0.01	0.03	0.86	0.29	0.40	0.06	0.89	×	0.00	0.03	0.01	0.03	0.00	0.01	0.07	0.01	0.06	0.01	0.89	0.99	0.00	0.01	0.20	0.81
1.47	1.07	-1.03	1.30	1.12	1.56	-0.99	-1.01	1.50	1.15	1.70	-1.25	-1.29	2.40	-1.09	-1.38	-1.56	-1.27	1.05	-1.16	1.05	-1.07	1.08	-1.34	1.10	1.04	0.99	-1.11	1.10	-1.19	-1.27	-1.00	-1.77	1.16	-1.14	-1.96	-1.86	-1.59	1.06
0.02	0.23	0.52	0.24	0.03	0.07	0.67	0.88	0.00	0.71	0.02	0.15	0.05	0.00	0.07	0.03	0.09	0.03	0.57	0.79	0.83	0.79	0.76	0.12	0.04	0.67	0.67	0.75	0.67	0.79	0.08	0.78	0.01	0.14	0.67	0.00	0.00	0.04	0.85
1.34	1.02	-1.32	0.15	1.35	N/A	-1.76	0.80	1.79	N/A	0.08	-1.25	-1.44	2.24	-0.08	-0.24	-1.99	-1.16	-0.09	-1.73	-0.28	-1.03	1.09	-1.91	-0.04	-0.14	-0.19	-1.11	-0.16	-1.52	-1.45	-1.41	-2.45	1.06	-0.05	-2.06	-1.66	-1.39	0.01
0.14	0.05	0.29	1.15	0.23	N/A	0.78	1.81	0.29	N/A	1.63	0.01	0.15	0.16	1.01	1.14	0.43	0.12	1.14	0.57	1.33	0.04	0.01	0.57	1.14	1.18	1.17	0.00	1.26	0.33	0.18	0.42	0.68	0.10	1.09	0.10	0.21	0.21	1.06
-1.32	-1.80	1.41	1.47	3.21	3.29	-1.38	-3.79	-2.92	-1.14	-2.14	-1.81	-5.72	1.58	0.75	-2.49	-3.17	1.55	1.30	-3.74	-2.22	-1.88	1.26	-3.81	-1.19	-2.82	-3.33	-2.96	-4.07	-6.24	-2.62	-7.07	-39.50	-3.10	-1.60	-1.58	-1.56	-1.09	1.47
0.02	0.00	0.00	0.38	0.00	0.00	0.15	0.00	0.00	0.18	0.02	0.00	0.00	0.01	0.00	0.00	0.00	0.03	0.11	0.28	0.03	0.13	0.38	0.00	0.33	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.35	0.58	0.04	0.01	0.44	0.13
1.19	1.50	-1.01	1.21	1.24	-1.09	-1.35	-1.08	-1.14	х	-1.34	1.19	1.94	1.53	-1.18	1.19	1.12	-1.04	-1.02	-0.99	1.18	-0.99	-1.09	-1.02	1.22	1.13	1.25	-1.04	-1.01	-0.99	-1.22	-1.07	1.22	1.36	1.51	-1.46	-1.27	- 1.45	1.23
0.23	0.00	0.75	0.08	0.00	0.68	0.06	0.08	0.04	x	0.08	0.14	0.00	0.01	0.01	0.03	0.52	0.51	0.93	0.99	0.43	0.92	0.79	0.92	0.19	0.31	0.14	0.95	0.97	0.97	0.50	0.80	0.01	0.03	0.18	0.01	0.01	0.07	0.34
-0.07	-0.15	0.20	1.34	2.23	1.10	-1.37	-2.44	-2.03	N/A	-1.74	-0.31	-1.89	1.56	-0.22	-0.65	-1.03	0.26	0.14	-2.37	-0.52	-1.44	0.09	-2.42	0.02	-0.85	-1.04	-2.00	-2.54	-3.61	-1.92	-4.07	-19.14	-0.87	-0.05	-1.52	-1.42	-1.27	1.35
1.26	1.65	1.21	0.13	0.98	2.19	0.01	1.36	0.89	N/A	0.40	1.50	3.83	0.03	0.97	1.84	2.15	1.30	1.16	1.37	1.70	0.45	1.18	1.40	1.21	1.98	2.29	0.96	1.53	2.63	0.70	3.00	20.36	2.23	1.56	0.06	0.14	0.18	0.12
1.76	1.34	1.02	1.68	-1.02	1.18	-1.26	-1.63	-0.98	-1.35	-1.24	1.07	-1.13	1.39	1.08	-1.31	-1.17	-1.59	-1.34	-1.15	1.15	-1.08	-1.18	-1.36	-1.03	1.25	1.16	1.15	1.06	-1.18	-1.35	-0.96	1.43	1.23	-1.08	-1.88	-1.49	-1.43	-2.13
0.00	0.00	0.91	0.00	0.75	0.15	0.00	0.00	0.77	0.18	0.01	0.43	0.34	0.00	0.65	0.01	0.56	0.01	0.25	0.78	0.49	0.78	0.55	0.28	0.95	0.01	0.17	0.30	0.54	0.78	0.31	0.56	0.00	0.10	0.76	0.00	0.01	0.00	0.00
1.34	1.56	-1.43	1.32	-1.12	-1.14	-1.19	-2.01	-1.36	x	-1.46	1.55	1.98	1.11	-1.14	-1.27	1.73	-1.81	-1.97	1.31	-0.97	1.31	-0.95	1.77	-1.33	1.15	1.35	-0.89	-1.01	1.53	1.57	-1.77	-1.02	1.16	1.12	-1.86	-1.48	-1.63	-2.02
0.01	0.00	0.00	0.03	0.41	0.98	x	0.00	0.02	×	0.01	0.05	0.01	0.36	0.00	0.03	0.04	0.00	0.04	0.68	0.70	0.22	0.32	0.01	0.49	0.06	0.11	0.27	0.65	0.50	0.07	×	0.36	0.12	0.38	0.00	0.00	0.01	0.02
1.55	1.45	-0.21	1.50	-1.07	0.02	-1.23	-1.82	-1.17	N/A	-1.35	1.31	0.43	1.25	-0.03	-1.29	0.28	-1.70	-1.66	0.08	0.09	0.12	-1.07	0.21	-1.18	1.20	1.26	0.13	0.03	0.18	0.11	-1.37	0.21	1.20	0.02	-1.87	-1.49	-1.53	-2.08
0.21	0.11	1.23	0.18	0.05	1.16	0.04	0.19	0.19	N/A	0.11	0.24	1.56	0.14	1.11	0.02	1.45	0.11	0.32	1.23	1.06	1.20	0.11	1.57	0.15	0.05	0.10	1.02	1.04	1.36	1.46	0.41	1.23	0.04	1.10	0.01	0.01	0.10	0.05

## Chapter 10: Appendix

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SPO0683	SPO0682	SPO0681	SPO0680	SPO0679	SPO0678	SPO0677	SPO0676	SPO0675	SPO0674	SPO0673	SPO0672	SPO0671	SPO0670	SPO0669	SPO0668	SPO0667	SPO0666	SPO0665	SPO0664	SPO0663	SPO0662	SPO0661	SPO0660	SPO0659	SPO0658	SPO0657	SPO0656	SPO0655	SPO0654	SPO0653	SPO0652	SPO0651	SPO0650	SPO0649	SPO0648	SPO0647	SPO0646
				hmgC			tauC	tauB	tauA	tpa				hisG	hisZ	hisS			naaC'	naaC	naaB'	naaB	naaA	naaR	naaS	naaT	dnaE		xdhA	xdhB	xdhC						
metallo-beta-lactamase	FAD-dependent oxidoreductase	hypothetical protein	glyoxalase	maleylacetoacetate isomerase (EC:5.2.1.2)	agmatinase	acyl-CoA synthetase (EC:2.3.1.86)	taurine ABC transporter permease	taurine ABC transporter ATP-binding protein	taurine ABC transporter substrate-binding protein	taurinepyruvate aminotransferase (EC:2.6.1.77)	hypothetical protein	hypothetical protein	DNA polymerase III subunit alpha	ATP phosphoribosyltransferase (EC:2.4.2.17)	ATP phosphoribosyltransferase	histidyl-tRNA synthetase (EC:6.1.1.21)	enoyl-CoA hydratase (EC:4.2.1.17)	SlyX protein	n-acetynaunne ABC transporter ATF-omung protein	protein	N-acetyltaurine ABC transporter permease N-acetyltaurine ABC transporter ATP-binding	N-acetyltaurine	protein	LysR family transcriptional regulator	N-acetyltaurine amidohydrolas	metallochaperone	DNA polymerase III subunit alpha (EC:2.7.7.7)	hypothetical protein	xanthine dehydrogenase, A subunit (EC:1.17.1.4)	xanthine dehydrogenase subunit B (EC:1.17.1.4)	xanthine dehydrogenase accessory factor	sugar ABC transporter ATP-binding protein	sugar ABC transporter permease	sugar ABC transporter permease	bmp family protein	hypothetical protein	hypothetical protein
-1.46	-1.16	1.03	-1.21	-2.24	1.00	-1.06	-2.14	-3.37	-4.95	2.01	-1.93	Х	-1.98	1.40	-2.75	0.87	1.82	-1.04	-0.97	-1.23	-1.07	-1.44	-1.56	-2.47	1.13	-2.66	1.31	1.14	-1.87	-1.54	-2.21	-1.59	-1.41	-2.16	1.12	-1.72	-2.33
0.01	0.01	0.81	0.02	0.00	0.61	0.72	0.00	0.08	0.00	0.00	х	x	0.07	0.05	0.00	0.00	0.01	0.57	0.91	0.81	0.95	0.02	0.29	0.01	0.06	0.00	0.02	0.62	0.00	0.01	0.00	0.00	0.00	0.00	0.86	0.06	0.00
1.07	-1.28	-1.32	1.02	-1.16	-1.17	-1.09	1.11	-1.42	-1.38	1.21	-1.07	-1.16	-1.39	2.21	1.89	1.08	1.14	-1.12	-1.14	-1.04	1.04	-1.19	-1.23	-1.16	-1.08	-1.29	-1.16	1.65	1.11	1.11	1.04	1.24	1.32	-1.66	0.98	-1.15	1.21
0.78	0.30	0.07	0.81	0.03	0.36	0.83	0.08	0.45	0.00	0.05	0.86	0.61	0.51	0.00	0.00	0.82	0.48	0.07	0.86	0.90	0.84	0.38	0.56	0.70	0.82	0.03	0.12	0.19	0.01	0.39	0.47	0.12	0.03	0.01	0.57	0.19	0.20
-0.20	-1.22	-0.15	-0.10	-1.70	-0.09	-1.08	-0.52	-2.40	-3.17	1.61	-1.50	N/A	-1.69	1.81	-0.43	0.98	1.48	-1.08	-1.05	-1.14	-0.02	-1.32	-1.40	-1.82	0.02	-1.98	0.08	1.40	-0.38	-0.22	-0.59	-0.18	-0.04	-1.91	1.05	-1.44	-0.56
1.27	0.06	1.18	1.12	0.54	1.09	0.02	1.63	0.98	1.79	0.40	0.43	N/A	0.29	0.41	2.32	0.10	0.34	0.04	0.09	0.10	1.06	0.13	0.17	0.66	1.11	0.69	1.24	0.26	1.49	1.33	1.63	1.42	1.37	0.25	0.07	0.28	1.77
-1.58	1.11	1.41	1.24	-1.91	-1.72	-1.55	-1.09	-4.47	-15.30	1.26	-2.03	1.41	-2.76	2.32	-3.34	-2.60	-1.57	1.88	-1.09	-1.52	-1.32	-2.42	-3.69	-3.55	-2.01	-3.95	2.94	-1.18	-5.18	-3.53	-5.21	-3.30	-2.78	-4.21	0.87	1.05	1.29
3 0.02	0.38	0.57	0.02	0.00	0.04	0.15	0.82	0.05	0 0.00	0.24	0.01	0.00	0.03	0.00	0.00	0.00	0.32	0.00	0.69	0.72	0.96	0.07	0.00	0.00	0.01	0.00	0.00	-	s 0.00	0.00	0.00	0.00	s 0.00	0.00	0.00	0.74	0.00
1.26	-1.11	1.09	1.03	1.12	. 1.30	1.04	-1.22	-1.15	-1.08	. 0.97	-1.1	-			-1.76	-1.15	2.08	1.05	-1.29	1.06	X	-1.30	1.23		-1.09	_	1.06		1.34	-0.97	1.23	1.03		1.34	1.31		-1.00
6 0.28	0.41	9 0.59	3 0.77	2 0.23	0 0.31	4 0.91	0.11	5 0.74	0.28	7 0.18	0 X	6 0.31	0.97	5 0.00	6 <u>0.00</u>	5 0.26	8 0.00	5 0.75	9 0.47	6 0.67	X	0 0.38	3 0.40	7 0.68	9 0.56	5 0.71	6 0.59	1 0.79	4 0.06	0.79	3 0.02	3 0.98	0.76	4 0.05	1 0.02	-	0 0.89
-0.	-1 0.00	9 1.25	7 1.14	-0.40	-0.21	-0.26	1 -1.16	-2.81	.8 -8.19	8 1.12	-1.57	0.13	-1.90	0 0.39	0 -2.55	-1.88	0 0.26	5 1.47	.7 -1.19	-0.23	N/A	8 -1.86	-1.23					-1.15		·9 -2.25	2 -1.99		6 -1.91				.9 0.15
16 1.																																					
1.42 - 1	1.11	0.16 -1	0.11 -1	1.52 -1	1.51 -1	1.30 1	0.06 1	1.66 - ]	7.11 1	0.14 1	0.47 1	1.29 1	0.86 - ]	1.94 1	0.79 - 1	0.73 1	1.83 2	0.42 - ]	0.10 -(	1.29 -1	N/A -]	0.56 - ]	2.46 1	1.19 1	0.46 -(	1.40 -1	0.94 1		3.26 - 1	1.28 1	3.22 1	2.17 1	0.88 - 1	2.78 -1	0.22 0		1.15 1
1.12 0	1.27 0	-1.40 0	-1.11 0	-1.42 0	-1.15 0	1.20 0	1.33 0	-1.12 0	1.36 0	1.17 0	1.12 0	1.17 0	-1.08 0	1.23 0	-1.14 0	1.08 0	2.19 0	-1.24 0		-1.03 0	-1.01 0		1.26 0	1.09 0	0.99 0	-1.03 0	1.07 0	1.37 0	1.14 0	1.21 0	1.04 0	1.27 0	1.16 0	-1.58 0			1.49 0
0.28	0.23	0.05	0.47 -	0.00	0.41	0.44	0.03	0.82	0.02	0.03	0.49	0.01	0.84	0.04	0.15 -	0.69	0.01	0.14	0.70	0.99	0.69	0.92	0.34	0.42	0.75	0.94	0.53	0.09 -	0.13 -	0.18	0.49	0.03	0.01	0.00		0.54	0.03
1.18	1.95	-1.41	-1.35	-1.15	1.04	1.08	-1.17	1.16	1.35	1.56	1.09	х	1.99	-1.84	-1.50	1.78	4.35	-1.70	1.72	1.62	Х	-1.22	1.23	1.20	1.31	1.66			-1.15	1.25	-1.20	1.64	-1.40	-1.15	1.52		-1.24
0.40	0.03	0.10	0.01	0.71	0.87	0.72	0.88	0.67	0.00	0.01	x	x	0.29	0.00	0.07	0.01	0.00	0.00	0.40	0.04	×	X	0.19	0.21	0.03	0.08	0.00	0.86	0.93	0.01	0.33	0.05	0.00	0.00	0.00		0.30
-1.15	1.61	-1.41	-1.23	-1.29	-0.05	1.14	0.08	0.02	1.36	1.37	1.11	N/A	0.46	-0.31	-1.32	1.43	3.27	-1.47	0.38	0.30	N/A	-1.13	1.25	1.15	0.16	0.32	-0.10	0.22	-1.15	1.23	-0.08	1.46	-1.28	-1.37	1.24	-0.16	0.13
0.03	0.34	0.01	0.12	0.14	1.10	0.06	1.25	1.14	0.01	0.20	0.02	N/A	1.54	1.54	0.18	0.35	1.08	0.23	1.34	1.33	N/A	0.09	0.02	0.05	1.15	1.35	1.17	1.15	0.01	0.02	1.12	0.18	0.12	0.22	0.28	1.23	1.37

Ν	S
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SPO0721	SPO0720	SPO0719	SPO0718	SPO0717	SPO0716	SPO0715	SPO0714	SPO0713	SPO0712	SPO0711	SPO0710	SPO0709	SPO0708	SPO0707	SPO0706	SPO0705	SPO0704	SPO0703	SPO0702	SPO0701	SPO0700	SPO0699	SPO0698	SPO0697	SPO0695	SPO0694	SPO0693	SPO0692	SPO0691	SPO0690	SPO0689	SPO0688	SPO0687	SPO0686	SPO0685	SPO0684
	etfB	etfA		bhbD						chvG	chvI	pckA								gap-1		modA	modB	modC					era					hmgA	hmgB	
glyoxalase	electron transfer flavoprotein subunit beta	electron transfer flavoprotein subunit alpha	hypothetical protein	3-hydrox yburyry1-CoA denydrogenase (EC:1.1.1.157)	hypothetical protein	phosphocarrier protein HPr	PTS system mannose subfamily IIA subunit	hypothetical protein	protein	sensor histidine kinase ChvG (EC:2.7.3) Hor serine kinase/bhosphatase domain-containing	DNA-binding response regulator ChvI	phosphoenolpyruvate carboxykinase (EC:4.1.1.49)	invasion protein IbeA	DeoR family transcriptional regulator	protein	oligopeptide ABC transporter permease	oligopeptide ABC transporter permease	oligopeptide ABC transporter ATP-binding protein	oligopeptide ABC transporter ATP-binding protein	EC:1.2.1)	molybdenum-binding transcriptional regulator	protein	molybdate ABC transporter permease	(EC:3.6.3.29)	hypothetical protein	hypothetical protein	isobutyryl-CoA dehydrogenase	multicopper oxidase	GTP-binding protein Era	hypothetical protein	hypothetical protein	guanylate cyclase	MarR family transcriptional regulator	homogentisate 1,2-dioxygenase (EC:1.13.11.5)	fumarylacetoacetase (EC:3.7.1.2)	glyoxalase
-1.81	1.24	2.45	-1.15	2.32	1.39	1.06	-1.20	1.53	-1.19	1.04	-1.64	1.10	10.90	-1.97	-1.40	-1.09	1.21	-1.68	1.11	1.82	1.16	-1.49	х	1.65	-1.58	1.14	-2.10	1.17	1.31	x	-1.07	1.03	-3.22	-1.98	-1.33	-1.38
0.00	0.61	0.00	0.60	0.00	0.24	0.11	0.00	0.00	0.01	0.76	0.07	0.99	0.00	0.00	0.00	0.66	0.56	0.12	0.04	0.00	0.63	0.13	x	0.03	0.00	0.21	0.00	0.17	0.48	x	x	0.99	0.00	0.00	0.02	0.27
1.19	1.76	2.23	-1.19	1.94	1.33	1.11	-1.36	1.03	-1.31	-1.16	-1.38	1.03	1.46	-1.57	-1.43	-1.23	1.10	-1.10	-1.08	1.96	-1.08	-1.14	-1.08	-1.12	1.29	-1.31	1.34	1.51	1.19	1.97	1.01	-1.09	-1.17	-1.22	-1.02	1.02
0.01	0.00	0.00	0.52	0.00	0.43	0.48	0.00	0.88	0.65	0.43	0.11	1.00	0.07	0.00	0.30	0.16	0.47	0.68	0.66	0.00	0.35	0.67	0.74	0.60	0.00	0.01	0.03	0.01	0.34	0.00	0.84	0.18	0.01	0.20	0.77	0.75
-0.31	1.50	2.34	-1.17	2.13	1.36	1.09	-1.28	1.28	-1.25	-0.06	-1.51	1.07	6.18	-1.77	-1.42	-1.16	1.16	-1.39	0.02	1.89	0.04	-1.32	N/A	0.27	-0.15	-0.09	-0.38	1.34	1.25	N/A	-0.03	-0.03	-2.20	-1.60	-1.18	-0.18
1.50	0.26	0.11	0.02	0.19	0.03	0.03	0.08	0.25	0.06	1.10	0.13	0.04	4.72	0.20	0.02	0.07	0.05	0.29	1.10	0.07	1.12	0.18	N/A	1.39	1.44	1.23	1.72	0.17	0.06	N/A	1.04	1.06	1.03	0.38	0.16	1.20
-7.43	1.84	4.03	-2.83	2.55	2.55	0.94	1.04	2.04	-1.16	1.03	-2.32	1.41	1.55	-6.57	-4.58	-1.88	-1.01	-8.57	-1.55	-1.88	1.07	1.13	-1.28	1.51	1.14	2.37	-1.08	-1.10	3.58	4.08	1.70	1.52	-3.84	-2.57	-1.52	-1.60
0.00	0.00	0.00	0.00	0.00	0.03	0.02	0.29	0.00	0.86	0.80	0.01	0.59	0.00	0.00	0.00	0.00	0.81	0.00	0.01	0.00	0.37	0.84	0.54	0.04	0.02	0.00	0.97	0.59	0.01	0.00	0.01	0.01	0.00	0.00	0.00	0.00
1.29	1.07	0.99	1.01	1.09	-1.23	1.21	1.07	-1.03	1.08	1.08	1.18	-2.59	-1.23	-1.12	0.98	-1.45	-1.42	-1.22	-1.21	13.40	1.99	1.09	-1.13	1.13	1.16	1.06	-1.01	-1.14	-1.17	-1.37	-1.22	-1.18	-1.26	1.17	-1.01	1.00
0.03	0.90	0.38	1.00	0.47	0.47	0.18	0.84	0.68	0.89	0.88	0.22	0.07	0.14	0.31	0.64	0.00	0.09	0.31	0.11	0.00	0.00	0.79	0.45	0.37	0.03	0.52	0.91	0.61	0.00	0.01	0.02	0.04	0.08	0.15	0.57	0.72
-3.07	1.46	2.51	-0.91	1.82	0.66	1.08	1.06	0.51	-0.04	1.06	-0.57	-0.59	0.16	-3.85	-1.80	-1.67	-1.22	-4.90	-1.38	5.76	1.53	1.11	-1.21	1.32	1.15	1.72	-1.05	-1.12	1.21	1.36	0.24	0.17	-2.55	-0.70	-1.27	-0.30
4.36	0.38	1.52	1.92	0.73	1.89	0.13	0.02	1.54	1.12	0.03	1.75	2.00	1.39	2.73	2.78	0.21	0.21	3.68	0.17	7.64	0.46	0.02	0.08	0.19	0.01	0.66	0.04	0.02	2.38	2.73	1.46	1.35	1.29	1.87	0.25	1.30
1.36	1.33	1.18	-1.04	-1.02	-1.02	1.08	-1.33	-1.10	-1.03	1.55	-1.19	-1.04	-1.07	-1.33	-1.07	-1.27	1.17	-1.04	-1.49	-2.12	1.07	-1.44	-1.06	-1.34	-1.14	1.04	1.16	1.12	-1.20	-1.04	-1.17	1.19	-1.56	-1.64	-1.16	-1.32
0.02	0.03	0.04	0.91	0.67	0.87	0.63	0.02	0.28	0.96	0.07	0.04	0.84	0.80	0.02	0.69	0.07	0.23	0.83	0.00	0.00	0.39	0.26	0.89	0.12	0.23	0.44	0.13	0.07	0.09	0.71	0.00	0.02	0.00	0.00	0.03	0.00
-1.09	1.22	-1.43	2.23	-1.32	-1.47	-1.05	-1.19	-1.13	-1.19	1.09	-1.16	-1.15	-1.35	0.98	1.32	-1.42	-1.26	-1.17	-1.64	-3.88	-0.97	1.09	-0.91	-1.22	-1.48	1.45	-1.25	1.60	-1.34	-3.24	-1.84	1.02	-1.60	-1.53	-1.17	-1.74
0.72	0.14	0.01	0.01	0.01	0.15	0.22	0.15	0.53	0.78	0.85	0.02	0.64	0.63	0.43	0.44	0.11	0.52	0.65	0.00	0.00	0.49	0.69	0.13	0.70	0.00	0.01	0.44	×	0.06	0.15	0.00	0.99	0.01	0.01	0.07	0.01
0.14	1.28	-0.13	0.60	-1.17	-1.25	0.02	-1.26	-1.12	-1.11	1.32	-1.18	-1.10	-1.21	-0.18	0.13	-1.35	-0.05	-1.11	-1.57	-3.00	0.05	-0.18	-0.98	-1.28	-1.31	1.25	-0.05	1.36	-1.27	-2.14	-1.51	1.11	-1.58	-1.59	-1.17	-1.53
1.23	0.06	1.31	1.64	0.15	0.22	1.07	0.07	0.01	0.08	0.23	0.02	0.05	0.14	1.15	1.20	0.08	1.22	0.06	0.08	0.88	1.02	1.27	0.08	0.06	0.17	0.20	1.21	0.24	0.07	1.10	0.34	0.09	0.02	0.05	0.01	0.21

SPO0733 SPO0732 SPO0731

glpD

glpR

SPO0734

paaZ

SPO0728 SPO0727 SPO0726 SPO0725 SPO0724 SPO0723 SPO0722

tuf-1

parC

SPO0729

## Chapter 10: Appendix

alkylhydroperoxidase	hypothetical protein	beta-ketoadipyl CoA thiolase (EC:2.3.1)	phenylacetate-CoA oxygenase subunit PaaA	phenylacetate-CoA oxygenase subunit PaaB	phenylacetic acid degradation protein PaaI	phenylacetic acid degradation protein PaaJ	phenylacetic acid degradation oxidoreductase Paa	phenylacetic acid degradation protein	guanylate cyclase	hypothetical protein	hypothetical protein	hypothetical protein	peroxidase	hypothetical protein	hypothetical protein	hypothetical protein	TetR family transcriptional regulator	phenylacetate-CoA ligase (EC:6.2.1.30)	phenylacetic acid degradation protein PaaD	enoyl-CoA hydratase (EC:4.2.1.17)	enoyl-CoA hydratase	TRAP dicarboxylate transporter subunit DctP	TRAP dicarboxylate transporter subunit DctQ	TRAP dicarboxylate transporter subunit DctM	bitunicitoriai aluenyue denyulogenase/enoyi-COA hydratase	PaaX domain-containing protein	esterase	glycerol-3-phosphate dehydrogenase (EC:1.1.5.3)	glycerol-3-phosphate regulon repressor	hypothetical protein	elongation factor Tu (EC:3.6.5.3)	hypothetical protein	DNA topoisomerase IV subunit A (EC:5.99.1)	hypothetical protein	sitori chani denyai ogenase/reductase oxidoreductase	hypothetical protein	ATP:cob(I)alamin adenosyltransferase
-1.71	9.36	-1.24	0.99	1.15	1.17	1.38	1.30	1.19	-1.12	-1.91	-4.98	-2.07	Х	-1.09	-1.08	-1.22	-1.11	0.95	-1.18	-2.27	-2.29	-2.02	Х	-1.08	-1.44	-2.14	-1.20	-1.53	-1.18	-1.48	1.92	1.33	-1.46	Х	1.41	1.91	1.51
0.14	0.00	0.51	0.58	0.63	0.96	0.00	0.72	0.21	0.84	0.48	0.00	0.17	х	0.39	0.08	0.08	0.15	0.02	0.02	0.12	0.01	0.00	х	0.91	0.00	0.00	0.00	0.00	0.09	0.02	0.00	0.04	0.05	х	0.44	0.24	0.42
-1.42	11.80	1.89	1.12	1.78	1.53	1.22	1.22	1.15	-1.01	-1.12	-1.55	-1.17	-1.28	-1.26	-1.16	1.25	-1.18	-1.09	-1.03	-1.09	-1.31	-1.61	-1.13	-1.19	-1.06	-1.25	-1.03	-1.09	-1.03	1.09	-1.12	1.00	1.04	-1.21	1.51	1.33	1.72
0.56	0.00	0.00	0.40	0.01	0.19	0.02	0.69	0.47	0.98	0.87	0.02	0.66	0.09	0.01	0.36	0.32	0.41	0.35	0.66	0.80	0.04	0.00	0.27	0.75	0.77	0.21	0.42	0.68	0.78	0.59	0.12	0.87	0.83	0.25	0.06	0.08	0.18
-1.57	10.58	0.33	1.06	1.47	1.35	1.30	1.26	1.17	-1.07	-1.52	-3.27	-1.62	N/A	-1.18	-1.12	0.02	-1.15	-0.07	-1.11	-1.68	-1.80	-1.82	N/A	-1.14	-1.25	-1.70	-1.12	-1.31	-1.11	-0.20	0.40	1.17	-0.21	N/A	1.46	1.62	1.62
0.15	1.22	1.57	0.06	0.32	0.18	0.08	0.04	0.02	0.06	0.39	1.72	0.45	N/A	0.09	0.04	1.24	0.03	1.02	0.08	0.59	0.49	0.21	N/A	0.05	0.19	0.45	0.09	0.22	0.08	1.29	1.52	0.17	1.25	N/A	0.05	0.29	0.11
-3.40	32.00	1.26	-5.10	-3.18	-2.01	-3.53	-3.32	-2.00	-2.02	-1.77	-6.10	-1.98	-2.79	-2.40	-2.50	-1.75	-1.13	-2.98	-3.04	-3.89	-3.24	-2.20	-1.04	-1.13	-2.51	-3.14	2.35	-1.76	-0.96	1.94	7.16	2.35	-1.04	-2.04	1.46	3.44	2.03
0.01	0.00	0.19	0.00	0.00	0.23	0.00	0.07	0.00	0.47	0.58	0.00	0.34	0.01	0.00	0.00	0.07	0.78	0.00	0.00	0.00	0.00	0.00	0.01	0.95	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.81	0.01	0.27	0.02	0.08
-1.34	15.10	1.41	2.48	2.20	2.06	1.81	1.91	1.39	-1.00	1.19	×	-1.10	-1.14	1.52	1.31	1.57	-1.07	1.39	1.44	1.05	-1.49	-1.14	х	-1.11	1.34	1.23	1.14	-1.45	-1.54	-1.49	-1.65	-1.36	-1.06	1.33	-1.16	-1.14	-1.05
0.54	0.00	0.01	0.00	0.00	0.06	0.00	0.15	0.06	0.99	0.76	×	0.88	0.55	0.00	0.21	0.19	0.49	0.00	0.01	0.79	0.01	0.26	x	0.85	0.04	0.16	0.05	0.01	0.01	0.01	0.00	0.07	0.15	0.31	0.13	0.04	0.76
-2.37	23.55	1.34	-1.31	-0.49	0.03	-0.86	-0.71	-0.31	-1.51	-0.29	N/A	-1.54	-1.97	-0.44	-0.60	-0.09	-1.10	-0.80	-0.80	-1.42	-2.37	-1.67	N/A	-1.12	-0.59	-0.96	1.75	-1.61	-1.25	0.23	2.76	0.50	-1.05	-0.36	0.15		0.49
1.03	8.45	0.08	3.79	2.69	2.04	2.67	2.62	1.70	0.51	1.48	N/A	0.44	0.83	1.96	1.91	1.66	0.03	2.19	2.24	2.47	0.88	0.53	N/A	0.01	1.93	2.19	0.61	0.16	0.29	1.72	4.41	1.86	0.01	1.69	1.31	2.29	1.54
1.12	-2.70	-1.31	-1.39	-1.05	-1.31	-1.44	-1.11	-1.22	1.20	1.08	-1.45	-1.20	-1.07	-1.61	-1.68	1.06	-1.55	-1.00	-1.13	-1.12	-1.16	-1.45	-1.09	-1.21	-1.19	-1.30	-1.22	-1.09	-1.22	1.40	1.03	1.03	1.23	-1.45	-1.07	-1.26	1.07
0.79	0.00	0.06	0.07	0.52	0.32	0.00	0.72	0.12	0.65	0.89	0.27	0.36	0.87	0.00	0.02	0.77	0.00	0.88	0.62	0.77	0.22	0.02	0.72	0.71	0.27	0.02	0.00	0.19	0.04	0.00	0.79	0.99	0.14	0.00	0.65	0.00	0.87
-1.05	-4.97	-1.35	1.11	-1.14	-1.15	-1.23	1.22	1.12	-1.02	1.48	1.08	1.30	1.87	-1.29	1.19	-1.66	-2.14	-1.11	1.60	1.28	1.47	-1.13	-1.12	1.25	1.30	1.24	-1.19	-1.41	-1.47	-1.00	-1.55	-1.36	-1.06	Х	-1.48	-1.76	-1.62
0.98	0.00	0.24	0.57	0.63	0.80	0.17	0.55	0.49	0.95	0.51	×	×	0.06	0.06	0.42	0.05	0.00	0.23	0.01	0.50	0.03	0.93	×	0.30	0.14	0.22	0.02	0.16	0.08	0.88	0.00	0.00	0.33	×	0.04	0.00	0.09
0.04	-3.84	-1.33	-0.14	-1.10	-1.23	-1.34	0.05	-0.05	0.09	1.28	-0.19	0.05	0.40	-1.45	-0.25	-0.30	-1.85	-1.06	0.24	0.08	0.16	-1.29	-1.11	0.02	0.06	-0.03	-1.21	-1.25	-1.35	0.20	-0.26	-0.17	0.09	N/A	-1.28	-1.51	-0.28
1.09	1.14	0.02	1.25	0.04	0.08	0.11	1.17	1.17	1.11	0.20	1.27	1.25	1.47	0.16	1.44	1.36	0.29	0.06	1.37	1.20	1.32	0.16	0.02	1.23	1.25	1.27	0.02	0.16	0.13	1.20	1.29	1.20	1.15	N/A	0.21	0.25	1.35

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					cueR						gph-1				phnE	phoE	phnD	phoC	rpe																		
monooxygenase domain-containing protein	bile acid transporter family protein	hypothetical protein	oxidoreductase	copper-translocating P-type ATPase (EC:3.6.3.4) short chain dehvdrogenase/reductase	Cu(I)-responsive transcriptional regulator	6-phosphogluconate dehydrogenase	hypothetical protein	hypothetical protein	hypothetical protein	metallo-beta-lactamase	phosphoglycolate phosphatase (EC:3.1.3.18)	thioesterase	2-dehydro-3-deoxygalactonokinase	chloramphenicol acetyltransferase	phosphonate ABC transporter permease	phosphonate ABC transporter permease	prospionare rice nanopores substate-onitaing	phosphonate ABC transporter ATP-binding protein	ribulose-phosphate 3-epimerase (EC:5.1.3.1)	long-chain-fatty-acidCoA ligase	enoyl-CoA hydratase (EC:4.2.1.17)	hypothetical protein	acyl-CoA dehydrogenase	acyl-CoA dehydrogenase	acetyl-CoA acyltransferase/thiolase	enoyl-CoA hydratase	hypothetical protein	hypothetical protein	hypothetical protein	IcIR family transcriptional regulator	FkbM family methyltransferase	(Fe-S)-binding protein	glutamine synthetase	cytochrome P450 family protein	glutamine amidotransferase	GntR family transcriptional regulator	cyclase
-1.46	1.12	-1.00	х	1.13	1.08	-1.02	1.21	х	Х	-1.29	1.38	2.10	1.75	-1.43	-1.13	-1.92	-1.19	x	-1.13	1.17	0.98	1.06	1.17	1.49	1.31	1.99	х	-2.17	-3.02	-1.76	-1.40	-1.09	-1.38	-1.48	-1.56	-1.57	-1.94
0.62	0.70	0.54	×	0.87	0.70	0.51	0.07	×	х	0.55	0.02	0.00	0.71	0.41	0.13	0.00	0.16	×	0.07	0.87	0.87	0.99	0.79	0.00	0.86	0.00	x	0.07	0.03	0.00	0.00	0.95	0.00	0.01	0.05	0.00	0.00
-1.79	-1.35	1.01	-1.39	1.41	1.03	1.35	1.44	1.22	1.09	-1.33	1.36	1.95	1.06	1.14	-1.23	-1.14	1.10	Х	1.62	-1.01	1.04	-1.17	-1.22	-1.27	-1.10	-1.10	-1.06	-1.54	-1.98	1.01	1.01	-1.03	-1.15	-1.01	-1.06	-1.05	-2.08
0.36	0.35	0.90	0.35	0.06	0.81	0.04	0.00	0.04	0.65	0.22	0.00	0.00	0.78	0.19	0.09	0.75	0.46	×	0.01	1.00	0.89	0.66	0.32	0.00	0.59	0.67	0.71	0.46	0.00	0.89	0.96	0.95	0.18	0.92	0.68	0.61	0.00
-1.63	-0.12	0.01	N/A	1.27	1.06	0.17	1.33	N/A	N/A	-1.31	1.37	2.03	1.41	-0.15	-1.18	-1.53	-0.04	N/A	0.25	0.08	1.01	-0.05	-0.03	0.11	0.11	0.45	N/A	-1.86	-2.50	-0.38	-0.20	-1.06	-1.27	-1.25	-1.31	-1.31	-2.01
0.16	1.24	1.01	N/A	0.14	0.03	1.19	0.12	N/A	N/A	0.02	0.01	0.08	0.35	1.29	0.05	0.39	1.15	N/A	1.38	1.09	0.03	1.12	1.20	1.38	1.21	1.55	N/A	0.32	0.52	1.39	1.21	0.03	0.12	0.23	0.25	0.26	0.07
-2.25	1.71	1.26	х	1.61	1.38	3.73	3.94	-1.24	-0.93	-3.49	2.59	2.39	1.75	11.80	1.61	-1.70	-0.96	-1.58	4.09	-2.48	-1.74	-3.33	-4.10	-4.07	-2.86	-3.78	-1.69	-3.30	-7.23	1.32	1.06	-1.32	-3.03	-4.80	-5.36	1.09	-9.45
0.46	0.00	0.01	x	0.44	0.04	0.00	0.00	0.25	0.12	0.07	0.00	0.00	0.66	0.00	0.03	0.03	0.02	×	0.00	0.22	0.41	0.01	0.00	0.00	0.00	0.00	0.01	0.15	0.00	0.01	0.89	0.96	0.00	0.00	0.00	0.09	0.00
-0.98	-1.19	-1.07	-1.39	1.81	-1.04	-1.12	-1.50	x	-1.09	1.40	-1.07	1.07	-1.30	-1.05	-1.30	-1.64	-0.96	x	-1.12	1.42	1.28	1.40	1.73	1.30	1.37	1.23	Х	-1.07	1.11	-1.27	-1.14	1.35	1.13	-1.14	-1.33	1.06	1.08
0.95	0.58	0.74	0.31	0.00	0.92	0.12	0.00	×	0.81	0.09	0.21	0.23	0.34	x	0.47	0.45	0.35	×	0.07	0.38	0.46	0.15	0.07	0.04	0.06	0.41	×	0.89	0.25	0.03	0.30	0.15	0.19	0.24	0.06	0.32	0.55
-1.61	0.26	0.10	N/A	1.71	0.17	1.31	1.22	N/A	-1.01	-1.05	0.76	1.73	0.23	5.38	0.16	-1.67	-0.96	N/A	1.49	-0.53	-0.23	-0.97	-1.19	-1.39	-0.75	-1.28	N/A	-2.19	-3.06	0.03	-0.04	0.02	-0.95	-2.97	-3.35	1.08	-4.19
0.64	1.45	1.17	N/A	0.10	1.21	2.43	2.72	N/A	0.08	2.45	1.83	0.66	1.53	6.43	1.46	0.03	0.00	N/A	2.61	1.95	1.51	2.37	2.92	2.69	2.12	2.51	N/A	1.12	4.17	1.30	1.10	1.34	2.08	1.83	2.02	0.02	5.27
-1.24	1.18	-0.97	-1.09	-1.10	-1.12	1.00	1.35	-1.00	-1.13	1.12	-1.38	-1.23	-1.12	-1.13	-1.30	-1.27	1.20	х	-1.32	1.13	1.25	1.22	1.29	1.15	1.26	1.27	1.22	-0.98	-1.32	-1.51	1.27	1.13	-1.15	-1.31	-1.43	1.27	-1.29
0.66	0.42	0.55	0.85	0.30	0.53	0.85	0.00	0.76	0.31	0.09	0.01	0.09	0.85	0.69	0.03	0.69	0.03	x	0.02	0.75	0.45	0.51	0.39	0.14	0.25	0.19	0.05	0.94	0.07	0.00	0.23	0.47	0.10	0.02	0.00	0.04	0.04
1.27	1.24	-1.09	1.37	-1.03	1.42	-1.08	-1.31	x	1.72	1.59	-1.69	-1.90	-1.04	Х	-0.92	1.14	-1.11	×	-1.84	1.57	1.74	1.35	1.50	1.44	2.24	1.74	х	1.43	1.16	-1.83	0.98	-1.28	-1.35	-1.14	-1.00	-0.91	1.09
0.68	0.47	0.78	0.62	0.83	0.20	0.19	0.01	x	0.19	0.14	0.00	0.00	0.98	x	0.42	0.86	х	х	0.00	0.15	0.12	0.25	0.16	0.00	0.00	0.05	x	0.43	0.09	0.00	0.64	0.25	0.28	0.98	0.23	0.09	0.60
0.02	1.21	-1.03	0.14	-1.07	0.15	-0.04	0.02	N/A	0.30	1.36	-1.54	-1.57	-1.08	N/A	-1.11	-0.07	0.04	N/A	-1.58	1.35	1.50	1.29	1.40	1.30	1.75	1.51	N/A	0.22	-0.08	-1.67	1.12	-0.08	-1.25	-1.23	-1.22	0.18	-0.10
1.26		0.06		0.04	1.27	1.04	1.33	N/A	1.43	0.24	0.16	0.34	0.04	N/A	0.19	1.21	1.16	N/A	0.26	0.22	0.24	0.07	0.11	0.15	0.49	0.24	N/A	1.21	1.24	0.16	0.15	1.21	0.10	0.09	_		1.19

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SPO0835	SPO0834	SPO0833	SPO0832	SPO0831	SPO0830	SPO0829	SPO0828	SPO0827	SPO0826	SPO0825	SPO0824	SPO0823	SPO0822	SPO0821	SPO0820	SPO0819	SPO0818	SPO0817	SPO0816	SPO0815	SPO0814	SPO0813	SPO0812	SPO0811	SPO0810	SPO0809	SPO0808	SPO0807	SPO0806	SPO0805	SPO0804	SPO0803	SPO0802	SPO0801	SPO0800	SPO0799
																		ychF		trpA			rplY	pth			trpB		trpF		ihfB		rpsA	fadD		
cyclase	formate dehydrogenase subunit alpha (EC:1.2.1.2)	formate dehydrogenase subunit beta (EC:1.2.1.2)	LysR family transcriptional regulator	xanthine dehydrogenase, medium subunit	xanthine dehydrogenase, small/large subunits	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	substrate-binding protein	branched-chain amino acid ABC transporter permease hear hed a bring amino acid ABC transporter	binding protein	binding protein	MarR family transcriptional regulator	CAIB/BAIF family protein	amidohydrolase	MAPEG family protein	GTP-dependent nucleic acid-binding protein EngD	hypothetical protein	tryptophan synthase subunit alpha (EC:4.2.1.20)	acyl dehydratase MaoC	L-lactate dehydrogenase	50S ribosomal protein L25	peptidyl-tRNA hydrolase (EC:3.1.1.29)	6-aminohexanoate-dimer hydrolase	hypothetical protein	tryptophan synthase subunit beta (EC:4.2.1.20)	hypothetical protein	N-(3-phosphorioosyi)anthranilate isometase (EC:5.3.1.24)	hypothetical protein	integration host factor subunit beta	NUDIX domain-containing protein (EC:6)	30S ribosomal protein S1	4-coumarateCoA ligase (EC:6.2.1.12)	choline sulfatase	TetR family transcriptional regulator
1.88	3.05	3.81	x	-2.06	-1.99	-1.79	-1.74	-1.19	-1.42	-2.44	-1.48	1.06	-1.35	1.08	1.25	0.98	-0.96	-0.99	-1.94	1.91	-1.45	1.43	-1.23	-1.35	1.48	1.33	1.96	1.45	1.62	1.45	-1.68	-1.30	-1.33	1.35	-0.99	-1.67
0.00	0.00	0.00	x	0.01	0.00	0.03	0.00	0.33	0.00	0.13	0.33	0.28	0.00	0.98	0.00	0.19	0.09	0.98	×	0.00	0.01	0.11	0.48	0.21	0.73	0.08	0.17	0.04	0.01	0.00	0.00	0.00	0.55	0.46	0.91	0.26
1.26	1.36	1.20	1.07	-1.17	1.03	1.10	1.51	1.25	1.25	-1.48	-1.16	1.02	1.09	1.25	1.25	2.14	1.27	1.38	-1.39	1.18	1.09	1.06	1.03	-1.08	1.04	1.20	1.27	1.06	1.53	1.67	1.50	-1.18	1.11	-1.08	-1.14	-1.02
0.02	0.04	0.21	0.12	0.01	0.87	0.68	0.00	0.01	0.00	0.06	0.43	0.73	0.02	0.56	0.11	0.01	0.00	0.10	0.24	0.14	0.32	0.68	0.92	0.46	0.88	0.13	0.23	0.81	0.01	0.00	0.00	0.48	0.72	0.58	0.75	0.99
1.57	2.21	2.51	N/A	-1.62	-0.48	-0.35	-0.12	0.03	-0.09	-1.96	-1.32	1.04	-0.13	1.17	1.25	1.56	0.16	0.19	-1.67	1.55	-0.18	1.25	-0.10	-1.22	1.26	1.27	1.62	1.26	1.58	1.56	-0.09	-1.24	-0.11	0.14	-1.07	-1.35
0.31	0.85	1.31	N/A	0.45	1.51	1.45	1.63	1.22	1.34	0.48	0.16	0.02	1.22	0.09	0.00	0.58	1.11	1.19	0.27	0.37	1.27	0.18	1.13	0.14	0.22	0.07	0.35	0.20	0.05	0.11	1.59	0.06	1.22	1.22	0.08	0.33
1.71	1.65	2.09	×	-4.29	-8.22	-2.98	-4.39	-3.13	-4.23	-11.80	-2.20	-2.19	-3.02	1.56	2.47	-8.19	2.00	4.05	-1.44	4.09	1.12	0.93	3.95	1.87	0.86	-1.73	2.75	-1.43	1.51	-1.48	-2.57	-1.28	3.87	-2.40	-2.22	-1.79
0.00	0.01	0.00	×	0.00	0.00	0.03	0.00	0.00	0.00	0.00	0.06	0.00	0.00	0.68	0.00	0.00	0.00	0.10	0.68	0.00	0.01	0.50	0.00	0.00	0.70	0.01	0.01	0.27	0.00	0.00	0.00	0.04	0.04	0.06	0.00	0.45
-1.05	-6.36	-7.13	1.23	-1.23	1.10	1.10	1.17	1.32	1.39	1.42	-1.19	-1.18	-1.22	-1.36	-1.47	1.29	-1.12	-1.54	-1.70	-1.27	1.48	2.85	-1.29	-1.23	1.11	1.26	1.12	-1.00	-1.17	1.23	1.36	-1.07	-1.98	1.48	1.60	-1.26
0.13	0.00	0.00	0.07	0.28	0.69	0.66	0.24	0.02	0.03	0.01	0.37	0.23	0.12	0.29	0.01	0.45	0.14	0.02	0.17	0.01	0.00	0.00	0.00	0.03	0.17	0.01	0.61	0.86	0.05	0.02	0.00	0.67	0.00	0.04	0.07	0.23
0.33	-2.36	-2.52	N/A	-2.76	-3.56	-0.94	-1.61	-0.91	-1.42	-5.19	-1.70	-1.69	-2.12	0.10	0.50	-3.45	0.44	1.26	-1.57	1.41	1.30	1.89	1.33	0.32	0.99	-0.24	1.94	-1.21	0.17	-0.13	-0.61	-1.18	0.95	-0.46	-0.31	-1.53
1.38	4.01	4.61	N/A	1.53	4.66	2.04	2.78	2.23	2.81	6.61	0.51	0.50	0.90	1.46	1.97	4.74	1.56	2.80	0.13	2.68	0.18	0.96	2.62	1.55	0.12	1.50	0.82	0.22	1.34	1.36	1.97	0.11	2.93	1.94	1.91	0.27
1.02	-1.21	-1.33	1.20	-0.97	-1.15	-0.98	-0.99	-1.06	-1.16	1.04	1.12	1.41	1.37	1.18	-1.71	1.16	-1.03	-1.91	-1.14	-1.38	-1.03	-1.10	-1.25	-1.37	1.13	1.10	1.00	1.30	1.52	1.69	1.15	1.05	1.05	-1.17	-1.17	1.14
0.94	0.02	0.01	0.03	0.70	0.43	0.86	0.32	0.35	0.01	0.58	0.53	0.03	0.01	0.70	0.00	0.10	0.84	0.00	0.69	0.02	0.67	0.45	0.00	0.00	0.38	0.13	0.81	0.02	0.00	0.00	0.19	0.85	0.98	0.26	0.18	0.66
1.02	-1.44	1.18	-1.13	1.60	1.43	1.18	Х	-1.37	-1.08	1.34	1.19	1.22	-0.98	1.01	-2.83	1.16	-1.68	-2.45	1.21	-2.04	-1.06	1.18	-1.36	-1.67	1.73	1.08	-1.18	-1.06	-1.09	1.49	1.31	1.46	-1.50	1.22	-0.93	-1.14
0.44	0.01	0.31	x	0.00	0.16	0.29	x	0.18	0.84	0.01	0.32	0.06	0.34	0.98	0.00	х	0.02	0.00	0.42	0.00	0.78	0.60	0.03	0.00	0.00	0.07	0.13	х	0.66	0.00	0.00	0.19	0.01	0.10	0.74	0.94
1.02	-1.33	-0.08	0.04	0.32	0.14	0.10	N/A	-1.22	-1.12	1.19	1.16	1.32	0.19	1.10	-2.27	1.16	-1.36	-2.18	0.04	-1.71	-1.05	0.04	-1.31	-1.52	1.43	1.09	-0.09	0.12	0.22	1.59	1.23	1.26	-0.23	0.03	-1.05	0.00
0.00	0.12	1.26	1.17	1.28	1.29	1.08	N/A	0.16	0.04	0.15	0.03	0.10	1.18	0.09		0.00	0.32	0.27	1.18	0.33	0.02	1.14	0.06	0.15	0.30	0.01	1.09	1.18	1.31	0.10	0.08	0.21	1.28	1.20	0.12	1.14

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SPO0874	SPO0873	SPO0872	SPO0871	SPO0870	SPO0869	SPO0868	SPO0867	SPO0866	SPO0865	SPO0864	SPO0863	SPO0862	SPO0861	SPO0860	SPO0859	SPO0858	SPO0857	SPO0856	SPO0855	SPO0854	SPO0853	SPO0852	SPO0851	SPO0850	SPO0849	SPO0848	SPO0847	SPO0846	SPO0845	SPO0844	SPO0843	SPO0842	SPO0841	SPO0840	SPO0839	SPO0838	SPO0837	SPO0836
	allA										xylG	xylH	xylF				galM	xylA	xylB																			
xanthine/uracil permease	ureidoglycolate hydrolase (EC:3.5.3.19)	polysaccharide deacetylase	transthyretin family protein	LysR family transcriptional regulator	hypothetical protein	OMP85 family outer membrane protein	hypothetical protein	gamma-glutamyltranspeptidase	Gfo/Idh/MocA family oxidoreductase	glucokinase	xylose ABC transporter ATP-binding protein	xylose ABC transporter permease	xylose ABC transporter substrate-binding protein	xylose repressor	hypothetical protein	methylamine utilization protein MauG	aldose 1-epimerase (EC:5.1.3.3)	xylose isomerase (EC:5.3.1.5)	xylulokinase (EC:2.7.1.17)	glycoside hydrolase	glycosyl transferase family protein	Gfo/Idh/MocA family oxidoreductase	WecB/TagA/CpsF family glycosyl transferase	hypothetical protein	non-ribosomal peptide synthase	glycosyl transferase family protein	non-ribosomal peptide synthetase	phosphopantetheinyl transferase PptA	hypothetical protein	hypothetical protein	glycosyl transferase family protein	glycoside hydrolase	polysaccharide biosynthesis protein	chain length determinant protein	protein	sugar transferase exonolysarcharide biosynthesis domain-containing	hypothetical protein	hypothetical protein
1.60	1.47	1.43	-1.13	-1.14	-1.27	1.01	1.72	1.71	1.04	1.21	1.04	1.75	1.17	0.97	-1.10	2.05	-1.18	-1.25	-1.69	1.02	-1.20	1.10	-1.20	0.99	1.04	-1.82	1.16	1.21	-1.14	-1.02	1.08	-0.94	-1.11	0.99	2.96	20.60	2.07	1.13
0.02	0.52	0.11	0.88	0.50	0.08	0.76	0.02	0.00	0.87	0.58	0.89	0.05	0.91	0.19	0.13	0.01	0.57	0.79	0.01	0.85	0.31	0.37	0.79	0.83	0.97	0.00	0.51	0.89	0.14	0.98	0.64	0.26	0.63	0.84	0.00	0.00	0.00	0.85
-1.06	1.79	1.42	1.51	-1.38	-1.27	-1.06	1.04	1.04	-1.10	1.28	-1.12	-1.22	-2.16	1.15	1.12	-1.10	-1.17	-1.22	-1.60	1.12	1.14	1.41	1.18	1.25	1.37	1.07	1.02	1.15	-1.05	1.31	1.71	1.19	1.41	1.15	2.17	3.27	2.09	1.32
0.59	0.02	0.06	0.27	0.01	0.74	0.79	0.56	0.66	0.90	0.22	0.17	0.66	0.00	0.21	0.49	0.82	0.62	0.53	0.23	0.28	0.14	0.25	0.66	0.56	0.23	0.77	0.91	0.59	0.58	0.04	0.05	0.55	0.00	0.32	0.00	0.00	0.00	0.10
0.27	1.63	1.43	0.19	-1.26	-1.27	-0.03	1.38	1.38	-0.03	1.25	-0.04	0.27	-0.50	1.06	0.01	0.48	-1.18	-1.24	-1.65	1.07	-0.03	1.26	-0.01	1.12	1.21	-0.38	1.09	1.18	-1.10	0.15	1.40	0.13	0.15	1.07	2.57	11.94	2.08	1.23
1.33	0.16	0.01	1.32	0.12	0.00	1.04	0.34	0.34	1.07	0.04	1.08	1.49	1.67	0.09	1.11	1.58	0.01	0.02	0.04	0.05	1.17	0.16	1.19	0.13	0.17	1.45	0.07	0.03	0.04	1.17	0.31	1.06	1.26	0.08	0.39	8.67	0.01	0.10
-2.42	-2.45	-2.40	-1.69	1.18	-1.84	-1.28	1.55	2.32	-1.91	1.31	-2.90	2.21	-8.32	-1.34	-1.45	2.03	-1.45	-3.10	-3.19	-1.39	-1.88	-1.26	1.11	-2.24	-1.28	-3.45	-2.14	-1.15	-1.17	-1.08	-1.00	-0.91	-1.00	-2.08	-2.90	2.07	-1.08	1.18
0.00	0.05	0.01	0.72	0.86	0.28	0.19	0.02	0.01	0.32	0.61	0.09	0.00	0.01	0.07	0.08	0.01	0.49	0.00	0.01	0.02	0.02	0.51	0.90	0.21	0.83	0.00	0.00	0.93	0.31	0.50	0.35	0.04	0.47	0.00	0.00	0.00	0.40	0.60
1.09	1.08	1.20	1.14	-1.32	1.18	-1.08	1.37	-1.01	-1.06	-1.37	-1.08	-1.18	-1.02	1.28	1.20	-1.16	-1.21	-1.42	-1.37	-1.03	-1.00	1.03	-1.40	1.24	1.16	1.11	1.22	1.21	-1.00	-1.06	-1.09	-1.41	1.06	1.10	1.22	-1.02	1.58	1.09
0.47	0.63	0.25	0.71	0.23	0.58	0.64	0.04	0.93	0.94	0.31	0.75	0.46	0.57	0.06	0.22	x	0.49	0.41	0.22	0.63	0.91	0.69	0.51	0.47	0.69	0.59	0.06	0.37	0.86	0.56	0.85	0.18	0.31	0.25	0.24	0.98	0.00	0.49
-0.67	-0.69	-0.60	-0.28	-0.07	-0.33	-1.18	1.46	0.66	-1.49	-0.03	-1.99	0.52	-4.67	-0.03	-0.13	0.44	-1.33	-2.26	-2.28	-1.21	-1.44	-0.12	-0.15	-0.50	-0.06	-1.17	-0.46	0.03	-1.09	-1.07	-1.05	-1.16	0.03	-0.49	-0.84	0.53	0.25	1.14
1.76	1.77	1.80	1.42	1.25	1.51	0.10	0.09	1.67	0.43	1.34	0.91	1.70	3.65	1.31	1.33	1.60	0.12	0.84	0.91	0.18	0.44	1.15	1.26	1.74	1.22	2.28	1.68	1.18	0.09	0.01	0.05	0.25	1.03	1.59	2.06	1.55	1.33	0.04
1.19	-1.17	-1.25	-0.97	-1.09	-1.32	1.04	-1.06	-1.00	-1.00	1.20	-1.05	-1.28	1.11	1.25	1.08	-1.33	-1.15	-1.01	-1.43	-1.98	-2.02	-1.53	1.49	-1.44	-1.15	-1.86	-1.60	-1.32	-1.40	1.15	1.78	1.40	1.15	-1.00	-1.27	-1.17	-1.16	-1.13
0.04	0.34	0.16	0.82	0.69	0.79	0.88	0.29	0.96	0.98	0.45	0.83	0.74	0.50	0.11	0.13	0.14	0.62	0.92	0.27	0.01	0.01	0.06	0.46	0.12	0.52	0.12	0.01	0.16	0.01	0.06	0.06	0.18	0.07	0.77	0.17	0.10	0.19	0.25
1.76	-1.53	-1.57	-1.54	1.34	-1.09	1.11	-1.02	-1.46	1.30	-0.93	-1.00	1.10	-1.07	-1.15	-1.28	х	1.35	-1.06	1.24	-2.45	-1.55	-1.58	-0.97	-1.17	-1.17	-1.25	-1.43	-1.17	1.04	-0.95	-1.00	-1.07	1.16	1.16	-1.18	-1.84	-1.40	0.98
0.05	0.10	0.05	0.36	0.07	0.84	0.44	0.53	0.09	0.77	0.51	0.83	0.87	0.50	0.28	0.10	x	0.28	0.76	0.60	0.01	0.11	0.16	0.87	0.51	0.65	0.62	0.00	0.70	0.65	0.09	0.54	0.75	0.07	0.17	0.37	0.05	0.44	0.81
1.48	-1.35	-1.41	-1.25	0.13	-1.21	1.08	-1.04	-1.23	0.15	0.14	-1.03	-0.09	0.02	0.05	-0.10	N/A	0.10	-1.04	-0.10	-2.22	-1.79	-1.56	0.26	-1.31	-1.16	-1.56	-1.52	-1.25	-0.18	0.10	0.39	0.17	1.16	0.08	-1.23	-1.51	-1.28	-0.08
0.29	0.18	0.16	0.29	1.22	0.12	0.04	0.02	0.23	1.15	1.07	0.03	1.19	1.09	1.20	1.18	N/A	1.25	0.03	1.34	0.24	0.24	0.03	1.23	0.14	0.01	0.30	0.09	0.08	1.22	1.05	1.39	1.24	0.01	1.08	0.05	0.34	0.12	1.05

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SPO0911	SPO0910	SPO0909	SPO0908	SPO0907	SPO0906	SPO0905	SPO0904	SPO0903	SPO0902	SPO0901	SPO0900	SPO0899	SPO0898	SPO0897	SPO0896	SPO0895	SPO0894	SPO0893	SPO0892	SPO0891	SPO0890	SPO0889	SPO0888	SPO0887	SPO0886	SPO0885	SPO0884	SPO0883	SPO0882	SPO0881	SPO0880	SPO0879	SPO0878	SPO0877	SPO0876	SPO0875
proC			lgt					trxB			sat							purE	purK					groEL	groES								gcvH-1			gap-2
pyrroline-5-carboxylate reductase (EC:1.5.1.2)	hypothetical protein	hypothetical protein	prolipoprotein diacylglyceryl transferase (EC:2.4.99)	hypothetical protein	hypothetical protein	hypothetical protein	leucine-responsive regulatory protein	thioredoxin-disulfide reductase (EC:1.8.1.9)	hypothetical protein	guanylate cyclase	bitunctional sulfate adenylyltransferase subunit l/adenylylsulfate kinase (EC:2.7.1.25 2.7.7.4)	hypothetical protein	hypothetical protein	PhzF family phenazine biosynthesis protein	hypothetical protein	heat shock protein 20	hypothetical protein	pnospnoritos ytaminoimidazoje carboxytase catalytic subunit (EC:4.1.1.21)	ATPase subunit (EC:4.1.1.21)	alkylphosphonate utilization protein PhnM	lipase	hypothetical protein	acetyltransferase	molecular chaperone GroEL	co-chaperonin GroES	alpha/beta hydrolase	hypothetical protein	acyl-CoA dehydrogenase	glycine cleavage system H protein	hypothetical protein	hypothetical protein	glyceraldehyde-3-phosphate dehydrogenase, type I (EC:1.2.1)				
2.19	1.96	1.52	1.06	-1.33	-1.78	3.58	-2.25	0.98	1.39	-1.04	1.31	2.38	1.18	1.07	1.23	-1.27	-1.47	1.95	1.40	Х	Х	1.04	1.42	1.06	-1.27	-1.19	-1.07	1.32	-4.40	-2.70	1.08	2.09	1.58	-1.14	1.43	1.44
0.18	0.11	0.77	0.99	0.17	0.06	0.00	0.01	0.45	0.90	0.66	0.67	0.46	0.11	0.63	0.61	0.06	0.00	0.03	0.07	х	х	0.97	0.17	0.81	0.04	0.78	0.10	0.15	0.12	0.26	0.99	0.08	0.79	0.81	0.26	0.00
1.28	1.60	1.34	-1.71	-1.22	1.09	1.81	-1.36	1.06	1.28	-1.08	0.98	1.60	-1.04	1.14	-1.32	-1.95	1.21	1.48	1.85	1.11	1.18	1.02	-1.15	-1.96	-3.46	-1.03	1.22	1.08	1.12	-1.04	1.13	1.01	1.09	1.45	-1.77	-1.20
0.37	0.14	0.34	0.01	0.44	0.63	0.01	0.10	0.63	0.16	0.90	0.89	0.02	0.91	0.40	0.22	0.00	0.30	0.09	0.01	0.28	0.11	0.98	0.53	0.04	0.01	0.94	0.33	0.32	0.86	0.78	0.57	0.92	0.68	0.32	0.10	0.48
1.74	1.78	1.43	-0.33	-1.28	-0.35	2.70	-1.81	1.02	1.34	-1.06	1.14	1.99	0.07	1.11	-0.05	-1.61	-0.13	1.72	1.63	N/A	N/A	1.03	0.14	-0.45	-2.37	-1.11	0.08	1.20	-1.64	-1.87	1.11	1.55	1.34	0.16	-0.17	0.12
0.46	0.18	0.09	1.39	0.06	1.44	0.88	0.45	0.04	0.05	0.02	0.17	0.39	1.11	0.03	1.28	0.34	1.34	0.24	0.23	N/A	N/A	0.01	1.29	1.51	1.10	0.08	1.15	0.12	2.76	0.83	0.02	0.54	0.25	1.30	1.60	1.32
1.80	-2.27	1.77	1.53	-1.13	-1.47	-1.23	1.49	2.22	2.42	-2.28	7.68	-1.57	1.29	1.71	1.63	1.29	-2.39	5.06	2.36	2.88	5.62	1.48	1.88	7.89	5.38	1.38	1.36	2.29	-8.69	-2.16	-1.73	-1.44	1.01	-1.04	-3.15	1.48
0.59	0.01	0.62	0.05	0.68	0.24	0.14	0.03	0.02	0.68	0.00	0.05	0.08	0.08	0.02	0.04	0.46	0.00	0.00	0.01	0.00	0.00	0.90	0.09	0.00	0.01	0.68	0.24	0.01	0.05	0.07	0.09	0.04	0.82	0.91	0.02	0.04
1.29	2.29	-1.07	-1.33	1.15	1.21	-1.01	-1.06	-1.16	-1.17	1.46	1.44	1.94	-1.41	-1.20	-1.24	-1.27	1.34	-1.03	-1.06	-1.52	-1.56	-1.04	1.06	-9.38	-10.60	1.07	1.05	1.19	-1.18	-1.40	1.91	1.85	1.44	1.32	-1.32	-1.21
0.35	0.00	0.06	0.11	0.37	0.13	0.77	0.22	0.13	0.13	0.04	0.14	0.01	0.21	0.08	0.11	0.24	0.16	0.44	0.48	0.02	0.01	0.90	0.86	0.01	0.00	0.91	0.94	0.37	0.88	0.13	0.00	0.04	0.19	0.45	0.37	0.03
1.55	0.01	0.35	0.10	0.01	-0.13	-1.12	0.22	0.53	0.63	-0.41	4.56	0.19	-0.06	0.26	0.20	0.01	-0.53	2.02	0.65	0.68	2.03	0.22	1.47	-0.75	-2.61	1.23	1.21	1.74	-4.94	-1.78	0.09	0.21	1.23	0.14	-2.24	0.14
0.26	2.28	1.42	1.43	1.14	1.34	0.11	1.28	1.69	1.80	1.87	3.12	1.76	1.35	1.46	1.44	1.28	1.87	3.05	1.71	2.20	3.59	1.26	0.41	8.64	7.99	0.15	0.16	0.55	3.76	0.38	1.82	1.65	0.21	1.18	0.92	1.35
-1.25	1.41	1.55	1.20	-1.19	1.05	-1.70	-1.13	1.25	1.32	1.34	-1.32	1.48	1.22	1.03	1.61	1.40	1.04	-1.13	1.07	-1.87	-1.57	1.21	1.73	1.97	-1.59	1.08	1.91	1.55	-1.04	1.06	1.10	1.50	1.37	1.53	1.27	-1.35
0.15	0.03	0.19	0.66	0.29	0.71	0.01	0.26	0.22	0.09	0.08	0.15	0.05	0.19	0.91	0.01	0.05	0.83	0.02	0.28	0.09	0.02	0.77	0.03	0.00	0.07	0.89	0.06	0.12	0.99	0.79	0.17	0.20	0.15	0.28	0.49	0.08
-1.10	1.83	1.26	1.30	1.25	-1.19	-1.46	-1.42	1.23	1.07	1.16	-1.91	1.57	1.38	-1.06	1.49	1.39	-1.12	-1.83	-1.68	-1.92	-2.40	1.80	1.16	1.47	-1.70	1.13	1.43	1.39	-1.12	-1.44	1.86	2.85	1.32	-1.11	1.32	-1.38
0.36	0.03	0.33	0.42	0.42	0.08	0.08	0.06	0.48	0.94	0.36	0.10	0.03	0.39	0.27	0.06	0.03	0.35	0.02	0.04	0.21	0.03	0.37	0.23	0.33	0.04	0.65	0.12	0.11	х	0.03	0.01	0.02	0.06	0.98	0.53	0.02
-1.18	1.62	1.41	1.25	0.03	-0.07	-1.58	-1.28	1.24	1.20	1.25	-1.62	1.53	1.30	-0.02	1.55	1.40	-0.04	-1.48	-0.31	-1.90	-1.99	1.51	1.45	1.72	-1.65	1.11	1.67	1.47	-1.08	-0.19	1.48	2.18	1.35	0.21	1.30	-1.37
0.08	0.21	0.15	0.05	1.22	1.12	0.12	0.14	0.01	0.13	0.09	0.30	0.05	0.08	1.05	0.06	0.01	1.08	0.35	1.38	0.02	0.42	0.30	0.29	0.25	0.05	0.02	0.24	0.08	0.04	1.25	0.38	0.68	0.03	1.32	0.03	0.01

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SPO0949	SPO0948	SPO0947	SPO0946	SPO0945	SPO0944	SPO0943	SPO0942	SPO0941	SPO0940	SPO0939	SPO0938	SPO0937	SPO0936	SPO0935	SPO0934	SPO0933	SPO0932	SPO0931	SPO0930	SPO0929	SPO0928	SPO0927	SPO0926	SPO0925	SPO0924	SPO0923	SPO0922	SPO0921	SPO0920	SPO0919	SPO0918	SPO0917	SPO0916	SPO0915	SPO0914	SPO0913	SPO0912
kpsT	kpsE	kdsA	algC		rimO																		aspS		pabB	carB					fabH-1				tdk		csaA
capsutar polysaccnaride export ATF-binding protein (EC:3.6.3.38)	capsule polysaccharide exporter	2-чепушо-э-чеохурнозрноостопате атиолазе (EC:2.5.1.55)	2 debudeo 3 deoxymboschoootomate addolase	AsmA family protein	30S ribosomal protein S12 methylthiotransferase	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	M48 family peptidase	hypothetical protein	hypothetical protein	hypothetical protein	nitroreductase	hypothetical protein	hypothetical protein	methylmalonyl-CoA epimerase (EC:5.1.99.1)	response regulator	hypothetical protein	type II DNA modification methyltransferase	hypothetical protein	hypothetical protein	aspartyl-tRNA synthetase (EC:6.1.1.12)	hypothetical protein	aminodeoxychorismate synthase (EC:2.6.1.85)	carbamoyl phosphate synthase large subunit (EC:6.3.5.5)	hypothetical protein	AcrB/AcrD/AcrF family transporter	RND family efflux transporter MFP subunit	MarR family transcriptional regulator	3-oxoacyl-ACP synthase (EC:2.3.1.41)	glyoxalase	hypothetical protein	hypothetical protein	thymidine kinase (EC:2.7.1.21)	D-isomer specific 2-hydroxyacid dehydrogenase	chaperonin csaA
1.06	1.39	1.31	1.35	1.29	-1.30	1.46	1.65	1.91	1.45	1.34	-1.33	1.11	1.02	1.00	1.11	3.59	5.45	1.45	1.94	-2.33	-1.57	1.15	-1.20	1.10	-1.48	1.63	-1.11	1.59	0.98	1.11	-1.32	х	1.17	1.01	-1.16	1.60	2.00
0.84	0.03	0.40	0.11	0.24	0.86	0.68	0.26	0.01	0.06	0.71	0.06	0.93	0.03	0.47	0.90	0.00	0.00	0.07	0.00	0.00	0.02	x	×	0.87	0.10	0.07	0.43	0.16	0.09	0.31	0.00	x	0.70	0.10	0.35	0.01	0.02
-1.54	-1.09	-1.29	1.07	-1.34	-1.02	1.13	2.01	1.45	1.20	-1.09	-1.34	1.18	1.12	-1.33	1.08	2.08	1.76	1.08	1.51	-1.23	-1.20	1.42	1.30	1.37	1.16	1.41	-1.10	-1.18	-1.10	-1.42	-1.41	-1.07	1.21	-1.05	-1.02	1.30	1.09
0.30	0.25	0.28	0.54	0.21	0.94	0.54	0.14	0.03	0.40	0.32	0.16	0.69	0.06	0.23	0.66	0.10	0.04	0.75	0.04	0.63	0.21	0.19	0.06	0.09	0.20	0.10	0.48	0.61	0.37	0.40	0.18	0.68	0.66	0.47	0.79	0.03	0.34
-0.24	0.15	0.01	1.21	-0.03	-1.16	1.30	1.83	1.68	1.33	0.13	-1.34	1.15	1.07	-0.17	1.10	2.84	3.61	1.27	1.73	-1.78	-1.39	1.29	0.05	1.24	-0.16	1.52	-1.11	0.21	-0.06	-0.16	-1.37	N/A	1.19	-0.02	-1.09	1.45	1.55
1.30	1.24	1.30	0.14	1.32	0.14	0.16	0.18	0.23	0.13	1.22	0.01	0.03	0.05	1.16	0.02	0.76	1.85	0.19	0.21	0.55	0.19	0.14	1.25	0.14	1.32	0.11	0.01	1.39	1.04	1.27	0.04	N/A	0.02	1.03	0.07	0.15	0.46
-1.30	-1.11	1.87	1.78	0.88	1.40	1.24	2.89	8.20	1.04	0.96	1.00	-1.03	-4.41	1.71	1.80	5.18	8.10	1.53	-2.43	-1.74	-2.46	4.69	8.33	1.71	5.09	2.03	-1.64	2.55	-1.73	2.61	0.89	-0.85	1.34	-1.74	4.75	1.56	2.73
0.35	0.78	0.10	0.01	0.01	0.80	0.65	0.24	0.00	0.82	0.71	0.16	0.89	0.00	0.10	0.33	0.01	0.00	0.08	0.01	0.02	0.01	0.02	0.00	0.47	0.00	0.01	0.02	0.03	0.03	0.00	0.05	0.04	0.30	0.01	0.01	0.07	0.00
1.13	- 1.05	1.14	1.00	1.32	-1.22	2.23	1.39	-1.25	1.11	1.12	1.36	-1.00	1.61	-1.84	-1.55	1.32	-1.08	1.54	2.28	1.43	-1.03	-1.79	-1.97	-1.10	-1.59	-1.12	1.40	-1.41	-1.15	-1.21	1.70	1.10	1.21	1.08	-1.36	-1.10	1.24
0.84	0.45	0.64	0.76	0.21	0.43	0.02	0.52	0.23	0.60	0.83	0.22	0.99	0.01	0.00	0.02	0.05	0.14	0.18	0.00	0.33	0.73	0.02	0.05	0.66	0.04	0.12	0.32	0.07	0.28	0.36	0.04	0.39	0.27	0.94	0.03	0.33	0.21
-0.09	-1.08	1.51	1.39	1.10	0.09	1.74	2.14	3.48	1.08	1.04	1.18	-1.01	-1.40	-0.07	0.13	3.25	3.51	1.54	-0.08	-0.16	-1.75	1.45	3.18	0.31	1.75	0.46	-0.12	0.57	-1.44	0.70	1.30	0.12	1.28	-0.33	1.70	0.23	1.99
1.22	0.03	0.37	0.39	0.22	1.31	0.50	0.75	4.73	0.04	0.08	0.18	0.02	3.01	1.78	1.68	1.93	4.59	0.01	2.36	1.59	0.72	3.24	5.15	1.41	3.34	1.58	1.52	1.98	0.29	1.91	0.40	0.98	0.07	1.41	3.06	1.33	0.75
-1.54	-1.72	-1.43	1.11	1.81	-1.63	1.44	1.23	1.42	1.16	-1.11	1.33	1.41	-1.38	1.17	1.43	-1.19	-1.47	-1.35	1.42	-1.02	-1.22	1.00	-1.07	-1.36	-1.22	-1.26	-1.04	-1.14	-1.22	-1.34	-1.14	-1.08	1.25	-1.06	-1.63	1.14	1.05
0.24	0.04	0.02	0.33	0.03	0.22	0.27	0.48	0.03	0.22	0.65	0.08	0.50	0.04	0.24	0.05	0.09	0.01	0.05	0.07	0.85	0.10	0.79	0.64	0.06	0.17	0.04	0.66	0.26	0.06	0.17	0.16	0.64	0.26	0.52	0.01	0.06	0.91
-1.32	-2.02	-1.22	-1.18	2.26	-1.95	1.33	-1.09	-1.13	-1.05	1.00	1.21	-1.13	-1.45	1.48	1.21	-1.48	-1.74	1.53	2.31	1.00	-1.00	-2.34	-2.38	-1.75	-2.27	-1.55	1.11	-1.29	-1.77	-1.28	1.57	-1.14	1.13	1.01	-2.69	-1.10	-1.14
0.32	0.02	0.11	0.24	0.11	0.18	0.30	х	0.37	0.58	0.80	0.19	0.91	0.12	0.24	0.30	0.15	0.02	0.07	0.03	0.94	0.97	0.02	0.00	0.05	0.02	0.05	0.59	0.26	0.06	0.63	0.01	х	0.75	0.26	0.01	0.24	0.04
-1.43	-1.87	-1.33	-0.03	2.04	-1.79	1.39	0.07	0.15	0.05	-0.06	1.27	0.14	-1.42	1.33	1.32	-1.34	-1.61	0.09	1.87	-0.01	-1.11	-0.67	-1.73	-1.56	-1.75	-1.41	0.04	-1.22	-1.50	-1.31	0.22	-1.11	1.19	-0.03	-2.16	0.02	-0.04
0.11	0.15	0.11	1.15	0.22	0.16	0.05	1.16	1.28	1.11	1.05	0.06	1.27	0.04	0.15	0.11	0.14	0.14	1.44	0.45	1.01	0.11	1.67	0.65	0.19	0.52	0.15	1.08	0.08	0.27	0.03	1.36	0.03	0.06	1.04	0.53	1.12	1.10

## Chapter 10: Appendix

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	znuA	zur	znuC	znuB	lepA		hisD							rpmB		hrpB		apqZ			panE-1		argC-1	argF	argD												
hypothetical protein	zinc ABC transporter substrate-binding protein	zinc uptake regulation protein	zinc ABC transporter ATP-binding protein	zinc ABC transporter permease	GTP-binding protein LepA	hypothetical protein	histidinol dehydrogenase (EC:1.1.1.23)	LysR family transcriptional regulator	iow molecular weign phosphotyrosine protein phosphatase (EC:3.1.3.48)	NAD-dependent deacetylase	ormunne cychoreannnase/mu-crystannn rannny protein	hypothetical protein	hypothetical protein	50S ribosomal protein L28	arginine/ornithine transport system ATPase (EC:2.7)	ATP-dependent helicase HrpB	GMC family oxidoreductase	aquaporin Z	NUDIX family hydrolase	phosphate transporter family protein	2-dehydropantoate 2-reductase (EC:1.1.1.169)	acetyltransferase	vacetygamma-guuamy-pnospnate reductase (EC:1.2.1.38)	ornithine carbamoyltransferase (EC:2.1.3.3)	acetylornithine transaminase (EC:2.6.1.11)	hypothetical protein	MerR family transcriptional regulator	hypothetical protein	hypothetical protein	ABC transporter permease	hypothetical protein	amidohydrolase	hypothetical protein	glutathione S-transferase	hypothetical protein	hypothetical protein	uracil-DNA glycosylase
-1.27	-11.90	-5.15	-6.66	-2.81	-1.32	-1.71	-1.96	1.23	1.07	1.08	1.31	3.67	2.58	-1.08	1.11	х	-1.17	1.85	-1.26	-1.66	1.16	-1.67	1.09	1.59	3.18	-2.92	-1.64	Х	2.06	-0.97	-1.13	1.10	1.01	1.51	Х	1.14	-2.01
0.80	0.00	0.00	0.01	0.24	0.60	0.07	0.72	0.89	0.50	0.92	0.01	0.00	0.01	0.92	0.92	×	0.86	0.07	0.02	x	0.37	0.24	0.71	0.50	0.13	0.02	0.01	x	0.42	0.30	0.26	0.72	0.24	0.03	x	0.99	0.23
-1.03	-1.30	-1.33	-1.09	1.07	1.51	1.07	-1.17	1.05	1.58	1.63	1.68	3.64	1.37	1.10	2.17	-1.01	-1.12	-1.21	-1.36	1.42	-1.13	1.32	1.44	1.39	1.25	-1.04	-1.24	1.24	1.10	1.14	1.31	-1.04	1.50	1.05	Х	2.06	-1.16
0.95	0.03	0.19	0.25	0.60	0.39	0.76	0.91	0.65	0.05	0.14	0.04	0.01	0.50	0.83	0.00	0.99	0.52	0.21	0.03	0.09	0.59	0.52	0.08	0.04	0.30	0.81	0.38	0.46	0.85	0.65	0.41	0.25	0.07	0.29	х	0.23	0.37
-1.15	-6.60	-3.24	-3.88	-0.87	0.10	-0.32	-1.57	1.14	1.33	1.36	1.50	3.66	1.98	0.01	1.64	N/A	-1.15	0.32	-1.31	-0.12	0.02	-0.18	1.27	1.49	2.22	-1.98	-1.44	N/A	1.58	0.09	0.09	0.03	1.26	1.28	N/A	1.60	-1.59
0.12	5.30	1.91	2.79	1.94	1.42	1.39	0.40	0.09	0.25	0.28	0.18	0.01	0.61	1.09	0.53	N/A	0.02	1.53	0.05	1.54	1.15	1.50	0.18	0.10	0.97	0.94	0.20	N/A	0.48	1.06	1.22	1.07	0.25	0.23	N/A	0.46	0.43
-2.47	-3.48	-1.20	-1.04	1.10	2.15	-1.26	-2.02	1.67	1.80	-1.19	-1.05	3.29	1.86	5.44	1.69	2.65	0.97	2.23	-2.62	2.49	-2.26	-1.28	1.24	1.28	1.78	-3.33	-1.28	2.25	-1.23	2.00	1.96	-3.74	-2.28	1.69	х	1.41	-2.92
0.08	0.00	0.30	0.38	0.85	0.18	0.32	0.76	0.62	0.06	0.76	0.87	0.00	0.03	0.00	0.13	0.02	0.98	0.03	0.00	0.01	0.01	0.93	0.02	0.79	0.10	0.00	0.27	0.06	0.44	0.16	0.11	0.01	0.02	0.04	х	0.88	0.03
1.15	-1.21	1.16	-1.10	-1.28	-1.40	2.05	1.10	-1.07	-1.12	1.65	1.37	1.34	-1.04	-1.52	1.10	-1.20	1.37	2.52	-2.32	-1.62	1.39	-1.68	-1.45	1.18	1.35	1.24	1.11	1.20	1.28	-1.91	-1.31	2.39	1.15	1.03	Х	1.15	-1.06
0.84	0.18	0.46	0.38	0.39	0.50	0.01	0.96	0.49	0.18	0.02	0.05	0.07	x	0.07	0.62	0.53	0.03	0.01	0.01	0.05	0.07	0.42	0.19	0.21	0.14	0.46	0.48	0.59	0.69	0.12	0.06	0.00	0.19	0.87	х	0.73	0.70
-0.66	-2.35	-0.02	-1.07	-0.09	0.38	0.40	-0.46	0.30	0.34	0.23	0.16	2.32	0.41	1.96	1.40	0.73	1.17	2.38	-2.47	0.44	-0.44	-1.48	-0.11	1.23	1.57	-1.05	-0.09	1.73	0.03	0.05	0.33	-0.68	-0.57	1.36	N/A	1.28	-1.99
1.81	1.14	1.18	0.03	1.19	1.78	1.66	1.56	1.37	1.46	1.42	1.21	0.98	1.45	3.48	0.29	1.93	0.20	0.15	0.15	2.06	1.83	0.20	1.35	0.05	0.22	2.29	1.20	0.52	1.26	1.96	1.64	3.07	1.72	0.33	N/A	0.13	0.93
1.38	-1.29	-1.16	-1.03	1.31	-1.15	1.37	-1.05	1.09	1.05	1.41	1.53	1.78	-1.41	-1.49	-1.07	-1.32	-1.01	1.48	1.04	-1.36	-1.30	-1.03	-1.50	-1.27	-1.51	1.25	1.43	-1.10	1.04	1.29	1.20	1.01	-1.15	1.59	x	-1.01	1.04
0.51	0.04	0.15	0.78	0.38	0.71	0.31	0.97	0.30	0.56	0.03	0.13	0.00	0.14	0.10	0.19	0.08	0.89	0.05	0.96	0.14	0.08	0.97	0.05	0.01	0.02	0.38	0.09	0.69	0.97	0.30	0.56	1.00	0.51	0.05	х	0.95	0.85
1.50	-1.00	1.05	-1.16	1.27	-1.96	1.07	2.51	-1.91	-2.05	1.45	1.35	5.25	1.36	-1.60	1.05	-1.69	1.04	1.71	1.05	-1.41	-1.09	-0.95	-2.27	-1.83	-1.52	1.33	-1.22	-1.46	2.13	-1.42	-1.27	1.36	-1.46	1.35	Х	-1.16	2.22
0.61	0.42	0.93	0.24	0.48	0.32	0.74	0.10	0.05	0.01	0.02	0.08	0.01	x	0.13	0.75	0.17	0.50	0.18	0.19	0.48	0.93	0.88	0.03	0.01	0.14	0.49	0.46	0.15	0.19	0.40	0.14	0.01	0.03	0.17	×	0.77	0.13
1.44	-1.15	-0.05	-1.10	1.29	-1.56	1.22	0.73	-0.41	-0.50	1.43	1.44	3.52	-0.02	-1.55	-0.01	-1.51	0.02	1.60	1.05	-1.39	-1.20	-0.99	-1.89	-1.55	-1.52	1.29	0.11	-1.28	1.59	-0.06	-0.04	1.19	-1.31	1.47	N/A	-1.09	1.63
0.06	0.15	1.11	0.06	0.02	0.41	0.15	1.78	1.50	1.55	0.02	0.09	1.74	1.39	0.06	1.06	0.19	1.03	0.12	0.01	0.02	0.11	0.04	0.39	0.28	0.01	0.04	1.33	0.18	0.55	1.36	1.24	0.18	0.16	0.12	N/A	0.08	0.59

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SPO0980 SPO0981 SPO0982 SPO0983 SPO0984 SPO0985 SPO0986 SPO0986 SPO0987

SPO0977 SPO0978 SPO0979 SPO0973 SPO0974 SPO0975 SPO0976 SPO0968 SPO0970 SPO0971 SPO0972

SPO0964 SPO0965 SPO0966 SPO0967 SPO0959 SPO0960 SPO0961 SPO0962 SPO0963

SPO0957 SPO0958

SPO0956

SPO0950 SPO0951 SPO0952 SPO0953 SPO0954 SPO0955

SPO1023	SPO1022	SPO1021	SPO1020	SPO1019	SPO1018	SPO1017	SPO1016	SPO1015	SPO1014	SPO1013	SPO1012	SPO1011	SPO1010	SPO1009	SPO1008	SPO1007	SPO1006	SPO1005	SPO1004	SPO1003	SPO1002	SPO1001	SPO1000	SPO0999	SPO0998	SPO0997	SPO0996	SPO0995	SPO0994	SPO0993	SPO0992	SPO0991	SPO0990	SPO0989
												accB	accC	aat					clpX	clpP		soxF	soxE	soxD	soxC	sox B	soxA	soxZ	soxY	sox X	soxW	soxV	soxS	
response regulator	transcriptional regulator	branched-chain amino acid ABC transporter substrate-binding protein	branched-chain amino acid ABC transporter permease	permease	binding protein branched-chain amino acid ABC transporter	branched-chain amino acid ABC transporter ATP- binding protein branched-chain amino acid ABC transporter ATP-	hypothetical protein	3-hydroxybutyrate dehydrogenase	AMP-binding protein	acetyl-CoA acetyltransferase (EC:2.3.1.9)	LuxR family transcriptional regulator	acetyl-CoA carboxylase, biofin carboxyl carrier protein (EC:6.4.1.2)	acetyI-CoA carboxylase biotin carboxylase subunit (EC:6.4.1.2)	(EC:2.3.2.6)	hypothetical protein	hypothetical protein	NADH dehydrogenase (EC:1.6.99.3)	endoribonuclease L-PSP	A 17-dependent protease A 17-binding subunit ClpX	ATP-dependent Clp protease proteolytic subunit (EC:3.4.21.92)	AraC family transcriptional regulator	sulfur oxidation F protein	diheme cytochrome c SoxE	diheme cytochrome c SoxD	sulfur oxidation molybdopterin C protein	sulfur oxidation B protein	diheme cytochrome c SoxA	sulfur oxidation Z protein	sulfur oxidation protein SoxY	monoheme cytochrome c SoxX	thioredoxin	sulfur oxidation V protein	regulatory protein SoxS	transcriptional regulator SoxR
-1.11	5.16	4.59	7.83	8.42	9.26	6.01	-1.41	-1.73	0.94	1.01	-1.77	-1.36	-1.14	-1.03	1.69	1.50	2.13	2.04	1.01	1.43	-1.05	-2.10	-2.81	-1.37	-1.31	-1.43	-2.55	-2.29	-1.92	-3.25	1.05	-1.87	-2.14	-1.73
0.71	0.02	0.12	0.02	0.00	0.00	0.00	0.06	0.02	0.01	0.03	0.50	0.06	0.01	0.98	0.44	0.01	0.01	0.01	0.26	0.51	0.38	0.11	0.00	0.52	0.08	0.12	0.00	0.01	0.25	0.07	0.39	0.23	0.03	0.01
-1.28	-1.07	-2.82	-1.87	-2.77	-1.50	-1.01	1.14	1.63	2.42	1.55	-1.38	-1.37	-1.07	-1.08	2.34	1.67	1.38	1.14	1.09	0.98	1.39	-2.12	-1.62	-1.02	-1.33	-1.49	-1.95	-2.17	-1.45	-2.33	-2.59	-1.36	-1.43	-1.07
0.53	0.66	0.09	0.05	х	0.59	0.86	0.11	0.12	0.00	0.17	0.30	0.18	0.17	0.62	0.12	0.23	0.22	0.49	0.70	0.63	0.22	0.04	0.04	0.96	0.09	0.19	0.02	0.03	0.37	0.01	0.01	0.33	0.04	0.53
-1.20	2.05	0.89	2.98	2.83	3.88	2.50	-0.14	-0.05	1.68	1.28	-1.58	-1.37	-1.11	-1.06	2.02	1.59	1.76	1.59	1.05	1.20	0.17	-2.11	-2.22	-1.20	-1.32	-1.46	-2.25	-2.23	-1.69	-2.79	-0.77	-1.62	-1.79	-1.40
0.09	3.12	3.71	4.85	5.60	5.38	3.51	1.28	1.68	0.74	0.27	0.20	0.01	0.03	0.03	0.33	0.09	0.38	0.45	0.04	0.23	1.22	0.01	0.60	0.18	0.01	0.03	0.30	0.06	0.23	0.46	1.82	0.25	0.35	0.33
-1.70	2.96	-8.57	-2.25	-1.74	-1.77	-1.47	-1.49	-1.46	-1.83	0.94	-1.48	1.51	2.38	2.94	2.09	1.50	1.93	2.16	1.39	2.26	-1.20	-5.90	-2.72	-1.66	-2.89	-3.98	-6.96	-8.62	-4.86	-14.50	-5.72	-3.45	-2.58	1.66
0.32	0.06	0.03	0.17	0.12	0.31	0.00	0.18	0.01	0.02	0.07	0.65	0.04	0.08	0.04	0.30	0.05	0.01	0.03	0.69	0.04	0.58	0.04	0.00	0.12	0.02	0.02	0.00	0.00	0.09	0.00	0.00	0.02	0.06	0.03
2.17	5.91	57.80	42.40	47.90	21.80	26.40	1.62	5.75	8.44	5.36	1.53	-1.08	-1.21	-1.08	1.05	1.17	0.97	-1.16	1.13	1.21	1.24	1.92	1.41	1.28	1.49	1.39	1.38	1.56	1.96	1.73	1.27	1.01	1.67	-1.45
0.15	0.00	0.00	0.00	0.01	0.01	0.00	0.07	0.00	0.00	0.00	0.11	0.34	0.05	0.10	0.99	0.45	0.47	0.07	0.84	0.28	0.23	0.04	0.10	0.22	0.07	0.15	0.07	0.11	0.05	0.04	0.22	0.78	0.02	0.06
0.24	4.44	24.62	20.08	23.08	10.02	12.47	0.07	2.15	3.31	3.15	0.03	0.22	0.59	0.93	1.57	1.34	1.45	0.50	1.26	1.74	0.02	-1.99	-0.66	-0.19	-0.70	-1.30	-2.79	-3.53	-1.45	-6.39	-2.23	-1.22	-0.46	0.11
1.94	1.48	33.19	22.33	24.82	11.79	13.94	1.56	3.61	5.14	2.21	1.51	1.30	1.80	2.01	0.52	0.17	0.48	1.66	0.13	0.53	1.22	3.91	2.07	1.47	2.19	2.69	4.17	5.09	3.41	8.12	3.50	2.23	2.13	1.56
1.62	1.40	2.49	2.42	2.27	1.98	2.06	1.05	1.97	3.44	2.48	1.15	-1.26	-1.23	-1.29	2.04	1.38	1.17	1.39	1.43	1.21	-1.06	1.59	-1.03	1.24	-0.99	-1.54	1.04	-1.50	1.08	-1.20	-1.59	0.99	-0.98	-1.19
0.04	0.17	0.01	0.01	0.16	0.06	0.01	0.84	0.01	0.01	0.02	0.69	0.17	0.04	0.14	0.07	0.03	0.54	0.07	0.30	0.30	0.91	0.01	_	0.29	0.73	0.21	0.71	0.08	0.85	0.03	0.03			
1.65	-1.53	3.57	2.58	x	2.66	1.54	-1.31	4.78	6.04	5.87	1.07	0.92	-1.76	-1.86	1.22	0.97	0.98	-1.20	1.08	1.42	-1.27	1.19	-1.43	-0.97	1.14	-0.93	1.07	0.95	1.45	1.16	-1.29			-1.86
0.42	3 0.06	0.00	0.04	x	0.10	0.12	0.44	0.00	0.00	0.02	0.86	0.26	5 0.03	5 0.03	0.36	0.42	0.52	0.11	0.88	0.13	0.61	0.10	0.10	0.31	0.11	0.34	0.54	0.37	0.31	0.55	0.10	0.02		5 0.01
1.64	-0.07	) 3.03	1 2.50	N/A	) 2.32	1.80	4 -0.13	3.38	4.74	4.18	5 1.11	-0.17	-1.50	-1.58	1.63	2 1.18	1.07	0.10	3 1.26	3 1.32	-1.17	1.39		0.13	0.08	-1.24		-0.28		-0.02	-1.44			-1.53
	7 1.47	0.54	0.08	N/A	0.34	0.26	3 1.18	1.41	1.30	1.70	0.04	1.09	0.27	3 0.28	0.41	0.20	0.10	1.30	0.18	0.11	0.11	0.20	3 0.20	1.11	1.06	4 0.30	0.02	3 1.22	0.19	1.18	4 0.15			3 0.34

SPO1024 SPO1025 SPO1026 SPO1027 SPO1027 SPO1028 SPO1029 SPO1030

## Chapter 10: Appendix

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<ul> <li>bypothetical protein</li> <li>bypothetical protein</li> <li>phage integrase site specific recombinase</li> <li>bypothetical protein</li> <li>c-5 cytosine-specific family DNA methylase</li> <li>phage integrase site specific recombinase</li> <li>bypothetical protein</li> </ul>	hypothetical protein sensor histidine kinase (EC:2.7.3) hypothetical protein hypothetical protein hypothetical protein metallo-beta-lactamase hypothetical protein
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SPO 1031 SPO 1032 SPO 1033 SPO 1033 SPO 1034 SPO 1036 SPO 1037 SPO 1040 SPO 1041 SPO 1042 SPO 1042 SPO 1043 SPO 1043 SPO 1045 SPO 1045 SPO 1055 SPO 1052 SPO 1055 SPO 1055

-1.11	-1.15	1.02	-0.99	-1.73	1.41	-1.49	1.09	Х	x	1.19	х	-1.98	х	-1.66	-1.30	1.08	0.96	0.91	-1.68	Х	-2.33	-2.22	-2.71	-2.86	х	1.38	-1.29	1.04	1.58	-2.13	-1.51	-1.30	-1.18	0.92	-1.16	-1.02	×	-1.55
0.33	0.18	0.86	0.98	0.03	0.02	0.01	0.65	x	х	0.14	×	0.79	x	0.01	0.55	0.86	0.36	0.84	0.29	х	0.01	0.13	0.00	Х	×	0.19	0.81	0.68	0.57	0.02	0.15	0.56	0.31	0.93	0.16	0.87	Х	0.01
-1.66	-1.75	-1.55	-1.28	-1.19	-1.06	-1.20	-1.50	x	х	1.12	x	-1.17	-1.07	-1.30	-1.05	-1.21	-1.18	-1.16	-1.18	-1.25	-1.79	-1.81	-1.43	-1.29	-1.29	1.18	-1.31	1.17	1.32	1.45	1.14	-1.98	-1.54	-1.35	1.12	-1.11	1.07	-1.30
0.05	0.02	0.07	0.05	0.43	0.37	0.50	0.44	x	x	0.41	×	0.83	0.81	0.05	0.96	0.26	0.73	0.23	0.67	0.60	0.13	0.43	0.36	0.65	0.57	0.15	0.61	0.47	0.46	0.05	0.58	0.01	0.03	0.62	0.40	0.52	0.29	0.19
-1.39	-1.45	-0.27	-1.14	-1.46	0.18	-1.35	-0.21	N/A	N/A	1.16	N/A	-1.58	N/A	-1.48	-1.18	-0.06	-0.11	-0.12	-1.43	N/A	-2.06	-2.02	-2.07	-2.08	N/A	1.28	-1.30	1.11	1.45	-0.34	-0.19	-1.64	-1.36	-0.21	-0.02	-1.07	N/A	-1.43
0.28	0.30	1.29	0.14	0.27	1.24	0.15	1.30	N/A	N/A	0.03	N/A	0.41	N/A	0.18	0.13	1.15	1.07	1.04	0.25	N/A	0.27	0.20	0.64	0.78	N/A	0.10	0.01	0.06	0.13	1.79	1.33	0.34	0.18	1.14	1.14	0.05	N/A	0.13
1.21	1.32	1.07	1.12	-1.91	3.58	-1.68	1.20	-1.02	-1.10	2.10	х	-2.48	х	-1.85	-1.33	-1.55	-2.40	-1.81	-2.23	-1.81	-1.99	-2.27	-3.73	-1.55	-1.17	-1.73	-2.26	1.13	4.06	1.19	2.73	-1.87	-1.37	0.91	0.92	-1.77	-1.35	-2.54
0.38	0.35	0.96	0.29	0.04	0.00	0.18	0.34	0.09	0.70	0.06	×	0.74	x	0.03	0.04	0.27	0.11	0.34	0.18	0.26	0.07	0.17	0.01	0.83	0.70	0.11	0.54	0.61	0.10	0.35	0.11	0.04	0.30	0.93	0.28	0.41	0.97	0.01
-9.18	-11.30	-11.10	-13.80	-7.34	-1.47	1.09	1.09	Х	х	-1.00	х	-1.04	-1.24	1.07	1.65	1.61	1.43	1.18	1.07	Х	1.48	1.04	-1.32	-1.49	-1.09	1.27	1.15	1.01	-1.32	-1.58	1.05	-1.15	-1.40	1.00	-1.28	1.35	х	2.47
0.00	0.00	0.00	0.00	0.02	0.03	0.61	0.83	x	х	0.97	×	0.98	х	0.86	0.04	0.14	0.45	0.77	0.86	x	0.01	0.96	0.69	х	x	0.26	0.64	0.80	0.22	0.00	0.87	0.22	0.02	0.89	0.32	0.58	х	0.01
-3.99	-4.99	-5.02	-6.34	-4.63	1.06	-0.30	1.15	N/A	N/A	0.55	N/A	-1.76	N/A	-0.39	0.16	0.03	-0.49	-0.32	-0.58	N/A	-0.26	-0.62	-2.53	-1.52	-1.13	-0.23	-0.56	1.07	1.37	-0.20	1.89	-1.51	-1.39	0.96	-0.18	-0.21	N/A	-0.03
5.20	6.31	6.09	7.46	2.72	2.53	1.39	0.05	N/A	N/A	1.55	N/A	0.72	N/A	1.46	1.49	1.58	1.92	1.50	1.65	N/A	1.74	1.66	1.21	0.03	0.04	1.50	1.71	0.06	2.69	1.39	0.84	0.36	0.01	0.04	1.10	1.56	N/A	2.51
-1.77	-1.60	-2.43	-2.30	-1.63	-1.11	-1.03	-1.39	-1.03	-1.35	-1.60	1.31	1.32	-1.11	-1.29	1.04	1.07	-1.07	1.32	-1.05	-1.02	-1.47	-1.45	-1.34	-1.04	-1.04	1.22	-1.13	1.17	1.05	-1.02	1.35	-1.17	1.15	1.70	1.00	-1.03	1.29	1.73
0.10	0.16	0.02	0.00	0.01	0.22	0.76	0.11	0.84	0.18	0.05	×	0.38	0.60	0.14	0.95	0.69	0.71	0.15	0.94	0.89	0.01	0.64	0.26	1.00	0.94	0.25	0.67	0.18	0.93	0.69	0.14	0.15	0.06	0.31	0.74	0.98	х	0.02
-2.59	-1.85	-3.17	-3.63	-2.12	-1.63	1.27	2.05	Х	Х	-1.94	х	1.27	Х	-1.08	1.21	1.09	0.96	1.10	-1.01	Х	-1.40	1.57	1.26	1.06	Х	-1.08	1.03	0.97	-1.50	-1.40	1.33	1.49	1.24	2.38	0.95	1.61	х	1.54
0.02	0.09	0.02	0.02	0.05	0.30	0.53	0.32	х	x	0.05	×	0.63	x	0.41	0.44	0.52	0.77	0.86	0.86	х	0.26	0.72	0.74	х	×	0.64	0.97	0.65	0.16	0.15	0.13	0.26	0.05	0.14	0.72	0.33	х	0.10
-2.18	-1.73	-2.80	-2.97	-1.88	-1.37	0.12	0.33	N/A	N/A	-1.77	N/A	1.30	N/A	-1.19	1.13	1.08	-0.06	1.21	-1.03	N/A	-1.44	0.06	-0.04	0.01	N/A	0.07	-0.05	1.07	-0.23	-1.21	1.34	0.16	1.20	2.04	0.98	0.29	N/A	1.64
0.41	0.13	0.37	0.67	0.25	0.26	1.15	1.72	N/A	N/A	0.17	N/A	0.03	N/A	0.11	0.09	0.01	1.02	0.11	0.02	N/A	0.04	1.51	1.30	1.05	N/A	1.15	1.08	0.10	1.28	0.19	0.01	1.33	0.05	0.34	0.02	1.32	N/A	0.10

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## Chapter 10: Appendix

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propionyl-CoA carboxylase subunit alpna (EC:6.4.1.3)	lipoprotein	hypothetical protein	hypothetical protein	hypothetical protein	lipoprotein	hypothetical protein	propionyl-CoA carboxylase subunit beta (EC:6.4.1.3)	Bcr/CfIA subfamily drug resistance transporter	LysR family transcriptional regulator	hypothetical protein	DNA-binding protein	hypothetical protein	choline dehydrogenase (EC:1.1.99.1)	hypothetical protein	TetR family transcriptional regulator	hypothetical protein	hypothetical protein	choline sulfatase (EC:3.1.6.6)	transcriptional regulator BetI	hypothetical protein	hypothetical protein	L-lysine exporter	chromosome replication initiation inhibitor protein	nitroreductase	TrkA domain-containing protein	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	glutamine amidotransferase (EC:6.3.5.2)	FAD-binding dehydrogenase	hypothetical protein	serine/threonine protein phosphatase				
6.30	-1.18	4.50	-2.80	3.41	0.97	6.27	10.00	-1.33	-1.65	1.90	1.23	-1.32	1.16	-1.42	-1.23	1.54	-1.26	1.20	-1.23	1.38	1.15	Х	-1.06	-0.97	-1.46	-1.09	-1.14	-3.40	1.13	x	-1.16	-1.20	-1.13	1.27	x	-1.39	-1.16
0.07	0.03	0.01	0.50	0.61	0.07	0.00	0.00	0.31	0.83	0.32	0.84	0.01	х	0.32	0.17	0.77	0.09	0.55	0.01	0.04	1.00	x	0.87	0.88	0.09	x	0.78	0.00	0.96	×	0.24	0.73	0.60	0.80	×	0.03	0.81
1.90	-1.85	1.51	-1.48	1.67	1.06	4.49	4.44	-1.43	-1.20	-1.18	-1.50	1.26	-1.34	-1.40	-1.07	1.26	1.18	-1.16	-1.19	1.13	-1.18	х	-1.16	-1.14	1.06	1.06	-1.09	-1.53	1.06	1.13	-1.73	-1.38	1.04	1.23	1.62	-1.21	-1.87
0.07	0.04	0.10	0.64	0.20	0.78	0.00	0.00	0.02	0.86	0.74	0.33	0.54	0.56	0.37	0.67	0.44	0.57	0.54	0.10	0.16	0.77	х	0.69	0.77	0.84	0.93	0.83	0.13	0.89	0.56	0.05	0.21	0.97	0.65	0.01	0.08	0.09
4.10	-1.52	3.01	-2.14	2.54	1.02	5.38	7.22	-1.38	-1.43	0.36	-0.14	-0.03	-0.09	-1.41	-1.15	1.40	-0.04	0.02	-1.21	1.26	-0.02	N/A	-1.11	-1.05	-0.20	-0.02	-1.12	-2.47	1.10	N/A	-1.45	-1.29	-0.04	1.25	N/A	-1.30	-1.52
2.20	0.33	1.50	0.66	0.87	0.04	0.89	2.78	0.05	0.23	1.54	1.37	1.29	1.25	0.01	0.08	0.14	1.22	1.18	0.02	0.13	1.17	N/A	0.05	0.09	1.26	1.08	0.02	0.94	0.03	N/A	0.29	0.09	1.09	0.02	N/A	0.09	0.36
5.43	2.86	5.31	-2.65	3.09	1.15	6.51	4.44	1.10	-1.79	-1.55	-1.43	5.28	1.44	-1.91	1.74	0.92	-1.46	1.34	1.12	2.46	1.17	Х	-1.20	1.47	1.18	-2.01	1.05	-15.10	1.98	2.90	-3.16	-1.28	-1.05	1.74	3.48	-0.98	1.26
0.07	0.01	0.01	0.61	0.45	0.05	0.01	0.00	0.23	0.76	0.78	0.88	0.00	0.43	0.43	0.01	0.26	0.33	0.13	0.19	0.08	0.98	х	0.95	0.31	0.63	x	0.95	0.00	0.36	0.00	0.01	0.66	0.92	0.30	0.01	0.17	0.59
8.86	0.95	11.80	-1.20	6.60	1.43	24.10	25.00	-1.50	-1.44	x	-1.33	-1.22	-1.14	-1.16	1.15	-1.07	1.18	-1.33	-1.38	-1.07	-1.25	х	-1.36	1.13	1.03	1.03	-1.24	-1.33	-1.18	-1.50	1.79	-4.94	-8.24	-1.06	-2.07	-4.20	-10.40
0.01	0.33	0.00	0.78	0.01	0.01	0.00	0.00	0.04	0.68	×	0.34	0.58	0.80	0.67	0.15	0.40	0.78	0.12	0.07	0.55	0.64	Х	0.41	0.74	0.96	x	0.54	0.17	0.10	0.03	0.07	0.02	0.03	0.58	0.03	0.00	0.02
7.15	1.90	8.56	-1.93	4.85	1.29	15.31	14.72	-0.20	-1.62	N/A	-1.38	2.03	0.15	-1.54	1.45	-0.07	-0.14	0.01	-0.13	0.70	-0.04	N/A	-1.28	1.30	1.11	-0.49	-0.10	-8.22	0.40	0.70	-0.69	-3.11	-4.65	0.34	0.71	-2.59	-4.57
1.72	0.96	3.25	0.73	1.76	0.14	8.80	10.28	1.30	0.17	N/A	0.05	3.25	1.29	0.38	0.30	1.00	1.32	1.34	1.25	1.77	1.21	N/A	0.08	0.17	0.08	1.52	1.15	6.89	1.58	2.20	2.48	1.83	3.60	1.40	2.78	1.61	5.83
1.16	1.54	1.06	-1.01	1.26	1.18	1.74	1.07	1.38	1.17	1.12	-1.04	1.43	-1.34	-1.54	1.03	1.23	-1.02	-1.13	-1.16	1.18	-1.12	-1.48	-1.36	-1.18	1.29	-1.09	1.39	1.40	-1.30	-1.14	1.24	-2.05	-2.39	-1.23	-1.23	-1.25	-2.63
0.29	0.06	0.18	0.99	0.51	0.12	0.01	0.43	0.01	0.88	0.12	0.72	0.15	0.40	0.28	0.85	0.07	0.93	0.31	0.28	0.28	0.83	×	0.24	0.54	0.43	0.90	0.32	0.19	0.02	0.44	0.20	0.05	0.06	0.18	0.12	0.06	0.02
1.11	1.32	-1.31	1.59	1.17	-2.00	1.74	1.95	1.41	1.21	x	-1.05	1.18	1.47	-1.07	-1.14	-1.01	x	-1.50	-1.72	-1.20	1.54	Х	-1.29	-1.24	1.19	1.35	1.44	1.66	-1.91	-1.89	2.20	-1.75	-4.30	-1.13	-2.73	-1.64	-3.95
0.46	0.35	0.38	0.41	0.91	0.00	0.01	0.02	0.04	0.84	×	0.98	0.65	0.39	0.96	0.03	0.75	×	0.20	0.01	0.79	0.59	х	0.57	0.66	0.53	x	0.13	0.02	0.00	0.10	0.03	0.15	0.08	0.50	0.01	0.04	0.03
1.14	1.43	-0.13	0.29	1.22	-0.41	1.74	1.51	1.40	1.19	N/A	-1.05	1.31	0.06	-1.31	-0.05	0.11	N/A	-1.32	-1.44	-0.01	0.21	N/A	-1.33	-1.21	1.24	0.13	1.42	1.53	-1.61	-1.52	1.72	-1.90	-3.35	-1.18	-1.98	-1.45	-3.29
0.02	0.11	1.19	1.30	0.05	1.59	0.00	0.44	0.02	0.02	N/A	0.01	0.13	1.41	0.23	1.09	1.12	N/A	0.19	0.28	1.19	1.33	N/A	0.04	0.03	0.05	1.22	0.03	0.13	0.31	0.38	0.48	0.15	0.96	0.05	0.75	0.20	0.66

SPO 1064 SPO 1065 SPO 1066 SPO 1067 SPO 1067 SPO 1070 SPO 1071 SPO 1072 SPO 1073 SPO 1073 SPO 1075 SPO 1075 SPO 1075 SPO 1077 SPO 1077 SPO 1077 SPO 1081 SPO 1082 SPO 1083 SPO 1084 SPO 1085 SPO 1085 SPO 1086 SPO 1086 SPO 1087 SPO 1087 SPO 1088 SPO 1087 SPO 1090 SPO 1090 SPO 1091 SPO

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SPO1102 SPO1103 SPO1104 SPO1106 SPO1106 SPO1107 SPO1107 SPO1110 SPO1110 SPO1111 SPO1112 SPO1114 SPO1115 SPO1116 SPO1116 SPO1117 SPO1120 SPO1121

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doeB	doeX	doeC	doeD												lolD	lolE		proS																			
Na-acetyl-L-2,4-diaminobutyric acid deacetylase	transcriptional regulator	aspariate-semiatuenyde denydrogenase (EC:1.2.1.16)	diaminobutyric acid transaminase	hypothetical protein	NnrU family protein	glycine betaine/proline ABC transporter permease	glycine betaine/proline ABC transporter A1P- binding protein (EC:3.6.3.32)	giyone betame/prome ABC transporter substrate- binding protein	choice between the ABC transporter substrate	hypothetical protein	hypothetical protein	cytochrome c family protein	cytochrome c family protein	hypothetical protein	lipoprotein releasing system ATP-binding protein	lipoprotein releasing system transmembrane protein LoIE	hypothetical protein	prolyl-tRNA synthetase (EC:6.1.1.15)	hypothetical protein	hypothetical protein	hypothetical protein	Ppx/GppA phosphatase	hypothetical protein	hypothetical protein	TRAP dicarboxylate transporter subunit DctM	TRAP transporter, transmembrane protein	TRAP transporter, periplasmic protein	hypothetical protein	LysR family transcriptional regulator	hypothetical protein	DnaJ-like protein Dj1A	phosphinothric in N-acetyltransferase	hypothetical protein	methylmalonyl-CoA mutase (EC:5.4.99.2)	hypothetical protein	hypothetical protein	lipoprotein
-1.47	2.68	-1.38	-1.16	-1.19	-1.08	-1.52	-1.71	-1.22	-1.76	1.40	1.67	-1.39	1.53	-1.55	-1.27	-1.55	1.61	1.76	1.07	0.99	1.61	2.86	-1.57	-1.45	-2.64	1.09	-1.03	х	1.35	1.15	1.45	-1.44	-1.18	5.91	5.51	0.96	3.83
0.01	x	0.35	0.08	0.58	0.85	0.02	0.00	0.06	0.35	0.87	0.02	0.78	0.05	х	0.03	0.01	0.00	0.54	0.98	0.33	0.03	0.05	0.51	0.06	0.07	0.17	0.88	x	0.69	0.09	0.03	0.22	0.75	0.00	0.01	0.47	0.36
1.30	1.30	1.18	1.35	1.16	1.01	-1.31	-1.72	-1.81	-1.10	1.08	1.04	1.35	1.12	1.07	-1.07	-1.03	1.44	1.13	-1.08	1.06	1.07	1.61	-1.97	1.37	1.44	1.63	1.67	1.69	-1.05	1.99	1.34	-1.39	1.14	2.65	1.78	-1.94	1.89
0.04	0.25	0.59	0.42	0.24	0.91	0.19	0.06	0.00	0.72	0.90	0.72	0.34	0.33	0.74	0.46	0.90	0.03	0.48	0.79	0.16	0.57	0.03	0.34	0.09	0.47	0.09	0.01	0.00	0.95	0.02	0.27	0.20	0.32	0.00	0.03	0.06	0.05
-0.09	1.99	-0.10	0.10	-0.02	-0.04	-1.42	-1.72	-1.52	-1.43	1.24	1.36	-0.02	1.33	-0.24	-1.17	-1.29	1.53	1.45	-0.01	1.03	1.34	2.24	-1.77	-0.04	-0.60	1.36	0.32	N/A	0.15	1.57	1.40	-1.42	-0.02	4.28	3.65	-0.49	2.86
1.39	0.69	1.28	1.26	1.18	1.05	0.11	0.01	0.29	0.33	0.16	0.32	1.37	0.20	1.31	0.10	0.26	0.09	0.32	1.08	0.03	0.27	0.63	0.20	1.41	2.04	0.27	1.35	N/A	1.20	0.42	0.05	0.03	1.16	1.63	1.87	1.45	0.97
-2.71	1.62	-2.12	-1.19	1.21	1.53	1.08	-1.44	-1.69	-1.21	1.13	1.17	-2.85	2.33	1.72	3.41	2.97	-1.31	3.36	1.36	1.00	2.39	1.55	-1.81	-1.80	-1.26	-1.12	1.39	-0.92	1.39	2.14	2.07	-2.16	-1.04	4.04	3.03	-1.23	3.95
0.00	0.05	0.31	0.67	0.12	0.27	0.03	0.07	0.00	0.84	0.95	0.21	0.32	0.01	0.04	0.00	0.01	0.07	0.24	0.77	0.96	0.01	0.11	0.44	0.13	0.98	0.59	0.00	0.16	0.57	0.02	0.04	0.18	0.55	0.00	0.02	0.02	0.31
-1.01	1.49	-0.99	-1.05	-1.30	1.31	1.10	-1.06	1.30	1.25	1.40	1.58	1.36	-1.00	-1.59	-1.60	-1.51	1.10	-1.05	-0.99	-1.02	-1.55	1.66	1.45	1.60	1.34	1.67	2.04	1.30	1.29	1.29	1.23	1.23	4.50	15.20	13.60	1.16	7.62
0.91	0.12	0.78	0.76	0.25	0.26	0.32	0.63	0.05	0.60	0.45	0.12	0.53	0.76	0.23	0.05	0.01	0.77	0.20	0.87	0.57	0.06	0.02	0.58	0.05	0.51	0.07	0.04	0.40	0.35	0.17	0.26	0.49	0.09	0.00	0.00	0.10	0.01
-1.86	1.56	-1.55	-1.12	-0.05	1.42	1.09	-1.25	-0.20	0.02	1.27	1.38	-0.75	0.67	0.06	0.91	0.73	-0.11	1.16	0.18	-0.01	0.42	1.61	-0.18	-0.10	0.04	0.28	1.72	0.19	1.34	1.72	1.65	-0.47	1.73	9.62	8.32	-0.04	5.79
0.85	0.07	0.57	0.07	1.26	0.11	0.01	0.19	1.50	1.23	0.14	0.21	2.11	1.66	1.66	2.51	2.24	1.21	2.21	1.18	1.01	1.97	0.05	1.63	1.70	1.30	1.40	0.33	1.11	0.05	0.43	0.42	1.70	2.77	5.58	5.29	1.20	1.84
1.27	1.15	1.07	-1.07	1.37	1.16	1.83	1.23	1.79	-1.15	1.15	-1.04	2.02	-1.03	1.33	1.27	-1.06	-1.53	-1.14	-1.13	-1.11	-1.39	-1.12	1.66	-0.98	1.49	1.50	1.20	1.25	1.07	1.33	1.08	1.20	-0.97	1.39	-2.03	1.14	1.23
0.33	0.37	0.63	0.49	0.09	0.31	0.00	0.08	0.07	0.55	0.87	0.72	0.17	0.97	0.29	0.02	0.39	0.00	0.25	0.18	0.06	0.21	0.34	0.40	0.66	0.36	0.02	0.15	0.19	0.35	0.11	0.67	0.63	0.71	0.06	0.01	0.13	0.19
1.06	1.11	1.35	-0.94	-1.08	1.10	-0.98	1.31	2.04	-0.99	1.16	1.09	1.65	-1.48	-1.30	-1.48	-1.69	-1.24	-1.29	-1.35	-1.37	-1.83	0.98	1.67	1.32	1.24	1.30	1.70	-1.24	-1.02	-1.33	1.19	1.50	1.20	1.46	-1.34	1.52	1.13
0.82	×	Х	0.25	0.81	0.46	0.60	0.10	0.08	0.96	0.62	0.04	0.14	0.23	0.56	0.01	0.01	0.17	0.13	0.18	0.02	0.01	0.43	0.41	х	0.72	0.05	0.04	х	0.83	0.58	0.19	0.41	0.06	0.04	0.06	0.02	0.51
1.17	1.13	1.21	-1.01	0.15	1.13	0.43	1.27	1.92	-1.07	1.16	0.03	1.84	-1.26	0.02	-0.11	-1.38	-1.39	-1.22	-1.24	-1.24	-1.61	-0.07	1.67	0.17	1.37	1.40	1.45	0.01	0.03	0.00	1.14	1.35	0.12	1.43	-1.69	1.33	1.18
0.11	0.02	0.14	0.06	1.23	0.03	1.40	0.04	0.13	0.08	0.01	1.07	0.18	0.23	1.32	1.38	0.32	0.14	0.08	0.11	0.13	0.22	1.05	0.01	1.15	0.13	0.10	0.25	1.25	1.05	1.33	0.05	0.15	1.08	0.04	0.34	0.19	0.05

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SPO1137 SPO1138 SPO1139 SPO1132 SPO1133 SPO1134 SPO1135 SPO1136 SPO1131

SP01123 SP01124 SP01125 SP01126 SP01127 SP01127 SP01128 SP01128 SP01130

### Chapter 10: Appendix

SPO1181	SPO1180	SPO1179	SPO1178	SPO1177	SPO1176	SPO1175	SPO1174	SPO1173	SPO1172	SPO1171	SPO1168	SPO1166	SPO1165	SPO1164	SPO1163	SPO1162	SPO1161	SPO1160	SPO1159	SPO1158	SPO1157	SPO1156	SPO1155	SPO1154	SPO1153	SPO1152	SP01151	SPO1150	SPO1148	SPO1147	SPO1146	SPO1145	SPO1144	SPO1143	SPO1142	SPO1141	SPO1140
		mraW								pyc					hisB	hisH				hisA		hisF	hisE						gntR	uehA	uehB	uehC					doeA
penicillin-binding protein	hypothetical protein	MraW (EC:2.1.1)	cell division protein MraZ	hypothetical protein	Ser/Thr protein phosphatase	hypothetical protein	DNA helicase II	diguanylate cyclase	rivity-dependent alpha-nyurox y acid dehydrogenase	carboxylase (EC:6.4.1.1)	hypothetical protein	aminotransferase	hypothetical protein	hypothetical protein	dehydratase (EC:4.2.1.19)	уусстогриохриате зуншахе хирини гизга (EC:2.4.2)	AraC family transcriptional regulator	hypothetical protein	hypothetical protein	imidazole-4-carboxamide isomerase (EC:5.3.1.16)	hypothetical protein	glycerol phosphate synthase subunit HisF	pyrophosphatase (EC:3.6.1.31)	lipoprotein	CoA-binding domain-containing protein	win-arginine transiocation pattiway signal sequence domain-containing protein	hypothetical protein	RNA methyltransferase	family transcriptional regulator	TRAP transporter large integral membrane protein	TRAP transporter small integral membrane protein	TRAP transporter periplasmic binding protein	universal stress protein family protein	ectoine utilization protein EutA	threonine dehydratase (EC:4.3.1.19)	ectoine utilization protein EutC (EC:4.3.1.12)	ectoine hydrolase
-1.13	-1.16	-1.26	0.99	10.60	2.48	x	-1.51	-2.30	1.12	-1.16	-1.22	1.64	-1.87	1.76	1.07	1.26	-1.39	1.44	1.29	2.14	-1.24	1.27	1.86	1.24	-1.04	1.04	-0.98	-1.32	-1.60	1.12	1.99	x	-1.33	1.08	1.50	-1.52	2.37
0.14	0.08	0.37	0.86	0.04	0.01	×	0.23	х	0.35	0.30	0.03	0.01	0.14	0.37	0.89	0.20	0.06	0.07	0.41	0.35	0.01	0.30	0.09	0.30	0.97	0.99	0.67	0.41	0.16	0.92	0.03	×	0.07	0.80	0.04	0.28	0.25
-1.19	-1.34	-1.40	-1.85	4.53	1.64	1.34	-1.19	-1.73	-1.18	1.22	1.30	1.65	-1.12	1.24	1.11	1.28	-1.12	1.41	-2.04	2.22	2.43	1.15	1.63	-1.11	-1.04	1.60	1.03	1.13	-1.30	-1.22	-1.32	-1.38	-1.21	-1.00	1.10	-1.25	1.36
0.17	0.11	0.23	0.09	0.03	0.07	0.06	0.27	x	0.30	0.49	0.05	0.12	0.67	0.07	0.70	0.03	0.44	0.02	0.04	0.17	0.10	0.53	0.10	0.36	0.73	0.04	0.81	0.62	0.42	0.87	0.59	0.76	0.28	0.88	0.03	0.09	0.44
-1.16	-1.25	-1.33	-0.43	7.57	2.06	N/A	-1.35	-2.02	-0.03	0.03	0.04	1.65	-1.50	1.50	1.09	1.27	-1.26	1.43	-0.38	2.18	0.60	1.21	1.75	0.06	-1.04	1.32	0.03	-0.10	-1.45	-0.05	0.34	N/A	-1.27	0.04	1.30	-1.39	1.87
0.03	0.09	0.07	1.42	3.04	0.42	N/A	0.16	0.29	1.15	1.19	1.26	0.01	0.38	0.26	0.02	0.01	0.14	0.02	1.67	0.04	1.84	0.06	0.12	1.18	0.00	0.28	1.00	1.23	0.15	1.17	1.66	N/A	0.06	1.04	0.20	0.14	0.50
0.88	0.96	1.33	1.56	35.90	-1.21	1.95	1.03	-3.48	4.73	0.79	-1.54	1.56	-6.31	1.57	2.05	3.70	-1.18	2.04	-2.57	3.87	-2.07	3.36	4.10	1.94	1.17	1.54	2.51	2.00	-2.08	-2.11	-1.35	-1.38	-2.61	-1.55	-1.26	-3.03	-1.15
0.05	0.24	0.37	0.72	0.01	0.35	0.04	0.92	х	0.00	0.02	0.13	0.14	0.02	0.09	0.19	0.02	0.08	0.02	0.03	0.31	0.00	0.01	0.00	0.03	0.73	0.08	0.02	0.11	0.27	0.50	0.05	0.96	0.04	0.21	0.09	0.01	0.91
-1.11	-1.05	1.06	1.05	3.13	1.07	1.33	-1.18	-1.13	1.22	1.23	1.41	-1.62	-1.09	1.09	-1.28	-1.42	1.34	1.83	1.08	1.05	2.07	-1.04	-1.22	0.99	1.02	1.17	-1.16	1.11	1.16	-1.04	-1.18	-1.25	-0.99	-1.24	-1.15	-1.12	1.26
0.21	0.43	0.94	0.91	0.03	0.65	0.27	0.12	x	0.26	0.30	0.07	0.04	0.74	0.43	0.26	0.03	0.03	0.05	0.83	0.98		0.39	0.11	0.45	0.96	0.30	0.31	0.74	0.76	0.96	×	0.85	0.81	0.25	0.12		0.62
-0.12	-0.05	1.20	1.31	19.52	-0.07	1.64	-0.08	-2.31		1.01		-0.03		1.33	0.39	1.14	0.08	1.94	-0.75	2.46	0.00	1.16	1.44	1.47	1.10	1.36					-1.2		-1.80	-1.40			0.06
0.99				2 16.39		0.31		1.18	1.76	0.22	1.48	3 1.59	2.61	0.24	1.67	2.56	1.26	0.11	1.83	1.41	2.07	2.20	2.66	0.47	0.08	0.19	1.84		5 1.62	3 0.54	0.09	0.06	0.81	0.16	0.06		1.21
9 1.12			1.14		4 -1.47	1 1.12	1 1.07	3 1.02	6 0.97	2 1.25	3 1.28	9 1.15	-1.21	4 -1.22	-1.80	-1.14	5 1.45	1 1.27	3 2.07	-1.03		-1.24	5 1.09			-1.07						-1.27					-1.20
2 0.59			4 0.78		~	2 0.29	7 0.57	2 X	7 0.28		8 0.11	5 0.54	1 0.15	2 0.06	0 0.00	4 0.21	5 0.16	7 0.28	7 0.01	3 0.90		4 0.05	9 0.49	6 0.48	0 0.97	7 0.39		6 0.39	6 0.28	0 0.65	6 0.95	.7 0.68			_	3 0.22	
9 1.07				8 1.12	4 -1.59	8 8	7 1.46	1.70	8 -1.13		1 1.27	4 1.05	5 1.57				6 1.31	8 -1.05	1 3.09	0 -1.22	2 1.58	5 1.29				9 -1.76		9 -1.10						2 -1.39			7 -0.99
7 0.91			9 0.42	2 0.95	0.07	x	6 0.03	0 X	0.08		7 0.13	5 0.92	7 0.11	0.29	0 0.06	0.00	1 0.11	0.58	9 0.00	0.50		9 0.55	5 0.10	0.14	0.84	<sup>76</sup> 0.02			0.80	1.00	27 X	X	0.39	X 68			0.95
91 1.10				95 0.05	.1.53	K N/A	03 1.27	K 1.36	0.08 -0.08	)3 1.26		92 1.10			06 -1.90	00 -1.54	11 1.38	0.11	2.58	50 -1.13		55 0.03	-0.28	14 -1.09		02 -1.42	46 1.29							-1.40			95 -1.10
	35 0.20		0.08	05 1.07		'A N/A		36 0.34	08 1.05	26 0.01		0 0.05		0.03	90 0.10	54 0.40	38 0.07	11 1.16	58 0.51	0.10	57 0.02		28 1.37	09 0.03	03 0.03	42 0.35			30 0.16	23 0.17	17 0.11	13 1.14	52 0.18				10 0.10
03	20	1.20	80	07	06	'A	19	34	05	01	01	05	1.39	03	10	40	07	16	51	10	02	1.27	37	03	03	35	15	03	16	17	11	14	18	01	07	1.64	10

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												recN		lpxC	ftsZ	ftsA		ddl	murB				murC	murG				gst				murD			mraY	murF	murE
phage integrase site specific recombinase	phage integrase site specific recombinase	DNA-binding protein	hypothetical protein	carboxypeptidase (EC:3.4.17.11)	hippurate hydrolase (EC:3.5.1.32)	oligopeptide ABC transporter ATP-binding protein	oligopeptide ABC transporter permease	oligopeptide ABC transporter permease	ongopeptude ABC transporter substrate-binding protein	MIrC domain-containing protein	voltage-gated chloride channel family protein	repair protein RecN	competence lipoprotein ComL	N-acetylglucosamine deacetylase (EC:3.5.1)	division protein FtsZ	division protein FtsA	cell division protein FtsQ	ligase (EC:6.3.2.4)	reductase (EC:1.1.1.158)	hypothetical protein	hypothetical protein	amino acid permease	ligase (EC:6.3.2.8)	beta-N-acetylglucosaminyltransferase (EC:2.4.1.227)	cell division protein FtsW	4-oxalocrotonate tautomerase	hypothetical protein	S-transferase (EC:2.5.1.18)	transcriptional regulator family protein	hypothetical protein	hypothetical protein	synthetase (EC:6.3.2.9)	hypothetical protein	hypothetical protein	(EC:2.7.8.13)	ligase (EC:6.3.2.10)	ligase (EC:6.3.2.13)
-1.11	-1.16	-2.09	-1.66	-1.20	-1.31	-1.35	-1.52	-1.98	-2.15	1.27	1.23	1.20	1.18	3.42	-1.38	-1.54	-2.05	-2.92	-1.31	1.18	-1.30	1.34	-1.09	-1.26	0.96	1.14	1.26	1.01	-2.61	2.83	-1.05	-1.52	-1.16	1.21	-1.49	1.62	1.12
0.66	0.21	0.29	x	0.83	0.01	0.16	0.57	0.01	0.00	0.54	0.62	0.57	0.63	0.17	0.03	0.04	0.00	0.00	0.22	0.59	0.42	0.93	0.51	0.00	0.31	0.62	0.39	0.30	0.04	0.02	0.91	0.69	0.25	0.41	0.06	0.00	0.40
1.01	-1.12	1.33	-1.35	1.07	0.99	1.09	1.06	-1.10	-1.35	-1.03	1.02	1.00	1.02	1.28	-1.28	-1.36	-1.73	-1.58	-1.02	1.06	1.36	1.33	-1.28	-1.46	-1.40	-1.30	-2.49	-2.71	-1.19	1.34	-1.15	1.29	1.16	1.15	-1.12	-1.06	-1.14
0.81	0.37	0.12	0.47	0.82	0.92	0.83	0.89	0.81	0.03	0.88	0.93	0.95	0.99	0.47	0.38	0.05	0.01	0.02	0.89	0.88	0.39	0.80	0.39	0.02	0.05	0.29	0.05	0.00	0.28	0.20	0.25	0.27	0.16	0.09	0.65	0.64	0.40
-0.05	-1.14	-0.38	-1.51	-0.06	-0.16	-0.13	-0.23	-1.54	-1.75	0.12	1.13	1.10	1.10	2.35	-1.33	-1.45	-1.89	-2.25	-1.17	1.12	0.03	1.34	-1.19	-1.36	-0.22	-0.08	-0.62	-0.85	-1.90	2.09	-1.10	-0.12	0.00	1.18	-1.31	0.28	-0.01
1.06	0.02	1.71	0.16	1.14	1.15	1.22	1.29	0.44	0.40	1.15	0.11	0.10	0.08	1.07	0.05	0.09	0.16	0.67	0.14	0.06	1.33	0.01	0.10	0.10	1.18	1.22	1.88	1.86	0.71	0.75	0.05	1.41	1.16	0.03	0.18	1.34	1.13
-4.30	-1.79	-1.21	-1.53	-1.89	-1.62	-1.31	-1.82	-2.15	-3.20	-1.55	-2.05	-1.68	1.72	0.92	0.93	0.86	0.88	-1.67	1.07	-1.43	1.23	1.62	1.22	0.82	1.04	3.06	2.74	2.34	-1.35	-4.16	2.07	-1.55	-1.45	-1.44	1.46	1.20	1.53
0.04	0.02	0.81	0.12	0.36	0.00	0.06	0.41	0.01	0.00	0.04	0.23	0.00	0.05	0.23	0.01	0.10	0.07	0.00	0.24	0.05	0.70	0.77	0.77	0.01	0.63	0.01	0.00	0.00	0.21	0.02	0.03	0.26	0.08	0.14	0.01	0.39	0.02
-1.52	-1.08	1.16	1.29	1.54	1.89	1.36	1.23	1.22	1.29	1.48	-1.14	-1.14	-1.15	-2.57	1.09	-1.83	-1.37	-1.13	1.09	1.12	-1.20	-1.12	-1.27	-1.26	-1.17	1.12	1.21	-1.01	1.56	-1.41	-1.34	-1.18	1.51	1.52	-1.31	1.01	-1.58
0.06	0.80	0.88	0.32	0.10	0.04	0.10	0.64	0.31	0.05	0.18	0.62	0.23	0.14	0.00	0.98	0.01	0.04	0.16	0.45	0.70	0.58	0.87	0.44	0.28	0.09	0.17	0.16	0.79	0.10	0.10	0.08	0.44	0.17	0.02	0.39	0.90	0.08
-2.91	-1.44	-0.03	-0.12	-0.18	0.14	0.03	-0.30	-0.47	-0.96	-0.04	-1.60	-1.41	0.29	-0.83	1.01	-0.49	-0.25	-1.40	1.08	-0.16	0.02	0.25	-0.03	-0.22	-0.06	2.09	1.98	0.67	0.11	-2.79	0.37	-1.37	0.03	0.04	0.08	1.11	-0.03
1.39	0.36	1.19	1.41	1.72	1.76	1.34	1.53	1.69	2.25	1.52	0.46	0.27	1.44	1.74	0.08	1.34	1.13	0.27	0.01	1.28	1.22	1.37	1.25	1.04	1.11	0.97	0.77	1.68	1.46	1.38	1.71	0.19	1.48	1.48	1.39	0.10	1.56
-1.24	-1.01	1.39	-1.18	1.45	1.22	-1.04	1.38	-1.19	2.00	1.55	1.06	-1.01	1.18	1.33	1.93	1.13	1.35	1.13	-1.69	1.14	1.52	1.36	1.03	-1.12	1.22	1.58	1.37	1.64	1.03	1.05	-1.70	1.13	-1.16	-1.08	-1.03	1.13	-1.12
0.23	0.83	0.15	0.42	0.15	0.46	0.74	0.40	0.02	0.00	0.15	0.49	0.71	0.10	0.33	0.01	0.95	0.01	0.25	0.18	0.62	0.29	0.75	0.99	0.41	0.03	0.03	0.05	0.01	0.70	0.77	0.04	0.65	0.35	0.13	0.87	0.54	0.05
-1.75	Х	-0.96	X	1.45	1.59	1.19	1.39	-1.06	2.60	1.47	1.44	1.61	1.23	-1.59	1.84	-1.12	-1.11	-1.22	-1.78	-1.01	-1.18	1.50	-1.57	-1.29	1.05	1.88	1.89	2.14	1.16	1.57	-2.33	-1.10	-1.13	0.99	1.72	1.51	-2.24
0.05	×	0.66	x	0.16	0.22	0.36	0.46	0.09	0.00	0.13	0.04	0.03	0.29	0.01	0.18	0.47	0.02		0.12	0.94	0.73	0.75	0.32	0.49	0.32	0.05	0.02		0.51	0.03	0.02	0.90	0.25	0.54		0.25	0.01
-1.50	N/A	0.22	N/A	1.45	1.41	0.08	1.39	-1.13	2.30	1.51	1.25	0.30	1.21	-0.13		0.00		-0.05		0.06		1.43			1.14	1.73	1.63		1.10	1.31		0.01			0.35		-1.68
0.26	N/A	1.18	N/A	0.00	0.19	1.12	0.01	0.06	0.30	0.04	0.19	1.31	0.03	1.46	0.04	1.13	1.23	1.18	0.05	1.08	1.35	0.07	1.30	0.09	0.09	0.15	0.26	0.25	0.06	0.26	0.31	1.12	0.02	1.03	1.38		0.56
0.26	N/A	1.18	N/A	0.00	0.19	1.12	0.01	0.06	0.30	0.04	0.19	1.31	0.03	1.46	0.04	1.13	1.23	1.18	0.05	1.08	1.35	0.07	1.30	0.09	0.09	0.15	0.26	0.25	0.06	0.26	0.31	1.12	0.02	1.03	1.38	U.19	0.10

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SPU1260	SPO1259	SPO1258	SPO1257	SPO1256 ppk2	SPO1255	SPO1254	SPO1253 parE	SPO1252	SPO1251	SPO1250	SPO1249	SPO1247	SPO1246	SPO1245	SPO1244 efp	SPO1243	SPO1242 cobQ	SPO1241	SPO1240	SPO1239 pcm-1	SPO1237	SPO1236	SPO1235	SPO1234	SPO1233	SPO1232	SPO1231	SPO1230	SPO1229	SPO1228	SPO1227	SPO1226	SPO1225	SPO1224	SPO1223	SPO1222	SPO1221
MATE ettiux family protein	hypothetical protein	alpha/beta hydrolase	glutathione S-transferase	2 kinase (EC:2.7.4.1)	alkylhydroperoxidase	hypothetical protein	<i>E</i> topoisomerase IV subunit B (EC:5.99.1)	hypothetical protein	hypothetical protein	lipoprotein	paraquat-inducible protein	glycosyl transferase family protein	glycine cleavage system protein T	ABC transporter permease	factor P	hypothetical protein	Q acid synthase	hypothetical protein	TolC family type I secretion outer membrane protein	O-methyltransferase (EC:2.1.1.77)	hypothetical protein	H-NS family DNA-binding protein	hypothetical protein	lipoprotein	hypothetical protein	hypothetical protein	hypothetical protein	resolvase site-specific recombinase	hypothetical protein								

1.18	1.15	0.95	1.21	1.17	-1.08	0.97	-1.04	-1.10	1.48	-1.18	1.10	-1.12	1.19	1.62	1.48	1.33	2.34	1.10	1.62	1.71	1.07	-1.23	1.19	1.33	1.63	-1.39	1.23	1.21	1.42	-0.95	-1.34	1.08	x	×	x	-2.07	1.34	-1.01
0.26	0.97	0.73	0.68	0.95	0.35	0.78	0.42	0.11	0.35	0.59	0.38	0.91	0.46	0.21	0.18	0.03	0.03	0.59	0.18	0.45	0.49	0.49	0.82	0.67	0.60	0.11	0.89	0.95	0.11	0.56	0.36	0.90	×	×	x	0.15	0.83	0.91
1.46	-1.29	-1.20	1.20	1.14	-1.12	-1.22	1.03	1.00	-1.04	1.25	1.34	1.14	-1.11	-1.11	1.67	0.98	1.64	1.20	-1.39	1.17	-1.16	-1.36	-1.27	1.01	1.38	-1.36	-1.12	-1.03	1.47	-1.13	1.00	1.01	-1.05	х	1.07	-1.29	1.18	1.43
0.20	0.26	0.28	0.66	0.59	0.27	0.38	0.84	0.99	0.92	0.21	0.26	0.67	0.12	0.40	0.38	0.70	0.09	0.43	0.05	0.47	0.19	0.56	0.23	1.00	0.73	0.22	0.91	0.75	0.04	0.31	0.99	0.97	x	x	0.52	0.75	0.52	0.26
1.32	-0.07	-0.13	1.21	1.16	-1.10	-0.13	-0.01	-0.05	0.22	0.04	1.22	0.01	0.04	0.26	1.58	1.15	1.99	1.15	0.12	1.44	-0.04	-1.30	-0.04	1.17	1.51	-1.38	0.05	0.09	1.45	-1.04	-0.17	1.05	N/A	N/A	N/A	-1.68	1.26	0.21
0.14	1.22	1.07	0.01	0.02	0.02	1.10	1.04	1.05	1.26	1.22	0.12	1.13	1.15	1.37	0.10	0.18	0.35	0.05	1.51	0.27	1.12	0.07	1.23	0.16	0.13	0.01	1.18	1.12	0.03	0.09	1.17	0.04	N/A	N/A	N/A	0.39	0.08	1.22
2.14	-2.33	-3.51	-1.74	3.15	1.57	-3.10	1.01	-1.89	1.67	-1.07	1.58	1.24	1.49	1.52	4.63	3.24	2.84	-1.76	1.58	3.23	1.38	1.23	-1.20	0.99	-1.78	-1.77	-1.35	-1.53	-1.04	1.16	0.99	-4.17	х	-2.42	х	-2.76	-1.42	-1.57
0.11	0.01	0.02	0.10	0.37	0.05	0.18	0.98	0.00	0.67	0.94	0.04	0.86	0.04	0.07	0.03	0.02	0.00	0.06	0.10	0.06	0.02	0.75	0.51	0.90	0.43	0.03	0.92	0.68	0.63	0.12	0.94	0.11	x	x	Х	0.21	0.74	0.51
-1.57	1.31	2.07	1.64	1.28	1.88	1.07	1.03	1.01	1.20	-1.16	1.03	1.10	1.27	1.31	-1.28	1.70	-1.18	1.51	1.10	1.09	1.21	1.27	1.60	1.99	2.43	1.94	1.11	1.88	1.81	1.04	1.56	2.17	×	×	x	-1.18	1.51	1.28
0.16	0.39	0.02	0.07	0.54	0.01	0.78	0.96	0.96	0.60	0.60	0.96	0.60	0.19	0.07	0.08	0.02	0.22	0.12	0.56	0.86	0.35	0.68	0.12	0.09	0.35	0.02	0.63	0.03	0.00	0.82	0.09	0.10	×	x	Х	0.81	0.17	0.49
0.29	-0.51	-0.72	-0.05	2.22	1.73	-1.02	1.02	-0.44	1.44	-1.12	1.31	1.17	1.38	1.42	1.68	2.47	0.83	-0.13	1.34	2.16	1.30	1.25	0.20	1.49	0.33	0.09	-0.12	0.18	0.39	1.10	1.28	-1.00	N/A	N/A	N/A	-1.97	0.05	-0.15
1.86	1.82	2.79	1.69	0.94	0.16	2.09	0.01	1.45	0.23	0.04	0.28	0.07	0.11	0.11	2.96	0.77	2.01	1.64	0.24	1.07	0.09	0.02	1.40	0.50	2.11	1.86	1.23	1.71	1.43	0.06	0.28	3.17	N/A	N/A	N/A	0.79	1.47	1.43
1.21	1.05	-1.62	1.48	-1.17	-1.25	-1.01	1.10	-1.21	1.25	1.49	1.21	1.40	-1.04	-1.05	1.07	-1.22	1.11	1.22	-1.15	1.16	-1.40	-1.05	-1.34	-1.14	1.14	-1.08	-1.13	1.09	1.30	-1.04	1.14	-1.15	х	х	Х	-1.19	1.03	-1.14
0.53	0.90	0.01	0.15	0.56	0.19	0.89	0.23	0.14	0.58	0.06	0.30	0.27	0.26	0.68	0.76	0.03	0.17	0.52	0.10	0.30	0.02	0.91	0.12	0.42	0.85	0.41	0.89	0.59	0.02	0.89	0.39	0.53	×	×	х	0.81	0.95	0.67
-1.21	1.77	1.03	1.67	-1.17	-1.10	1.63	-1.27	-1.17	1.17	1.17	-1.17	-0.98	-1.05	-1.12	-1.73	1.04	-1.24	1.27	1.10	1.00	-1.29	-1.24	-1.41	-1.16	1.60	1.55	-1.31	1.53	-1.06	-0.97	1.15	1.13	х	х	Х	-0.96	-1.03	-1.05
0.69	0.03	0.55	0.07	0.71	0.18	0.07	0.24	0.17	0.59	0.44	0.43	0.85	0.55	0.54	0.02	0.61	0.66	0.64	0.47	0.15	0.07	0.73	0.16	0.47	0.42	0.12	Х	0.14	0.64	0.48	0.29	0.59	x	x	Х	0.85	0.95	0.91
0.00	1.41	-0.30	1.58	-1.17	-1.18	0.31	-0.09	-1.19	1.21	1.33	0.02	0.21	-1.05	-1.09	-0.33	-0.09	-0.06	1.25	-0.02	1.08	-1.35	-1.15	-1.38	-1.15	1.37	0.24	-1.22	1.31	0.12	-1.00	1.15	-0.01	N/A	N/A	N/A	-1.07	0.00	-1.10
1.21	0.36	1.33	0.10	0.00	0.08	1.32	1.19	0.02	0.04	0.16	1.19	1.19	0.01	0.04	1.40	1.13	1.18	0.03	1.13	0.08	0.05	0.10	0.03	0.01	0.23	1.32	0.09	0.22	1.18	0.04	0.01	1.14	N/A	N/A	N/A	0.12	1.03	0.04

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SPO1301	SPO1300	SPO1299	SPO1298	SPO1295	SPO1294	SPO1293 phaP	SPO1292 phaC	SPO1291 phaZ	SPO1290	SPO1289	SPO1288	SPO1287	SPO1286	SPO1285	SPO1284	SPO1283	SPO1282 thrS	SPO1281	SPO1280	SPO1279	SPO1278	SPO1277	SPO1276	SPO1275	SPO1274	SPO1273 thyX	SPO1272	SPO1271	SPO1270	SPO1269	SPO1268	SPO1267	SPO1266	SPO1265	SPO1264 aspC-2	SPO1263	
glutamine amidotransferase	glutamine synthetase	LysR family transcriptional regulator	oxidoreductase, aldo/keto reductase	aminotransferase	polyhydroxyalkanoate synthesis repressor PhaR	PhaP	polymerase	depolymerase	hypothetical protein	alpha/beta hydrolase	hypothetical protein	glyoxalase	hypothetical protein	hypothetical protein	ArsR family transcriptional regulator	hypothetical protein	synthetase (EC:6.1.1.3)	hypothetical protein	ArsC family protein	cold shock family protein	hypothetical protein	thymidylate synthase (EC:2.1.1.148)	hypothetical protein	hypothetical protein	lactoylglutathione lyase	hypothetical protein	lipoprotein	MarR family transcriptional regulator	hypothetical protein	DNA-binding protein	aminotransferase (EC:2.6.1.1)	hypothetical protein					

-1.39	1.00	-1.26	1.74	3.77	-1.52	4.33	6.67	1.71	-1.09	1.07	1.04	1.22	-1.05	-1.28	-1.65	-1.45	х	1.74	-1.17	1.42	1.69	1.18	1.59	1.36	1.26	1.93	-1.56	-1.75	1.25	-1.21	-1.37	-1.25	2.67	2.74	1.25	-1.37	-1.57	1.45
0.02	0.30	0.26	0.05	0.18	0.01	0.17	0.00	0.15	0.83	0.92	0.77	0.17	0.54	0.43	0.01	0.08	x	0.37	0.39	0.26	0.02	0.38	0.03	0.42	0.69	0.35	0.21	0.40	0.48	0.31	0.05	0.11	0.01	0.38	0.08	0.06	0.28	0.10
1.10	1.60	1.03	1.19	-1.03	-1.20	1.19	1.19	-1.31	1.15	1.30	1.09	1.28	1.26	1.01	-1.45	1.00	1.26	1.39	1.30	1.12	1.14	1.15	-1.11	-1.30	-1.05	1.15	-1.45	-1.38	1.04	-1.02	-1.00	2.35	1.51	1.94	1.20	1.43	-1.16	-1.02
0.46	0.05	0.70	0.18	0.80	0.31	0.65	0.55	0.35	0.83	0.50	0.39	0.06	0.15	0.99	0.10	0.97	0.26	0.17	0.06	0.28	0.72	0.13	0.58	0.30	0.94	0.70	0.09	0.08	0.94	0.93	0.88	0.03	0.12	0.31	0.53	0.07	0.37	0.99
-0.15	1.30	-0.12	1.47	1.37	-1.36	2.76	3.93	0.20	0.03	1.19	1.07	1.25	0.11	-0.14	-1.55	-0.23	N/A	1.57	0.07	1.27	1.42	1.17	0.24	0.03	0.11	1.54	-1.51	-1.57	1.15	-1.12	-1.18	0.55	2.09	2.34	1.23	0.03	-1.37	0.22
1.25	0.30	1.15	0.28	2.40	0.16	1.57	2.74	1.51	1.12	0.12	0.03	0.03	1.16	1.15	0.10	1.23	N/A	0.18	1.24	0.15	0.27	0.02	1.35	1.33	1.16	0.39	0.06	0.19	0.11	0.10	0.19	1.80	0.58	0.40	0.03	1.40	0.21	1.24
-2.34	0.94	-1.16	-1.41	1.72	-2.14	12.30	1.65	1.73	-1.28	1.28	-1.15	-1.11	-1.15	-1.47	-3.53	-2.36	2.70	1.25	3.13	-1.65	0.94	1.34	1.98	1.66	-4.47	1.74	-2.86	-1.63	1.23	1.16	-1.29	-4.45	3.58	4.73	1.93	0.95	-1.64	2.02
0.01	0.68	0.90	0.66	0.56	0.01	0.01	0.01	0.03	0.65	0.80	0.73	0.52	0.56	0.31	0.00	0.01	0.01	0.85	0.01	0.09	0.29	0.13	0.03	0.12	0.00	0.03	0.04	0.19	0.72	0.15	0.16	0.01	0.01	0.24	0.01	0.27	0.30	0.01
1.33	-1.26	1.08	1.61	1.46	1.34	-3.79	1.86	1.72	2.07	1.44	1.66	1.81	1.62	1.25	-1.47	-1.27	-2.25	1.89	1.06	1.09	-1.06	-1.12	-1.04	1.50	1.44	1.17	-1.23	1.13	2.01	1.21	1.04	1.14	-1.04	1.48	1.13	-1.02	-1.20	-1.07
0.18	0.03	0.54	0.09	0.41	0.14	0.00	0.02	0.13	0.45	0.43	0.01	0.01	0.02	0.66	0.03	0.06	0.01	0.07	0.79	0.24	0.70	0.23	0.07	0.07	0.24	0.41	0.09	0.12	0.01	0.06	0.86	0.07	0.76	0.42	0.24	0.49	0.45	0.23
-0.51	-0.16	-0.04	0.10	1.59	-0.40	4.26	1.76	1.73	0.40	1.36	0.26	0.35	0.24	-0.11	-2.50	-1.82	0.23	1.57	2.10	-0.28	-0.06	0.11	0.47	1.58	-1.52	1.46	-2.05	-0.25	1.62	1.19	-0.13	-1.66	1.27	3.11	1.53	-0.04	-1.42	0.48
1.84	1.10	1.12	1.51	0.13	1.74	8.05	0.11	0.01	1.68	0.08	1.41	1.46	1.39	1.36	1.03	0.55	2.48	0.32	1.04	1.37	1.00	1.23	1.51	0.08	2.96	0.28	0.82	1.38	0.39	0.03	1.17	2.80	2.31	1.63	0.40	0.98	0.22	1.55
-1.85	-1.12	-1.38	1.06	1.10	-1.22	1.70	-1.29	-1.19	1.60	1.66	-1.25	1.24	1.65	1.05	-1.54	-1.09	-1.07	1.58	-1.24	-2.42	-1.27	1.53	-1.08	1.85	-1.01	-1.08	-1.54	-1.53	-1.03	1.03	-1.05	-1.19	-1.38	1.26	-1.26	1.15	-1.03	1.07
0.05	0.29	0.21	0.29	0.78	0.05	0.13	0.17	0.06	0.58	0.18	0.04	0.23	0.07	0.81	0.08	0.30	0.77	0.07	0.13	0.00	0.20	0.03	0.02	0.01	0.98	0.54	0.02	0.04	0.20	0.91	0.59	0.21	0.03	0.62	0.02	0.39	0.89	0.32
-1.81	-1.55	-1.65	-1.18	1.45	1.15	2.63	1.28	1.34	1.77	1.57	-1.63	-1.02	1.41	1.52	-1.22	-1.08	-1.48	1.42	-1.36	-2.41	-1.04	1.14	-1.64	2.42	1.10	-1.59	-1.44	-2.23	1.27	1.06	-0.93	-1.07	-1.73	1.12	-1.39	-1.38	1.61	-1.23
0.01	0.01	0.09	0.87	0.37	0.53	0.00	0.15	0.16	0.56	0.12	0.01	0.73	0.02	0.62	0.07	0.35	0.14	0.07	0.37	0.01	0.96	0.14	0.01	0.06	0.72	0.03	0.04	0.02	0.37	0.62	0.37	0.88	0.01	0.96	0.14	0.17	0.02	0.66
-1.83	-1.34	-1.52	-0.06	1.28	-0.04	2.17	-0.01	0.08	1.69	1.62	-1.44	0.11	1.53	1.29	-1.38	-1.09	-1.28	1.50	-1.30	-2.42	-1.16	1.34	-1.36	2.14	0.05	-1.34	-1.49	-1.88	0.12	1.05	-0.99	-1.13	-1.56	1.19	-1.33	-0.12	0.29	-0.08
0.02	0.22	0.14	1.12	0.18	1.19	0.47	1.29	1.27	0.09	0.04	0.19	1.13	0.12	0.23	0.16	0.01	0.21	0.08	0.06	0.00	0.12	0.20	0.28	0.29	1.06	0.26	0.05	0.35	1.15	0.02	0.06	0.06	0.18	0.07	0.06	1.27	1.32	1.15

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SPO1340	SPO1339	SPO1337	SPO1336	SPO1335	SPO1334	SPO1333	SPO1332	SPO1331	SPO1330	SPO1329	SPO1328	SPO1327	SPO1326	SPO1325	SPO1324	SPO1323	SPO1322	SPO1321	SPO1320	SPO 13 19	SPO1318	SPO 13 17	SPO1316	SPO1315	SPO1314	SPO1313	SPO1312	SPO1311	SPO1310	SPO1309	SPO1308	SPO1307	SPO1306	SPO1305	SPO1304	SPO1303	SPO1302
	ftsY		ispZ						hflC		gor	rpiA				sdaA					purA			nthB	nthA	secG	pyrG										
hypothetical protein	recognition particle-docking protein FtsY	hypothetical protein	septation protein A	Crp/Fnr family transcriptional regulator	hypothetical protein	serine protease	hypothetical protein	hypothetical protein	protein	HflK protein	reductase (EC:1.8.1.7)	isomerase A (EC:5.3.1.6)	MarR family transcriptional regulator	lipoprotein	glutathione S-transferase	ammonia-lyase (EC:4.3.1.17)	hypothetical protein	thiamine pyrophosphokinase (EC:2.7.6.2)	lipoprotein	hypothetical protein	synthetase (EC:6.3.4.4)	hypothetical protein	hypothetical protein	hydratase subunit beta (EC:4.2.1.84)	hydratase subunit alpha (EC:4.2.1.84)	translocase subunit SecG	synthetase (EC:6.3.4.2)	renal dipeptidase	hypothetical protein	hypothetical protein	hypothetical protein	ATP-binding protein	ubstrate-binding protein His/Glu/Arg/onine family ABC transporter	ABC transporter permease His/Clu/Clu/Arg/onine family ABC transporter	permease	fructosyl-amino acid oxidase His/Clu/Clu/Arg/oning family ABC transporter	glutamine synthetase
1.50	1.36	-1.13	1.28	0.98	-1.29	1.11	-0.98	-1.50	1.06	1.39	-1.18	-1.56	-1.45	-1.39	1.13	2.44	-1.46	-1.39	1.40	-1.28	1.00	-1.14	1.34	1.28	1.35	-1.13	2.25	1.22	0.97	1.79	1.18	-1.31	-2.57	-1.08	-1.26	1.22	-0.99
0.77	0.05	0.81	0.23	0.92	0.54	0.17	0.92	0.01	0.53	0.81	0.11	0.05	0.14	0.84	0.63	0.02	0.30	0.05	0.31	0.02	0.67	х	0.36	0.12	0.86	0.91	0.11	0.09	0.86	0.73	0.70	0.17	0.01	0.54	0.10	0.26	0.45
1.24	1.00	-1.10	-1.20	1.06	-1.71	-1.28	-1.09	1.00	1.36	-1.35	-1.70	-1.09	-1.46	-1.60	-1.24	-1.05	-1.07	1.10	1.53	1.02	1.86	2.10	1.29	1.17	-1.12	-1.09	1.75	-1.07	-1.04	1.30	-1.06	-1.34	-1.56	-1.11	-1.18	-1.02	1.18
0.48	0.81	0.38	0.40	0.89	0.19	0.31	0.82	0.98	0.17	0.02	0.01	0.85	0.04	0.02	0.47	0.43	0.92	0.68	0.11	0.85	0.02	0.04	0.09	0.05	0.78	0.60	0.49	0.23	0.41	0.41	0.76	0.25	0.07	0.50	0.41	0.86	0.12
1.37	1.18	-1.12	0.04	1.02	-1.50	-0.09	-1.04	-0.25	1.21	0.02	-1.44	-1.33	-1.46	-1.50	-0.06	0.70	-1.27	-0.15	1.47	-0.13	1.43	0.48	1.32	1.23	0.12	-1.11	2.00	0.08	-0.03	1.55	0.06	-1.33	-2.07	-1.10	-1.22	0.10	0.10
0.13	0.18	0.01	1.24	0.04	0.21	1.20	0.05	1.25	0.15	1.37	0.26	0.24	0.01	0.11	1.19	1.75	0.19	1.25	0.07	1.15	0.43	1.62	0.03	0.06	1.24	0.02	0.25	1.15	1.01	0.25	1.12	0.02	0.51	0.02	0.04	1.12	1.08
1.25	1.18	-0.94	2.90	1.24	-1.34	1.56	1.47	-1.06	2.29	2.62	2.50	1.68	2.05	-1.32	1.54	2.25	1.54	-1.36	-1.51	-1.44	2.31	3.41	2.56	2.41	1.84	5.26	2.48	1.18	1.00	2.16	1.12	1.36	-2.90	1.22	-0.94	2.29	-1.45
0.85	0.32	0.11	0.04	0.91	0.69	0.09	0.03	0.88	0.00	0.03	0.02	0.01	0.00	0.89	0.21	0.00	0.17	0.31	0.08	0.01	0.03	0.42	0.04	0.01	0.61	0.00	0.03	0.08	0.44	0.64	0.72	0.19	0.00	0.32	0.06	0.01	0.18
1.50	-1.05	-1.07	-1.19	1.25	1.04	1.47	-1.10	-1.01	-1.24	1.30	-1.35	-1.03	-1.53	-1.16	-1.06	-1.07	1.39	1.34	1.36	1.00	-1.26	-1.26	1.39	1.17	1.19	-1.27	-1.71	1.62	1.42	1.71	1.29	1.29	-1.11	1.13	- 1.05	-1.02	-2.11
0.13	0.55	0.60	0.06	0.29	0.94	0.02	0.40	0.84	0.07	0.10	0.02	0.15	0.04	0.54	0.64	0.23	0.68	0.15	0.17	0.59	0.20	0.23	0.21	0.30	0.79	0.11	0.03	0.01	0.07	0.11	0.42	0.05	0.41	0.76	0.39	0.89	0.00
1.38	0.06	-1.01	0.86	1.25	-0.15	1.52	0.19	-1.04	0.53	1.96	0.58	0.33	0.26	-1.24	0.24	0.59	1.47	-0.01	-0.08	-0.22	0.53	1.08	1.98	1.79	1.52	2.00	0.39	1.40	1.21	1.94	1.21	1.33	-2.01	1.18	-1.00	0.64	-1.78
0.13	1.12	0.06	2.05	0.01	1.19	0.05	1.29	0.03	1.77	0.66	1.93	1.36	1.79	0.08	1.30	1.66	0.08	1.35	1.44	1.22	1.79	2.34	0.59	0.62	0.33	3.27	2.10	0.22	0.21	0.23	0.09	0.04	0.90	0.05	0.06	1.66	0.33
-1.03	-1.42	-1.74	1.28	1.63	1.20	2.37	1.29	-1.08	1.63	1.19	-1.06	-1.31	-1.03	-1.01	-2.29	-2.20	-1.26	-1.03	1.14	1.17	1.04	-1.04	1.53	2.07	1.15	0.99	-1.17	1.69	1.06	1.58	-1.29	-1.63	-2.41	-1.24	-1.39	-1.16	-1.95
		4 0.00	0.27	0.19	0.67	0.01	0.12	3 0.54	0.05	0.06	5 0.97	0.02	3 0.79	0.97	0.01	0.00	5 0.66	3 0.81	0.38	0.05	. 0.99	4 0.68	0.08	0.00	0.69	0.71	7 0.02	0.01		0.14		3 0.03	0.32	4 0.39	_	5 0.12	
1.80	-1.12	-1.77	1.10	1.57	1.05	2.76	1.44	-1.07	1.02	1.58	1.29	-1.59	-2.00	-1.16	-1.80	-2.48	1.39	-1.16	1.19	1.15	1.12	-1.66	1.36	1.39	1.88	-1.29	-1.36	1.64	1.29	1.30	-1.25	-1.33	-1.58	-1.07	-1.31	-1.87	
			0.81	0.28	0.79	0.00	0.32	0.94	0.77	0.04	0.12	0.04	) 0.02	5 0.09		3 0.03	0.67	5 0.77	0.62	0.07	0.90	0.16	0.07	0.02	0.47	0.10	0.22	. 0.06	0.11	0.27	0.29	0.06	3 0.06	0.51	0.06	0.07	
		-1.76	1.19	3 1.60	1.13	) 2.57	1.37	-1.08	1.33	1.39	0.12		-1.52	-1.09	-2.05	-2.34	0.06	-1.10	1.17	1.16	1.08	-1.35	1.45	1.73	1.52	-0.15	-1.27	1.67	1.18			-1.48	-2.00	-1.16	-1.35		
		0.02	0.09	0.03	0.08	0.20	0.08	0.01	0.31	0.20	1.18	0.14	0.49	0.08	0.25	0.14	1.33	0.06	0.03	0.01	0.04	0.31	0.09	0.34	0.37	1.14	0.10	0.03	0.12	0.14	0.02	0.15	0.42	0.09			0.20

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SPO1379	SPO1378	SPO1377	SPO1376	SPO1375	SPO1374	SPO1373	SPO1372	SPO1371	SPO1370	SPO1369	SPO1368	SPO1367	SPO1366	SPO1365	SPO1364	SPO1363	SPO1362	SPO1361	SPO1360	SPO1359	SPO1358	SPO1357	SPO1356	SPO1355	SPO1354	SPO1353	SPO1352	SPO1351	SPO1350	SPO1349	SPO1348	SPO1347	SPO1346	SPO1345	SPO1344	SPO1343	SPO1342	SPO1341
		carA											ihvC		glmM	folP				ppdK	glyS	gltS		gly Q		trkH-1		metZ						purD	xseA			
prephenate dehydrogenase	GatB/YqeY domain-containing protein	phosphate synthase small subunit (EC:6.3.5.5)	glycosyl transferase family protein	GntR family transcriptional regulator	pyrimidine 5'-nucleotidase	hypothetical protein	hypothetical protein	2-octaprenyl-6-methoxyphenyl hydroxylase	aminotransferase	GntR family transcriptional regulator	AsnC family transcriptional regulator	AsnC family transcriptional regulator	reductoisomerase (EC:1.1.1.86)	hypothetical protein	mutase (EC:5.4.2.10)	synthase (EC:2.5.1.15)	dihydroneopterin aldolase	multicopper oxidase domain-containing protein	hypothetical protein	phosphate dikinase (EC:2.7.9.1)	synthetase subunit beta (EC:6.1.1.14)	symporter	FkbM family methyltransferase	synthetase subunit alpha (EC:6.1.1.14)	peptidoglycan binding domain-containing protein	system potassium uptake protein TrkH	GTP cyclohydrolase	sulfhydrylase (EC:4.2.99)	hypothetical protein	lipoprotein	(Fe-S)-binding protein	hypothetical protein	LuxR family transcriptional regulator	ligase (EC:6.3.4.13)	VII large subunit (EC:3.1.11.6)	glutathione S-transferase	hypothetical protein	alkane-1 monooxygenase
-2.15	1.20	1.59	-1.33	-1.15	1.34	-2.03	5.85	2.52	5.41	2.08	1.21	-1.52	1.10	-1.07	1.25	-1.79	1.13	1.66	-1.17	1.47	-1.00	1.03	1.10	1.34	-1.16	1.23	1.21	-3.17	Х	-1.27	1.29	-1.05	1.03	2.46	-1.24	1.14	-1.24	-1.00
0.57	0.97	0.01	0.00	0.09	0.23	0.16	0.00	0.08	0.00	0.02	0.98	0.52	0.25	0.88	0.13	0.30	0.43	0.09	0.02	0.71	0.96	0.97	1.00	0.29	0.58	0.66	0.24	0.00	×	0.32	0.44	0.60	0.85	0.01	0.73	0.05	0.13	0.89
-1.18	1.24	1.64	4.05	-1.77	-1.79	1.04	2.52	2.71	1.73	1.09	-1.08	-1.21	1.01	1.52	1.12	-1.21	1.27	-1.27	-1.31	1.19	1.28	1.14	-1.03	1.00	1.22	1.15	-1.28	-1.43	×	1.19	1.15	-1.03	1.07	1.94	-1.11	-1.15	1.01	1.18
0.90	0.24	0.01	0.00	0.10	0.02	0.61	0.01	0.07	0.00	0.69	0.83	0.34	0.93	0.06	0.33	0.52	0.57	0.09	0.14	0.66	0.35	0.57	0.24	0.91	0.44	0.61	0.03	0.03	х	0.74	0.65	0.69	0.83	0.03	0.83	0.13	0.91	0.12
-1.67	1.22	1.62	1.36	-1.46	-0.23	-0.50	4.19	2.62	3.57	1.59	0.06	-1.37	1.06	0.23	1.19	-1.50	1.20	0.20	-1.24	1.33	0.14	1.09	0.04	1.17	0.03	1.19	-0.04	-2.30	N/A	-0.04	1.22	-1.04	1.05	2.20	-1.18	-0.01	-0.12	0.09
0.48	0.02	0.02	2.69	0.31	1.57	1.54	1.67	0.10	1.84	0.50	1.15	0.16	0.05	1.30	0.06	0.29	0.07	1.47	0.07	0.14	1.14	0.05	1.07	0.17	1.19	0.04	1.25	0.87	N/A	1.23	0.07	0.01	0.02	0.26	0.06	1.15	1.13	1.09
-2.15	1.30	7.26	-6.87	0.97	3.02	-2.58	10.70	5.70	2.63	2.98	0.87	-1.55	-2.77	1.67	2.31	-2.28	1.97	2.33	2.69	1.55	1.50	1.62	0.87	5.16	1.35	2.98	1.85	10.20	x	-2.34	3.05	1.25	-1.47	7.44	1.41	1.24	-1.67	1.38
0.59	0.88	0.00	0.03	0.36	0.02	0.05	0.01	0.03	0.01	0.00	0.72	0.72	0.00	0.07	0.00	0.00	0.02	0.04	0.01	0.59	0.07	0.03	0.11	0.00	0.53	0.07	0.08	0.00	x	0.07	0.02	0.48	0.03	0.00	0.73	0.03	0.04	0.01
-1.47	1.15	-1.22	4.16	-1.17	-1.20	1.30	1.41	1.29	1.22	-1.33	-1.50	-2.35	-6.63	1.09	-1.10	-1.19	-1.16	1.75	1.87	1.10	-1.15	-1.08	-1.10	-1.53	1.02	-1.17	-1.17	2.34	x	1.24	-1.45	-1.27	1.35	-1.48	1.30	1.34	1.06	1.45
0.63	0.74	0.09	0.01	0.41	0.24	0.20	0.03	0.43	0.30	0.08	0.09	5 <u>0.05</u>	3 0.01	0.29	0.21	0.23	5 0.23	0.03	0.04	0.75	5 0.03	3 0.34	0.34	0.06	0.76	0.16	0.05	0.01	x	0.73	5 0.05	0.14	0.13	3 0.03	0.68			0.20
-1.81	1.23	3.02	-1.36	-0.10		-0.64	6.06	3 3.50	1.93	0.83	-0.31	-1.95	-4.70	1.38	0.61	-1.74	3 0.41	3 2.04	1 2.28	1.33	3 0.18	0.27	-0.12	1.82	1.19	0.91	0.34	6.27		-0.55	0.80	-0.01	-0.06	3 2.98	3 1.36			1.42
1 0.34	3 0.08	2 4.24	6 5.52	0 1.07	1 2.11	4 1.94	5 4.65	2.21	3 0.71	3 2.16	1 1.19	5 0.40	0 1.93	3 0.29	1 1.71	4 0.55	1 1.57	4 0.29	3 0.41	3 0.23	3 1.33	7 1.35	2 0.98	2 3.35	→ 0.17	1 2.08	4 1.51	7 3.93	N/A	5 1.79	) 2.25	1 1.26	6 1.41	3 4.46	5 0.05			0.04
	1.44	-1.38	62 -0.98	-1.30	-1.47	1.36	1.55		1.27				-1.12						-1.40		-1.29	-1.65	-1.59	-1.29		1.38						-1.21						-1.07
1.03 0.94	44 0.29	38 0.00	98 0.47	30 0.22	47 0.72	36 0.02	55 0.01	1.35 0.59	0.09	1.21 0.06	1.05 0.34	-	-	1.32 0.08	1.66 0.03	1.05 0.67	1.18 0.52	-1.27 0.09		1.09 0.73	29 0.12	65 0.08	59 0.04	29 0.11	1.38 0.14	38 0.03	1.76 0.05	-3.08 0.02	X X	1.42 0.36	1.10 0.70	21 0.33	1.50 0.16	1.43 0.01	1.07 0.94	1.47 0.01		07 0.46
	-															_																						
2.03 0	1.41 0	-2.99 0	Х	1.19 0	-1.24 0	-0.95 0	1.96 0	1.15 0	-1.03 0	-1.82 0	-1.04 0	-1.69 0	-1.60 0	-1.12 0	1.36 0	1.21 0	1.32 0	-1.27 0	1.33 0	1.06 0	-1.22 0	-2.67 0	-1.86 0	-1.72 0	1.51 0	1.28 0	1.42 0	-1.87 0	Х	1.03 0	1.00 0	0	1.37 0		1.29 0			-1.08 0
0.53 0	0.53 1	0.01 -2	X	0.14 -1	0.59 -1	0.16 0	0.01 1	0.58 1	0.64 0	0.04 -(	0.84 0	0.12 -1	0.17 -1	0.73 0	0.02 1	0.71 0	0.13 -(	0.24 -1	0.58 -(	0.76 1	0.24 -1	0.01 -2	0.02 - 1	0.01 -1	0.16 1	0.22 1	0.01 1	0.01 -2	X	0.98 1	0.15 1	0.36 -1	0.02 1	0.08 -(	0.74 1			0.98 -1
0.50	1.43	-2.19	N/A	1.25	-1.36	0.20	1.76	1.25	0.12	0.31	0.01		-1.36	0.10	1.51	0.08		-1.27			-1.26	-2.16	-1.73	-1.51	1.45	1.33	1.59	2.48		1.23	1.05	-1.16	1.44	0.12	1.18			-1.08
1.53	0.02	0.81	N/A	0.06	0.12	1.16	0.21	0.10	1.15	1.52	1.05	0.17	0.24	1.22	0.15	1.13	1.25	0.00	1.37	0.02	0.04	0.51	0.14	0.22	0.07	0.05	0.17	0.61	N/A	0.19	0.05	0.05	0.06	1.55	0.11	0.04	0.05	0.01

### Chapter 10: Appendix

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SPO 1380       by pothetical protein         SPO 1381       by pothetical protein         SPO 1382       cadD       c oxidase, aa3-type, submit I (EC: 1.9.3.1)         SPO 1383       cadD       c oxidase, aa3-type, submit I (EC: 1.9.3.1)         SPO 1384       mark family transcriptional regulator twin-arginine translocation pathway signal sequence domain-containing protein         SPO 1385       HIT family protein         SPO 1386       LuxR family transcriptional regulator twin-arginine translocation         SPO 1389       lipB         SPO 1390       LuxR family protein         SPO 1391       LuxR family protein         SPO 1392       alkylhydroperoxidase         SPO 1393       Rrf2 family protein         SPO 1394       beta-lactamase         SPO 1395       AraC family transcriptional regulator         SPO 1396       Rrf2 family protein         SPO 1397       Nard C family transcriptional regulator         SPO 1398       metG         SPO 1400       hydrophobe/amphiphile efflux-1 family protein         SPO 1401       metG         SPO 1402       metG         SPO 1403       metG         SPO 1404       typothetical protein         SPO 1405       metG         SPO 1406	S4 domain-containing protein		SPO1418
ctaD tipB rmetG rpoH-2	helicase		SPO1417
ctaD lipB rtuD rpoH-2	hypothetical protein		SPO1416
ctaD lipB metG rhuD rpoH-2	sterol carrier family protein		SPO1415
ctaD lipB rtuD rtuD rpoH-2	alpha/beta hydrolase		SPO1414
ctaD lipB rituD rpoH-2		pepF	SPO1413
ctaD lipB rluD rpoH-2	DNA modification methyltransferase domain- containing protein		SPO1412
ctaD lipB rtuD rpoH-2	renal dipeptidase		SPO1410
rtuD rtuD		rpoH-	SPO1409
metG		rluD	SPO1408
nne(G	hypothetical protein		SPO1407
nnetG	hypothetical protein		SPO1406
nnetG	tyrosinase domain-containing protein		SPO1405
nnet G	hypothetical protein		SPO1404
lipB lipB	synthetase (EC:6.1.1.10)	metG	SPO1403
ctaD lipB	hypothetical protein		SPO1402
ctaD lipB	hypothetical protein		SPO1401
ctaD lipB	hypothetical protein		SPO1400
lip B	AraC family transcriptional regulator		SPO1399
tipB	RND family efflux transporter MFP subunit		SPO1398
ipB ip	hydrophobe/amphiphile efflux-1 family proteir		SPO1397
ctaD lipB	transporter		SPO1396
lipB	AraC family transcriptional regulator		SPO1395
lipB	beta-lactamase		SPO1394
ctaD lipB	Rrf2 family protein		SPO1393
ctaD lipB	alkylhydroperoxidase		SPO1392
ctaD lipB	thioredoxin		SPO1390
ctaD	ligase B	lipB	SPO1389
ctaD	LuxR family transcriptional regulator		SPO1388
ctaD	cation efflux system protein		SPO1387
ctaD	HIT family protein		SPO1386
ctaD	twin-arginine translocation pathway signal sequence domain-containing protein		SPO1385
ctaD	MarR family transcriptional regulator		SPO1384
	c oxidase, aa3-type, subunit I (EC:1.9.3.1)	ctaD	SPO1383
	hypothetical protein		SPO1382
	hypothetical protein		SPO1381
	hypothetical protein		SPO1380

1.19	2.48	-2.36	1.40	0.99	1.48	-1.22	-2.67	1.53	-1.14	×	-1.05	1.26	0.97	-1.62	1.53	1.20	2.66	2.26	-1.58	1.69	1.26	1.35	x	-1.14	-1.27	1.07	2.61	1.97	-3.30	1.37	2.60	-1.36	-2.08	5.30	-1.36	1.66	1.64
0.97	0.00	0.03	0.02	0.61	0.39	0.65	0.30	0.57	0.02	х	0.39	0.07	0.04	0.30	0.48	0.97	0.01	0.02	0.00	0.01	0.35	0.65	×	0.55	0.25	0.81	0.34	0.21	0.00	0.64	0.00	0.84	0.10	0.05	0.27	0.03	0.70
1.96	1.13	-1.19	0.99	-1.56	-1.53	-1.23	-1.13	1.11	-1.92	1.12	1.19	1.32	1.27	-1.03	1.36	1.46	1.22	-1.12	-1.19	2.10	1.70	1.72	-1.12	-1.10	-1.62	-1.06	-1.50	1.00	1.02	1.15	1.35	-1.18	-1.10	1.37	-1.15	1.29	1.08
0.21	0.65	0.14	0.81	0.01	0.12	0.84	0.61	0.88	0.04	0.81	0.10	0.39	0.16	0.57	0.32	0.03	0.16	0.59	0.35	0.03	0.04	0.35	0.67	0.19	0.07	0.70	0.05	1.00	0.82	0.03	0.01	0.87	0.86	0.18	0.41	0.16	0.70
1.58	1.81	-1.78	1.19	-0.29	-0.03	-1.23	-1.90	1.32	-1.53	N/A	0.07	1.29	1.12	-1.33	1.45	1.33	1.94	0.57	-1.39	1.90	1.48	1.54	N/A	-1.12	-1.45	0.01	0.56	1.49	-1.14	1.26	1.98	-1.27	-1.59	3.34	-1.26	1.48	1.36
0.39	0.68	0.59	0.21	1.27	1.51	0.01	0.77	0.21	0.39	N/A	1.12	0.03	0.15	0.30	0.09	0.13	0.72	1.69	0.20	0.20	0.22	0.18	N/A	0.02	0.18	1.07	2.06	0.49	2.16	0.11	0.63	0.09	0.49	1.97	0.11	0.18	0.28
1.78	2.59	-1.51	1.80	1.81	2.83	-1.46	-3.84	-1.19	3.86	2.52	2.39	-1.35	-2.70	-1.94	4.63	-3.74	1.35	-0.98	-1.74	-2.77	-2.27	-1.43	1.83	1.60	-1.60	1.61	3.02	5.86	-1.28	1.42	2.93	1.07	1.15	3.84	0.91	2.70	1.17
0.76	0.01	0.15	0.01	0.09	0.06	0.85	0.01	1.00	0.00	0.04	0.01	0.27	0.02	0.01	0.02	0.00	0.11	0.08	0.11	0.01	0.00	0.64	×	0.03	0.33	0.12	0.31	0.02	0.35	0.58	0.00	0.97	0.03	0.06	0.49	0.00	0.86
-1.08	1.07	1.50	1.45	1.40	1.04	1.14	1.47	1.71	1.33	-1.35	-1.09	1.78	1.96	1.10	-1.14	1.62	1.38	1.26	1.15	1.16	-1.01	1.15	-0.98	-1.22	-1.09	-1.41	1.01	-1.14	1.29	-1.18	1.21	1.07	1.25	1.31	-1.04	-1.01	1.29
0.70	0.99	0.06	0.05	0.07	0.83	0.70	0.18	0.34	0.02	0.46	0.37	0.19	0.00	0.83	0.19	0.31	0.04	0.66	0.50	0.30	0.72	0.61	0.52	0.39	0.62	0.09	0.97	0.26	0.18	0.30	0.14	0.96	0.73	0.25	0.28	0.76	0.26
0.35	1.83	-0.01	1.63	1.61	1.94	-0.16	-1.19	0.26	2.60	0.59	0.65	0.22	-0.37	-0.42	1.75	-1.06	1.37	0.14	-0.30	-0.81	-1.64	-0.14	0.43	0.19	-1.35	0.10	2.02	2.36	0.01	0.12	2.07	1.07	1.20	2.58	-0.07	0.85	1.23
1.43	0.76	1.51	0.18	0.20	0.90	1.30	2.66	1.45	1.27	1.94	1.74	1.57	2.33	1.52	2.89	2.68	0.01	1.12	1.45	1.97	0.63	1.29	1.40	1.41	0.26	1.51	1.01	3.50	1.29	1.30	0.86	0.00	0.05	1.27	0.97	1.86	0.06
1.60	1.13	1.03	2.05	1.58	1.34	1.16	1.48	1.78	2.31	-1.49	1.03	1.84	1.60	-1.15	-1.08	1.00	-1.13	-1.11	-1.10	1.14	1.22	1.42	1.16	-1.23	-1.16	1.06	1.13	1.19	1.26	1.41	-1.21	-1.29	1.19	1.11	-1.27	-1.04	-1.38
0.44	0.80	0.90	0.00	0.00	0.11	0.61	0.16	0.14	0.02	0.23	0.87	0.29	0.01	0.54	0.20	0.14	0.19	0.57	0.22	0.09	0.56	0.10	0.24	0.09	0.28	0.55	0.83	0.32	0.08	0.09	0.14	0.81	0.58	0.76	0.12	0.77	0.37
1.22	1.43	1.22	2.39	1.68	1.55	1.30	1.63	2.78	2.44	-2.36	-1.21	1.55	2.07	1.85	-1.45	1.07	-1.47	1.39	1.46	1.62	1.50	1.60	-1.07	-0.93	-1.90	-1.03	-1.09	-1.28	-0.93	1.26	-1.18	1.14	-1.24	1.21	1.02	-1.66	-1.04
0.74	0.54	0.58	0.04	0.12	0.25	0.47	0.04	0.09	0.00	0.28	0.40	×	0.10	0.16	0.07	0.89	0.17	0.74	0.27	0.04	0.01	0.14	×	0.12	0.02	0.74	0.73	0.26	0.23	0.23	0.14	0.89	0.78	0.98	0.80	0.01	0.94
1.41	1.28	1.13	2.22	1.63	1.45	1.23	1.56	2.28	2.38	-1.93	-0.09	1.70	1.84	0.35	-1.27	1.04	-1.30	0.14	0.18	1.38	1.36	1.51	0.04	-1.08	-1.53	0.02	0.02	-0.05	0.17	1.34	-1.20	-0.08	-0.03	1.16	-0.13	-1.35	-1.21
0.19	0.15	0.10	0.17	0.05	0.11	0.07	0.08	0.50	0.06	0.44	1.12	0.15	0.24	1.50	0.19	0.04	0.17	1.25	1.28	0.24	0.14	0.09	1.12	0.15	0.37	1.05	1.11	1.24	1.09	0.08	0.02	1.22	1.22	0.05	1.15	0.31	0.17

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SPU1455	SPU1434	SDU1424	SPO1453	SPO1452	SPO1451	SPO1450	SPO1449	SPO1448	SPO1447	SPO1446	SPO1445	SPO1444	SPO1443	SPO1442	SPO1441	SPO1440	SPO1439	SPO1438	SPO1437	SPO1436	SPO1435	SPO1434	SPO1433	SPO1432	SPO1431	SPO1430	SPO1429	SPO1428	SPO1427	SPO1426	SPO1425	SPO1424	SPO1423	SPO1422	SPO1421	SPO1420	SPO1419
													rhlE						fabG								omp28			hppD			cobT				
TRAP dicarboxylate transporter subunit DctQ	TRAP dicarooxylate transporter subunit Detr	TD AD disorboxylata transportar subunit DotD	MarR family transcriptional regulator	oxidoreductase	aromatic 1,2-dioxygenase subunit alpha	aromatic 1,2-dioxygenase subunit beta	AMP-binding protein	hypothetical protein	oxidoreductase	cyclase	oxidoreductase	hypothetical protein	RNA helicase RhIE	hypothetical protein	fatty acid desaturase	hypothetical protein	DNA binding protein	hypothetical protein	reductase (EC:1.1.1.100)	dehydrogenase/thioesterase (EC:1.1.1.35)	hypothetical protein	AraC family transcriptional regulator	aldo/keto reductase	rhodanese domain-containing protein	O-acetylhomoserine aminocarboxypropyltransferase (EC:2.5.1.49)	antibiotic efflux protein	membrane protein, 28	hypothetical protein	hypothetical protein	dioxygenase (EC:1.13.11.27)	AsnC family transcriptional regulator	grutatione-regulated potassium-ernux system protein	phosphoribosyltransferase (EC:2.4.2.21)	cobalamin 5'-phosphate synthase (EC:2.7.8.26)	cation channel family protein	CarD family transcriptional regulator	(Fe-S)-binding protein
1.10	1.41	1 /1	1.06	-1.10	1.29	1.06	-1.39	-1.04	1.31	1.07	-1.20	1.46	1.51	-0.94	Х	7.46	-1.15	-1.68	3.46	16.30	11.90	2.03	1.13	-1.22	1.92	-1.09	0.99	-3.02	1.07	-3.11	-1.30	-1.16	х	Х	2.30	-1.61	1.09
0.58	0.87	0 07	0.97	0.83	0.64	0.36	0.02	0.35	0.92	0.92	0.18	0.14	0.11	0.39	×	0.01	0.92	0.27	0.10	0.00	0.00	0.04	0.58	0.71	0.08	0.59	0.09	0.02	0.18	0.01	0.21	0.05	×	x	0.24	0.01	0.73
-0.99	-1.19	1 10	1.01	-1.14	-1.19	-1.17	-1.08	1.15	1.28	1.19	-1.02	-1.21	-1.10	1.17	-1.05	1.26	-1.34	-1.26	1.47	2.21	1.95	-1.18	-1.15	-1.22	-1.28	1.16	-2.83	-1.34	1.21	-1.02	-1.42	1.26	2.14	1.33	2.57	-1.14	1.16
0.78	0.20	90.00	0.95	0.70	0.46	0.72	0.53	0.05	0.34	0.04	0.80	0.59	0.27	0.34	0.89	0.55	0.55	0.61	0.28	0.00	0.04	0.04	0.04	0.21	0.37	0.66	0.03	0.12	0.21	0.95	0.42	0.08	0.00	0.03	0.01	0.39	0.30
0.06	0.11	0 1 1	1.04	-1.12	0.05	-0.05	-1.24	0.05	1.30	1.13	-1.11	0.13	0.21	0.11	N/A	4.36	-1.25	-1.47	2.47	9.26	6.93	0.43	-0.01	-1.22	0.32	0.03	-0.92	-2.18	1.14	-2.07	-1.36	0.05	N/A	N/A	2.44	-1.38	1.13
1.04	1.50	1 20	0.03	0.02	1.24	1.12	0.16	1.10	0.02	0.06	0.09	1.34	1.31	1.06	N/A	3.10	0.10	0.21	1.00	7.05	4.98	1.61	1.14	0.00	1.60	1.13	1.91	0.84	0.07	1.05	0.06	1.21	N/A	N/A	0.14	0.24	0.03
-2.94	C0.7-	20 5	1.05	-4.34	-6.08	-3.67	-3.87	-1.87	-1.52	-2.60	-2.90	1.84	2.65	2.97	1.41	1.38	-1.05	-4.74	-1.38	-1.12	-1.55	4.12	2.36	-1.48	1.42	1.14	2.48	-3.64	-1.03	-7.49	-1.40	1.04	2.84	2.67	1.70	0.92	4.40
0.00	0.01	0 5 1	0.91	0.17	0.00	0.00	0.01	0.02	0.84	0.01	0.01	0.05	0.01	0.03	х	0.04	0.94	0.03	0.59	0.61	0.01	0.02	0.03	0.54	0.41	0.28	0.00	0.01	0.82	0.00	0.12	0.14	0.00	0.02	0.05	0.04	0.00
-1.02	1.20	1 20	1.17	-1.01	1.33	-1.01	1.14	1.13	1.36	1.24	1.54	1.43	1.03	-1.44	х	-1.26	-1.43	2.22	-1.02	-1.03	1.09	-1.16	1.39	1.26	1.16	1.49	1.05	-1.36	1.09	-1.25	-1.07	1.01	-1.37	-1.35	1.66	1.33	1.19
1.00	0.34	0.34	0.67	0.95	0.05	0.94	0.55	0.62	0.15	0.16	0.01	0.08	0.58	0.05	х	0.70	0.44	0.08	0.99	0.87	0.52	0.23	0.06	0.51	0.42	0.15	0.69	0.36	0.54	0.15	0.78	0.91	0.13	0.05	0.03	0.18	0.25
-1.98	-0.79	0.70	1.11	-2.68	-2.38	-2.34	-1.37	-0.37	-0.08	-0.68	-0.68	1.64	1.84	0.77	N/A	0.06	-1.24	-1.26	-1.20	-1.08	-0.23	1.48	1.88	-0.11	1.29	1.32	1.77	-2.50	0.03	-4.37	-1.24	1.03	0.74	0.66	1.68	1.13	2.80
0.96	2.07	2010 2010	0.06	1.67	3.71	1.33	2.51	1.50	1.44	1.92	2.22	0.21	0.81	2.21	N/A	1.32	0.19	3.48	0.18	0.05	1.32	2.64	0.49	1.37	0.13	0.18	0.72	1.14	1.06	3.12	0.17	0.02	2.11	2.01	0.02	0.21	1.61
1.26	c.'T	1 75	1.04	-1.21	-1.54	-1.17	-1.46	-1.01	1.03	-1.08	-1.43	-1.20	-1.09	1.74	-1.07	-0.98	1.10	1.09	-0.98	-1.29	-1.47	-1.49	-1.25	-1.54	-1.24	1.99	3.40	-1.44	1.16	-1.47	1.08	-1.05	-1.17	1.09	-1.17	1.01	-1.09
0.20	0.00	0.05	0.84	0.70	0.54	0.42	0.09	1.00	0.82	0.34	0.24	0.18	0.25	0.03	0.93	0.53	0.81	0.88	0.90	0.62	0.01	0.01	0.03	0.13	0.09	0.02	0.01	0.09	0.31	0.02	0.88	0.68	0.33	0.24	0.39	0.24	0.37
-1.43	CK'T	1 02	-1.12	-1.26	-1.48	1.06	-1.86	-1.36	-1.28	-1.39	-1.90	-1.17	-1.13	-1.35	х	1.18	-1.23	1.36	1.27	-1.90	-2.02	-2.07	-1.14	-1.27	-1.19	1.61	3.38	-1.11	1.07	-1.01	1.11	-1.18	-3.13	-1.61	-1.09	1.32	-1.34
0.08	0.04	0.04	0.88	0.67	0.03	0.86	0.02	0.05	0.23	0.08	0.06	0.79	0.03	0.05	х	0.88	0.82	0.53	0.27	0.01	0.05	0.01	0.12	0.36	0.03	0.16	0.01	x	0.29	0.62	0.84	0.08	0.00	0.09	0.55	0.16	0.15
-0.09	1.84	1 0/	-0.04	-1.24	-1.51	-0.05	-1.66	-1.19	-0.13	-1.24	-1.67	-1.19	-1.11	0.20	N/A	0.10	-0.06	1.23	0.15	-1.60	-1.75	-1.78	-1.20	-1.41	-1.22		3.39		1.12	-1.24	1.10	-1.12		-0.26			-1.22
1.35		0.00	1.08	0.03	0.03	1.12	0.20	0.18	1.16	0.16	0.23	0.02	0.02	1.55	N/A	1.08	1.17	0.14	1.12	0.31	0.27	0.29	0.06	0.14	0.03	0.19	0.01		0.04	0.23	0.02	0.06	0.98	1.35	0.04		0.13

### Chapter 10: Appendix

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SPO1492	SPO1491	SPO1490	SPO1489	SPO1488	SPO1487	SPO1486	SPO1485	SPO1484	SPO1483	SPO1482	SPO1481	SPO1480	SPO1479	SPO1478	SPO1477	SPO1476	SPO1475	SPO1474	SPO1473	SPO1472	SPO1471	SPO1470	SPO1469	SPO1468	SPO1467	SPO1466	SPO1465	SPO1464	SPO1463	SPO1462	SPO1461	SPO1460	SPO1459	SPO1458	SPO1457	SPO1456
															leuD-2	leuC-2				hyuA																
permease/ATP-binding protein	permease	regulatory protein	indole acetimide hydrolase (EC:3.5.1)	LuxR family transcriptional regulator	hypothetical protein	hypothetical protein	sodium:galactoside symporter family protein	hypothetical protein	cyclopropane-fatty-acyl-phospholipid synthase	hypothetical protein	hypothetical protein	snort chain denydrogenase/reductase oxidoreductase	transcriptional activator	RNA polymerase sigma factor RpoE	dehydratase small subunit (EC:4.2.1.33)	isomerase large subunit (EC:4.2.1.33)	GntR family transcriptional regulator	tautomerase	hydantoin utilization protein B	utilization protein A	isochorismatase	isocitrate lyase	3-ketosteroid dehydrogenase	aminotransferase	3-hydroxybutyryl-CoA dehydrogenase (EC:1.1.1.157)	opine dehydrogenase	TRAP dicarboxylate transporter subunit DctP	TRAP dicarboxylate transporter subunit DctQ	TRAP dicarboxylate transporter subunit DctM	OmpA family protein	hypothetical protein	(EC:1.2.7.8)	indolepyruvate oxidereductase, IorA subunit	MarR family transcriptional regulator	thioesterase	TRAP C4-dicarboxylate transport system permease DctM
1.06	Х	X	-1.62	-1.23	2.51	4.64	2.71	2.32	2.70	3.47	3.34	1.91	1.20	1.73	-2.05	-1.38	-1.02	-1.43	-1.13	-1.09	-1.03	-1.26	-1.21	-1.31	1.38	1.17	-3.13	-4.08	-1.53	-1.36	1.04	-1.25	-1.20	-1.63	1.10	-1.89
0.81	×	Х	×	0.58	0.02	0.01	0.02	0.01	0.21	0.01	0.00	0.49	0.62	0.02	0.12	0.07	0.96	0.02	0.55	0.13	0.87	0.31	×	0.23	0.00	0.69	0.02	0.00	0.01	0.03	0.99	0.07	0.41	0.22	0.52	0.01
-1.09	х	Х	-1.04	-1.05	-1.21	1.11	-1.23	-1.02	1.18	-1.27	-1.19	1.08	-1.22	-1.13	-1.16	-1.39	-1.14	-1.33	-1.33	-1.37	-1.27	-1.24	1.10	1.11	-1.02	-1.33	-1.84	-1.84	-1.23	1.10	-1.20	1.22	1.17	-1.03	1.14	-1.43
0.69	×	х	0.98	0.93	0.67	0.53	0.09	0.95	0.56	0.86	×	0.87	0.10	0.21	0.61	0.35	0.38	0.23	0.03	0.04	0.36	0.43	0.56	0.56	0.90	0.12	0.29	0.06	0.10	0.48	0.61	0.58	0.18	0.98	0.59	0.20
-0.02	N/A	N/A	-1.33	-1.14	0.65	2.88	0.74	0.65	1.94	1.10	1.08	1.50	-0.01	0.30	-1.61	-1.39	-1.08	-1.38	-1.23	-1.23	-1.15	-1.25	-0.05	-0.10	0.18	-0.08	-2.49	-2.96	-1.38	-0.13	-0.08	-0.02	-0.02	-1.33	1.12	-1.66
1.08	N/A	N/A	0.29	0.09	1.86	1.77	1.97	1.67	0.76	2.37	2.27	0.42	1.21	1.43	0.45	0.01	0.06	0.05	0.10	0.14	0.12	0.01	1.16	1.21	1.20	1.25	0.65	1.12	0.15	1.23	1.12	1.24	1.19	0.30	0.02	0.23
1.21	Х	-1.63	-2.82	-1.92	-1.35	-1.94	-3.95	-3.53	-4.12	-5.44	-1.56	-2.17	-2.80	-1.49	-1.85	-1.30	-0.99	-2.62	-2.37	-2.49	-2.28	-2.57	-1.09	-2.37	-1.61	-2.69	-8.27	-9.78	-2.59	-1.45	-2.43	-1.33	-1.71	-2.74	3.16	-1.73
0.19	×	0.45	0.02	0.22	0.50	0.03	0.01	0.02	0.13	0.02	0.11	0.69	0.00	0.01	0.18	0.61	0.56	0.00	0.06	0.01	0.07	0.02	0.16	0.03	0.23	0.18	0.01	0.00	0.01	0.05	0.10	0.30	0.12	0.11	0.01	0.07
-1.12	x	Х	-1.16	1.14	-1.02	-0.98	-1.06	Х	-1.00	-1.10	x	-0.97	-0.99	1.11	-0.99	1.11	1.19	-1.28	-1.25	-1.08	-0.99	1.03	х	-1.05	-1.32	-1.12	-1.01	-1.40	1.20	1.24	1.03	-1.32	-1.21	-1.36	-1.53	0.99
0.80	×	Х	x	0.89	×	0.79	0.93	х	0.91	0.95	×	0.91	0.92	0.67	0.98	0.51	0.35	0.13	0.39	0.59	0.82	0.77	x	0.83	x	0.72	0.83	0.25	0.32	0.36	0.93	0.49	0.11	0.78	0.02	0.67
0.04	N/A	N/A	-1.99	-0.39	-1.19	-1.46	-2.51	N/A	-2.56	-3.27	N/A	-1.57	-1.89	-0.19	-1.42	-0.10	0.10	-1.95	-1.81	-1.79	-1.64	-0.77	N/A	-1.71	-1.47	-1.91	-4.64	-5.59	-0.70	-0.11	-0.70	-1.33	-1.46	-2.05	0.82	-0.37
1.17	N/A	N/A	0.83	1.53	0.17	0.48	1.45	N/A	1.56	2.17	N/A	0.60	0.91	1.30	0.43	1.21	1.09	0.67	0.56	0.71	0.65	1.80	N/A	0.66	0.15	0.79	3.63	4.19	1.90	1.35	1.73	0.01	0.25	0.69	2.35	1.36
-1.26	Х	-0.98	-1.09	1.06	-0.99	-1.27	-1.54	-1.00	-0.96	-1.54	x	-0.99	-1.00	1.14	1.34	1.19	1.22	-0.99	1.16	-1.20	-1.25	-1.24	1.41	-1.18	1.08	1.27	1.28	-1.41	1.08	1.20	1.06	-0.97	-1.14	-1.15	1.07	-1.18
0.62	×	0.66	0.83	0.91	0.90	0.19	0.15	0.64	0.81	0.78	×	0.93	0.96	0.62	0.04	0.30	0.29	0.68	0.25	0.04	0.10	0.19	0.08	0.50	0.51	0.38	0.17	0.09	0.28	0.11	0.70	0.84	0.18	0.86	0.31	0.46
-1.07			Х	1.32	х	-1.05	-1.06	Х	-1.15	1.21	Х	-1.02	1.24	1.67	1.30	-0.92	-1.01	x	-1.24	-1.18	-1.71	-1.64	Х	-1.27	×	-0.93	2.64	2.05	1.35	1.30	1.29	-1.31	-1.12	-1.10	-1.67	2.39
7	×	Х		3		с,	9		<u>о</u> г				-																							
0.81	X X	Х Х	x	0.68	Х	0.75	6 0.68	х	5 0.53	0.88	x	x	0.24	0.11	0.33	0.58	0.60	Х	0.64	0.20	0.04	0.01	x	0.44	×	0.26	0.13	0.13	0.50	0.17	0.18	0.43	0.88		0.05	0.08
											X N/A								0.64 -0.04	0.20 -1.19		0.01 -1.44	X N/A	0.44 -1.23	X N/A	0.26 0.17	0.13 1.96	0.13 0.32	0.50 1.22	0.17 1.25	0.18 1.18	0.43 -1.14	-	0.97	0.05 -0.30	0.08 0.61

### Chapter 10: Appendix

01530	01529	01528	01527	01526	01525	01524	01523	01522	01521	01520	01519	01518	01517	01516	01515	01514	01513	01512	01511	01510	01509	01508	01507	01506	01505	01504	01503	01502	01501	01500	01499	01498	01497	01496	01495	01494	01493
										coxM-1	coxL-1	coxS-1	coxG													pqqA	pqqB	pqqC	pqqD	pqqE							
integral membrane protein MviN	hypothetical protein	hypothetical protein	hypothetical protein	RND family efflux transporter MFP subunit	AcrB/AcrD/AcrF family transporter	hypothetical protein	metallo-beta-lactamase	hypothetical protein	regulatory protein	carbon monoxide dehydrogenase, medium subunit	(EC:1.2.99.2)	dehydrogenase, small subunit (EC:1.2.99.2)	monoxide dehydrogenase operon G protein	protein	cytochrome c550	hypothetical protein	hypothetical protein	hypothetical protein	ABC transporter ATP-binding protein	ABC transporter permease	cytochrome c family protein	quinoprotein ethanol dehydrogenase (EC:1.1.99)	LuxR family transcriptional regulator	GfdT protein	EC:2.7.3)	PQQ biosynthesis protein A	quinone biosynthesis protein PqqB	quinone biosynthesis protein PqqC	PQQ synthesis protein D	quinone biosynthesis protein PqqE	alpha/beta hydrolase	fumarate hydratase, class I	RND family efflux transporter MFP subunit	ABC transporter permease	ABC transporter ATP-binding protein	2-hydroxychromene-2-carboxylate isomerase	branched-chain amino acid ABC transporter ATP- binding protein
1.03	-1.14	Х	1.90	1.48	1.33	-1.93	-1.97	1.24	-1.74	-1.46	0.91	-2.58	-1.31	1.15	-1.21	-1.60	1.19	1.02	1.03	-1.06	-2.11	-2.08	-1.17	-1.77	1.00	0.81	-1.16	1.04	-1.10	-1.16	1.35	1.21	2.76	4.93	4.19	-1.25	×
0.99	0.43	x	0.09	0.32	0.92	0.01	0.00	0.54	0.01	0.07	0.01	0.00	0.28	0.88	0.02	0.34	0.93	0.40	0.74	0.06	0.01	0.07	0.09	0.01	0.21	0.01	0.33	0.34	0.84	0.14	0.14	0.80	0.18	0.07	0.15	0.10	×
1.19	-1.16	х	-1.22	-1.17	1.02	2.80	1.97	1.48	3.31	1.56	1.79	1.56	4.37	3.97	7.22	4.34	6.07	2.58	2.22	1.76	1.39	2.96	3.53	-1.12	1.07	4.72	4.75	2.33	3.43	3.25	-1.17	2.26	1.28	2.20	2.19	1.21	-1.26
0.72	0.50	×	0.90	0.81	0.96	0.00	0.01	0.08	0.02	0.08	0.05	0.04	0.00	0.12	0.00	0.01	0.00	0.00	0.01	0.01	0.41	0.08	0.06	0.03	0.53	0.00	0.03	0.11	0.11	0.03	0.30	0.02	0.56	0.04	0.03	0.13	0.77
1.11	-1.15	N/A	0.34	0.16	1.18	0.44	0.00	1.36	0.79	0.05	1.35	-0.51	1.53	2.56	3.01	1.37	3.63	1.80	1.63	0.35	-0.36	0.44	1.18	-1.45	1.03	2.77	1.80	1.69	1.17	1.05	0.09	1.74	2.02	3.57	3.19	-0.02	N/A
0.08	0.01	N/A	1.56	1.33	0.16	2.37	1.97	0.12	2.53	1.51	0.44	2.07	2.84	1.41	4.22	2.97	2.44	0.78	0.60	1.41	1.75	2.52	2.35	0.33	0.04	1.95	2.96	0.65	2.27	2.21	1.26	0.53	0.74	1.37	1.00	1.23	N/A
-1.01	-2.47	Х	-3.18	1.59	1.60	-44.50	-29.80	2.29	-10.50	-16.10	-23.10	-30.10	-12.40	-12.00	-35.80	-21.80	-17.00	-11.50	-4.46	-9.02	-56.80	-65.20	-13.90	-7.45	-5.09	-47.00	-10.70	-8.58	-4.40	-13.70	1.23	1.22	1.42	2.80	2.36	-1.70	-1.53
0.91	0.05	х	0.05	0.10	0.84	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.01	0.00	0.00	0.00	0.00	0.09	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.04	0.00	0.27	0.00	0.02	0.38	0.62	0.12	0.18	0.01	0.45
1.24	1.04	х	х	-1.25	-0.99	-1.01	1.07	1.06	1.16	2.23	2.60	2.51	1.34	1.09	-1.50	-1.62	-1.47	1.13	1.12	1.06	-1.37	-1.01	-1.03	1.32	1.11	1.43	-1.16	-1.26	-1.18	1.07	1.43	1.04	1.26	1.49	1.48	1.26	-1.59
0.63	0.89	x	Х	0.79	0.88	0.99	0.38	0.77	0.23	0.01	0.01	0.01	0.14	0.93	0.08	0.33	0.17	0.20	0.16	0.21	0.15	0.87	0.83	0.10	0.78	0.10	0.73	0.39	0.77	0.68	0.11	0.93	0.65	0.30	0.07	0.21	×
0.12	-0.72	N/A	N/A	0.17	0.31	-22.76	-14.37	1.68	-4.67	-6.94	-10.25	-13.80	-5.53	-5.46	-18.65	-11.71	-9.24	-5.19	-1.67	-3.98	-29.09	-33.11	-7.47	-3.07	-1.99	-22.79	-5.93	-4.92	-2.79	-6.32	1.33	1.13	1.34	2.15	1.92	-0.22	-1.56
1.13	1.76	N/A	N/A	1.42	1.29	21.75	15.44	0.62	5.83	9.17	12.85	16.31	6.87	6.55	17.15	10.09	7.77	6.32	2.79	5.04	27.72	32.10	6.44	4.39	3.10	24.22	4.77	3.66	1.61	7.39	0.10	0.09	0.08	0.65	0.44	1.48	0.03
1.99	-1.19	-1.12	-1.18	-0.99	1.24	5.56	2.78	1.73	2.12	2.57	4.11	3.27	4.23	4.57	7.73	6.01	5.50	2.30	3.11	1.85	2.11	4.50	2.80	1.23	1.41	3.69	4.41	3.52	3.51	2.11	-1.05	1.57	1.75	1.67	1.33	-1.06	-1.18
0.32	0.85	0.34	0.71	0.97	0.40	0.00	0.01	0.02	0.08	0.02	0.02	0.01	0.00	0.09	0.00	0.02	0.00	0.00	0.00	0.01	0.02	0.06	0.02	0.23	0.05	0.00	0.03	0.00	0.06	0.02	0.70	0.03	0.28	0.09	0.27	0.84	0.33
1.52	1.11	Х	2.00	1.49	-0.88	2.81	2.08	1.41	-1.27	3.78	6.22	6.00	4.91	2.62	5.27	4.48	3.95	1.99	2.02	1.22	Х	5.52	2.17	1.09	1.33	7.03	3.30	2.30	2.65	1.97	1.05	1.20	2.84	1.73	1.50	-1.29	1.51
X	0.42	×	x	0.59	0.11	0.02	0.03	0.04	x	0.00	0.00	0.00	0.00	0.21	0.00	0.02	0.03	0.03	0.01	0.12	x	0.08	0.02	0.45	0.58	0.00	0.05	0.14	0.13	0.05	0.55	0.49	0.15	0.10	0.18	0.50	×
1.76	-0.04	N/A	0.41	0.25	0.18	4.19	2.43	1.57	0.43	3.18	5.17	4.64	4.57	3.60	6.50	5.25	4.73	2.15	2.57	1.54	N/A	5.01	2.49	1.16	1.37	5.36	3.86	2.91	3.08	2.04	0.00	1.39	2.30	1.70	1.42	-1.18	0.17
0.24	1.15	N/A	1.59	1.24	1.06	1.38	0.35	0.16	1.70	0.60	1.06	1.37	0.34	0.97	1.23	0.76	0.78	0.16	0.54	0.31	N/A	0.51	0.32	0.07	0.04	1.67	0.55	0.61	0.43	0.07	1.05	0.19	0.55	0.03	0.09	0.12	1.35

### Chapter 10: Appendix

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SPO1564	SPO1563	SPO1562	SPO1561	SPO1560	SPO1559	SPO1558	SPO1557	SPO1556	SPO1555	SPO1554	SPO1553	SPO1552	SPO1551	SPO1550	SPO1549	SPO1548	SPO1547	SPO1546	SPO1545	SPO1544	SPO1543	SPO1542	SPO1541	SPO1540	SPO1539	SPO1538	SPO1537	SPO1536	SPO1535	SPO1534	SPO1533	SPO1532	SPO1531
				folD			fhs-1																										
MOFRL domain-containing protein	hypothetical protein	glycine cleavage system T protein	hypothetical protein	5,10-methylene-tetrahydrofolate dehydrogenase/5,10-methylene-tetrahydrofolate cyclohydrolase	bifunctional 5, 10-methylene-tetrahydrofolate dehydrogenase/5, 10-methylene-tetrahydrofolate cyclohydrolase	chorismate mutase	ligase (EC:6.3.4.3)	formate dehydrogenase subunit alpha (EC:1.2.1.2)	formate dehydrogenase subunit beta (EC:1.2.1.2)	ammonium transporter	GntR family transcriptional regulator	ABC transporter substrate-binding protein	flavin-containing monooxygenase	glycine betaine/proline ABC transporter permease	glycine betaine/proline ABC transporter ATP- binding protein (EC:3.6.3.32)	glycine betaine/proline ABC transporter substrate- binding protein	behuue/opine/mexet upiake ABC transporter ATF-	binding protein binding protein		permease permease permetasino/nickal untaka ABC transporter	peptide/opine/nickel uptake ABC transporter substrate-binding protein nentide/opine/nickel uptake ABC transporter	renal dipeptidase	LysR family transcriptional regulator	glycosyl transferase family protein	O-antigen polymerase	chain length determinant protein	sequence domain-containing protein	group 1 family glycosyltransferase	polysaccharide deacetylase	sugar transferase	glycoside hydrolase	hypothetical protein	hypothetical protein
1.19	1.45	1.79	-0.97	1.08	11.70	5.55	11.90	2.34	3.84	-1.68	-1.63	-2.08	-1.06	-2.12	-2.01	-2.22	-3.31	-2.82	-2.17	-4.28	-3.79	-2.93	-4.93	1.16	1.32	1.26	1.88	1.43	1.36	1.30	-1.35	-1.02	-1.15
0.08	0.21	0.12	0.57	×	0.21	0.33	0.00	0.01	0.09	0.03	0.27	0.04	0.88	×	0.01	0.00	0.01	0.00	0.08	0.01	0.00	0.27	0.00	0.76	0.11	0.15	0.08	0.16	0.88	0.08	0.60	1.00	0.67
1.23	1.28	1.21	1.54	1.64	2.54	2.39	2.36	-1.03	1.34	-1.17	-1.13	-1.14	-1.48	-1.55	-1.86	-1.72	-2.81	-3.25	-3.08	-2.90	-6.49	-3.79	-2.05	1.13	-1.06	1.44	1.45	1.50	1.39	1.15	1.01	1.01	1.33
0.44	0.03	0.04	0.03	0.01	0.01	0.08	0.04	0.94	0.58	0.20	0.85	0.52	0.32	0.82	0.04	0.07	0.02	0.01	0.12	0.12	0.00	0.14	0.01	0.44	0.88	0.04	0.01	0.09	0.21	0.28	0.93	0.77	0.34
1.21	1.37	1.50	0.29	1.36	7.12	3.97	7.13	0.66	2.59	-1.43	-1.38	-1.61	-1.27	-1.84	-1.94	-1.97	-3.06	-3.04	-2.63	-3.59	-5.14	-3.36	-3.49	1.15	0.13	1.35	1.67	1.47	1.38	1.23	-0.17	-0.01	0.09
0.02	0.09	0.29	1.25	0.28	4.58	1.58	4.77	1.69	1.25	0.26	0.25	0.47	0.21	0.29	0.07	0.25	0.25	0.22	0.46	0.69	1.35	0.43	1.44	0.02	1.19	0.09	0.21	0.04	0.01	0.08	1.18	1.02	1.24
1.27	2.23	1.14	1.81	3.87	3.71	1.48	1.23	1.15	2.23	-1.62	1.06	-2.09	-1.29	-1.83	3.42	-6.14	-10.30	-8.41	-4.17	-13.00	-20.40	-10.40	-1.17	-1.21	-1.69	-1.01	-0.93	-1.32	1.14	-1.56	-1.10	1.16	-1.58
0.13	0.02	0.37	0.00	0.01	0.01	0.83	0.20	0.63	0.23	0.11	0.96	0.04	0.80	0.83	0.01	0.00	0.00	0.01	0.16	0.01	0.00	0.09	0.95	0.97	0.03	0.56	0.10	0.00	0.95	0.03	0.73	0.44	0.69
-4.03	1.19	1.36	-1.11	-1.32	-1.23	-1.30	-2.30	-1.25	-6.94	-1.03	1.22	1.13	-1.01	-1.28	-1.34	1.16	1.32	1.34	1.41	1.49	1.41	1.50	-1.55	1.50	1.49	1.91	1.79	1.63	1.24	1.28	-1.12	1.54	1.20
0.04	0.09	0.13	0.72	0.05	0.16	0.45	0.00	0.42	0.06	0.97	0.82	0.79	0.99	x	0.30	0.12	0.06	0.16	0.46	0.48	0.21	0.58	0.12	0.02	0.02	0.07	0.02	0.01	0.32	0.29	0.53	0.03	×
-1.38	1.71	1.25	0.35	1.28	1.24	0.09	-0.54	-0.05	-2.36	-1.33	1.14	-0.48	-1.15	-1.56	1.04	-2.49	-4.49	-3.54	-1.38	-5.76	-9.50	-4.45	-1.36	0.15	-0.10	0.45	0.43	0.16	1.19	-0.14	-1.11	1.35	-0.19
2.65	0.52	0.11	1.46	2.60	2.47	1.39	1.77	1.20	4.59	0.30	0.08	1.61	0.14	0.27	2.38	3.65	5.81	4.88	2.79	7.25	10.91	5.95	0.19	1.36	1.59	1.46	1.36	1.48	0.05	1.42	0.01	0.19	1.39
1.86	1.67	1.02	1.24	-1.25	-1.17	-1.27	-1.52	-1.55	1.09	-1.29	-1.04	-1.02	-1.02	1.16	-1.06	-1.23	1.12	1.29	1.26	-1.00	1.38	1.09	-1.41	-1.10	1.10	1.53	1.33	1.31	2.03	1.25	1.31	1.25	1.18
0.01	0.05	0.62	0.21	0.07	0.31	0.45	0.06	0.12	0.94	0.21	0.93	0.92	1.00	0.84	0.81	0.15	0.75	0.10	0.30	0.98	0.35	0.92	0.01	0.77	0.34	0.01	0.03	0.06	0.12	0.18	0.28	0.23	0.63
1.97	-1.01	-0.96	-1.55	-2.22	-1.70	-1.43	-1.80	-1.15	1.50	1.35	1.09	1.38	-0.95	1.56	1.64	-1.03	-1.74	-1.48	-1.26	-1.16	1.78	1.02	-1.24	1.21	1.20	1.32	-1.29	1.21	2.09	1.57	-1.45	-0.92	х
0.02		0.26	х	0.00		0.33	0.04	0.71	0.66	0.12	0.93	0.55	0.50	x	0.26	0.32	0.05	0.07	0.59	0.73	0.19	0.95	0.21	0.13	0.15	0.07	0.23	0.20	0.03	0.10	0.03	-	×
	0.33	0.03	-0.16	-1.74		-1.35		-1.35	1.30	0.03	0.03	0.18	-0.99	1.36	0.29	-1.13	-0.31	-0.10	0.00	-1.08	1.58	1.06	-1.33	0.05	1.15	1.43	0.02	1.26	2.06	1.41	-0.07		N/A
0.05	1.34	0.99	1.40	0.49		0.08		0.20	0.21	1.32	1.07	1.20	0.03	0.20	1.35	0.10	1.43	1.39	1.26	0.08	0.20	0.04	0.09	1.16	0.05	0.11	1.31	0.05	0.03	0.16	1.38		N/A

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SPO1603	SPO 1602	SPO1601	SPO1600	SPO1599	SPO1598	SPO1597	SPO1596	SPO1595	SPO1594	SPO1593	SPO1592	SPO1591	SPO1590	SPO1589	SPO1588	SPO 1587	SPO1586	SPO1585	SPO1584	SPO1583	SPO1582	SPO1581	SPO1580	SPO1579	SPO1578	SPO1577	SPO1576	SPO1575	SPO1574	SPO1573	SPO1572	SPO1571	SPO1570	SPO1569	SPO1568	SPO1567	SPO 1566	SPO1565
		panE-2				hemL	dddQ																							glnT	glyA	ppc			sucC			
malate/L-lactate dehydrogenase	trimethylamine methyltransferase	2-reductase (EC:1.1.1.169)	fumarylacetoacetate hydrolase	MmgE/PrpD family protein	hypothetical protein	2,1-aminomutase (EC:5.4.3.8)	lyase	mandelate racemase	mandelate racemase	alcohol dehydrogenase	glycine cleavage system protein T	SIS domain-containing protein	Rieske (2Fe-2S) domain-containing protein	carboxymuconolactone decarboxylase	sarcosine oxidase subunit beta	sarcosine oxidase subunit delta	sarcosine oxidase subunit alpha	sarcosine oxidase subunit gamma	AraC family transcriptional regulator	DNA-binding protein	hypothetical protein	oxidoreductase	hypothetical protein	glycine cleavage system protein T	ammonium transporter	hypothetical protein	glutamine amidotransferase	FwdC/FmdC family protein	glutamate synthase	synthetase (EC:6.3.1.2)	hydroxymethyltransferase (EC:2.1.2.1)	carboxylase (EC:4.1.1.31)	2-hydroxyacid dehydrogenase	succinyl-CoA synthetase subunit alpha (EC:6.2.1.5)	ligase subunit beta (EC:6.2.1.5)	aminotransferase	LuxR family transcriptional regulator	citrate lyase
-1.67	-1.11	-1.01	1.13	-1.90	2.75	2.40	2.84	-1.56	3.20	3.12	5.15	3.67	5.28	1.72	6.19	1.43	3.09	1.92	0.98	1.06	1.50	-1.03	1.03	-1.37	-1.41	-1.48	1.74	-1.16	1.08	×	3.16	-1.12	1.89	2.24	2.15	1.64	-1.36	1.46
0.09	0.54	x	0.15	0.34	0.01	0.17	0.02	0.14	0.06	0.01	0.04	0.00	0.15	0.01	0.01	0.88	0.00	0.84	0.95	0.92	0.37	0.44	0.58	0.29	×	0.38	0.61	0.89	0.81	x	0.00	0.74	0.04	0.03	0.02	0.52	0.21	0.07
-1.00	-1.36	-1.03	-1.27	-1.49	2.84	1.30	1.14	-1.30	1.47	1.05	1.24	1.07	1.27	1.12	1.19	1.07	-1.04	1.41	1.03	1.40	-1.01	-1.17	-1.17	-1.48	1.09	1.13	1.04	-1.13	-1.23	-1.20	2.03	-1.22	-1.17	-1.50	-1.38	-1.18	-1.34	2.01
0.92	0.73	1.00	0.23	0.34	0.03	0.22	0.22	0.68	0.40	0.50	0.14	0.13	0.09	0.10	0.21	0.82	0.90	0.24	0.92	0.23	0.99	0.14	0.03	0.47	0.79	0.17	0.88	0.25	0.37	0.24	0.07	0.35	0.39	0.22	0.05	0.36	0.54	0.00
-1.33	-1.24	-1.02	-0.07	-1.70	2.80	1.85	1.99	-1.43	2.34	2.09	3.20	2.37	3.28	1.42	3.69	1.25	1.03	1.67	1.00	1.23	0.25	-1.10	-0.07	-1.43	-0.16	-0.18	1.39	-1.15	-0.08	N/A	2.60	-1.17	0.36	0.37	0.39	0.23	-1.35	1.74
0.34	0.13	0.01	1.20	0.21	0.04	0.55	0.85	0.13	0.87	1.04	1.96	1.30	2.01	0.30	2.50	0.18	2.07	0.26	0.03	0.17	1.26	0.07	1.10	0.05	1.25	1.31	0.35	0.02	1.16	N/A	0.57	0.05	1.53	1.87	1.77	1.41	0.01	0.27
-2.22	2.27	-1.07	1.55	-2.75	2.85	-1.06	3.96	-2.70	-0.89	6.05	-1.29	-1.70	-1.23	1.29	1.33	-2.22	-0.89	1.23	-1.27	1.92	-1.13	-1.78	-1.94	-2.53	-2.27	-1.39	-1.13	-1.85	-1.66	-1.13	3.58	-1.20	-1.70	-2.17	-3.62	-1.76	-1.49	-1.11
0.05	0.04	0.44	0.02	0.08	0.01	0.99	0.00	0.01	0.55	0.00	0.72	0.10	0.58	0.02	0.26	0.70	0.05	0.90	0.88	0.22	0.56	0.02	0.04	0.12	x	0.10	0.88	0.05	0.02	0.42	0.01	0.55	0.11	0.10	0.01	0.52	0.20	0.83
1.44	-1.14	-1.08	-1.34	-1.13	1.57	1.16	1.18	-1.13	-1.45	-1.26	-1.60	-1.62	-1.79	1.15	х	2.69	1.36	2.15	-1.00	1.13	1.53	1.80	1.65	1.73	1.50	1.18	1.08	1.05	-1.06	-1.21	-1.58	-1.25	-1.52	-1.36	-1.18	-1.35	-1.61	-5.12
0.53	0.78	×	0.51	0.56	0.05	0.42	0.29	0.79	0.28	0.06	0.00	0.08	0.06	0.36	×	0.04	0.36	0.24	0.97	0.62	0.13	0.07	0.03	0.22	x	0.33	0.80	0.85	0.66	0.46	0.01	0.09	0.04	0.16	0.13	0.20	0.07	0.00
-0.39	0.57	-1.08	0.11	-1.94	2.21	0.05	2.57	-1.92	-1.17	2.40	-1.45	-1.66	-1.51	1.22	N/A	0.24	0.24	1.69	-1.13	1.53	0.20	0.01	-0.15	-0.40	-0.39	-0.11	-0.02	-0.40	-1.36	-1.17	1.00	-1.23	-1.61	-1.77	-2.40	-1.56	-1.55	-3.12
1.83	1.71	0.01	1.45	0.81	0.64	1.11	1.39	0.79	0.28	3.66	0.16	0.04	0.28	0.07	N/A	2.46	1.13	0.46	0.14	0.40	1.33	1.79	1.80	2.13	1.89	1.29	1.11	1.45	0.30	0.04	2.58	0.03	0.09	0.41	1.22	0.20	0.06	2.01
-1.38	1.17	1.23	-1.94	-1.38	-1.01	1.11	1.15	-1.04	1.12	-1.44	-0.98	-1.83	-1.17	1.04	-1.15	-1.23	-1.35	1.21	1.34	1.12	1.30	-1.03	1.02	-1.02	-0.96	-1.02	-1.56	-1.70	-1.35	1.17	-2.09	-0.99	1.07	-1.17	-1.26	-1.15	-1.42	1.37
0.29	0.52	0.27	0.07	0.64	0.92	0.03	0.25	0.79	0.61	0.06	0.58	0.03	0.50	0.74	0.57	0.52	0.53	0.42	0.32	0.61	0.32	0.54	0.89	0.92	0.85	0.95	0.16	0.02	0.24	0.31	0.01	0.85	0.61	0.41	0.05	0.63	0.09	0.12
-0.97	1.67	×	-2.56	1.93	1.20	-1.02	-1.06	2.11	-1.35	-1.32	-1.55	-2.51	-1.82	-1.12	×	-1.74	-0.96	x	1.25	-1.58	1.23	-1.03	1.10	1.16	x	-1.43	-1.52	-1.57	1.29	-1.06	-2.06	1.30	-1.00	-1.16	-1.12	1.42	-0.97	1.62
0.92	0.59	x	0.17	0.53	0.61	0.97	0.99	0.06	0.49	0.50	0.31	0.01	0.09	0.31	×	0.17	0.49	x	0.39	0.05	0.18	0.47	0.05	0.66	x	0.37	0.24	0.01	0.50	x	0.01	х	0.54	0.69	0.72	0.36	0.28	0.01
-1.17	1.42	N/A	-2.25	0.28	0.10	0.05	0.04	0.54	-0.12	-1.38	-1.26	-2.17	-1.50	-0.04	N/A	-1.49	-1.15	N/A	1.30	-0.23	1.27	-1.03	1.06	0.07	N/A	-1.23	-1.54	-1.64	-0.03	0.05	-2.08	0.16	0.04	-1.17	-1.19	0.14	-1.20	1.50
0.21	0.25	N/A	0.31	1.66	1.11	1.07	1.11	1.58	1.24	0.06	0.29	0.34	0.33	1.08	N/A	0.26	0.20	N/A	0.05	1.35	0.04	0.00	0.04	1.09	N/A	0.20	0.02	0.06	1.32	1.12	0.01	1.14	1.03	0.01	0.07	1.29	0.22	0.13

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SPO1642	SPO1641	SPO1640	SPO1639	SPO1638	SPO1637	SPO1636	SPO1635	SPO1634	SPO1633	SPO1632	SPO1631	SPO1630	SPO1629	SPO1628	SPO1627	SPO1626	SPO1625	SPO1624	SPO1623	SPO1622	SPO1621	SPO1620	SPO1619	SPO1618	SPO1617	SPO1616	SPO1615	SPO1614	SPO1613	SPO1612	SPO1611	SPO1610	SPO1609	SPO1608	SPO1607	SPO1606	SPO1605	SPO1604
					ssb		aroB	aroK		xerD																							potA	potB		potD		
hypothetical protein	beta-lactamase	LysR family transcriptional regulator	FadB domain-containing protein	transglycosylase	binding protein	SREBP protease	synthase (EC:4.2.3.4)	kinase (EC:2.7.1.71)	hypothetical protein	recombinase XerD	hypothetical protein	hypothetical protein	hypothetical protein	glycine cleavage system protein T	hypothetical protein	type I secretion target repeat-containing protein	serine protease	DNA-binding response regulator	sensor histidine kinase (EC:2.7.3)	cytochrome P450 family protein	TfoX domain-containing protein	glyoxalase	hypothetical protein	hypothetical protein	type I secretion target repeat-containing protein	transcriptional regulator	cyclic nucleotide-binding protein	hypothetical protein	hypothetical protein	twin-arginine translocation pathway signal sequence domain-containing protein	hypothetical protein	LuxR family transcriptional regulator	ABC transporter ATP-binding component	ABC transporter permease	spermidine/putrescine ABC transporter permease	ABC transporter substrate-binding protein	hypothetical protein	LysR family transcriptional regulator
-2.03	-1.21	-1.46	Х	-1.15	1.19	-1.36	-1.30	1.17	-2.19	-1.38	1.02	Х	-1.35	6.71	1.76	4.49	-2.43	-1.19	-1.75	5.05	-1.12	1.38	Х	-1.82	Х	Х	5.33	1.15	Х	-2.53	1.17	Х	-1.22	1.13	-1.40	-1.28	-1.04	1.62
0.12	0.27	0.01	×	0.17	0.97	0.71	0.31	0.37	×	0.02	0.99	×	0.06	0.01	0.05	0.00	0.24	0.10	0.02	0.01	0.11	0.16	×	0.08	×	×	0.04	0.98	×	×	0.97	×	0.72	0.27	0.05	0.24	0.15	0.21
-1.41	-1.32	-1.15	1.32	1.02	-1.26	1.01	1.95	2.28	-1.38	-1.81	1.00	-1.12	-1.06	2.76	1.31	2.89	-1.44	-1.25	-1.30	2.27	-1.04	1.08	-1.04	-1.38	-1.02	-1.52	3.35	1.14	-1.29	1.03	-1.08	1.07	-1.34	-1.01	-2.08	-1.61	1.02	-1.04
0.40	0.29	0.35	0.31	1.00	0.27	0.99	0.05	0.08	0.68	0.02	0.69	0.72	0.88	0.03	0.34	0.10	0.16	0.33	0.54	0.02	0.88	0.87	0.95	0.63	×	0.05	0.00	0.69	0.34	0.88	0.95	0.12	0.66	0.98	0.14	0.02	0.72	0.94
-1.72	-1.27	-1.31	N/A	-0.06	-0.04	-0.18	0.33	1.73	-1.79	-1.60	1.01	N/A	-1.21	4.74	1.54	3.69	-1.94	-1.22	-1.53	3.66	-1.08	1.23	N/A	-1.60	N/A	N/A	4.34	1.15	N/A	-0.75	0.04	N/A	-1.28	0.06	-1.74	-1.45	-0.01	0.29
0.31	0.06	0.16	N/A	1.09	1.23	1.19	1.63	0.56	0.41	0.21	0.01	N/A	0.15	1.98	0.22	0.80	0.50	0.03	0.23	1.39	0.04	0.15	N/A	0.22	N/A	N/A	0.99	0.01	N/A	1.78	1.13	N/A	0.06	1.07	0.34	0.17	1.03	1.33
-2.13	-2.01	-1.13	2.42	2.65	1.91	1.50	0.98	2.04	-2.23	1.23	1.43	1.77	-1.36	-2.77	1.29	-0.99	8.94	6.62	1.81	1.82	-1.54	2.32	Х	-1.62	-1.11	2.09	5.38	1.53	-1.05	-2.14	1.61	2.24	-2.59	-1.62	-2.67	-3.16	-3.07	-1.84
0.12	0.20	0.72	0.02	0.00	0.11	0.62	0.80	0.01	0.56	0.39	0.17	0.00	0.05	0.06	0.26	0.43	0.01	0.00	0.11	0.03	0.12	0.03	х	0.10	×	0.00	0.01	0.83	0.64	0.34	0.83	0.07	0.37	0.49	0.08	0.09	0.01	0.30
-1.02	1.09	-1.42	-1.17	0.99	1.08	-1.05	-1.26	1.23	-1.17	1.01	-1.28	-1.42	-1.17	-1.36	1.21	1.65	2.12	1.03	1.17	1.66	1.39	-2.12	Х	-1.54	х	-1.52	1.51	-1.32	Х	х	-1.35	-1.31	1.62	1.55	1.72	1.70	1.78	1.28
0.89	0.79	0.07	0.56	0.84	0.97	0.90	0.20	0.31	0.81	0.32	0.11	0.09	0.69	0.54	0.41	0.03	0.02	0.99	0.61	0.01	0.13	0.02	Х	0.60	x	0.05	0.07	0.64	Х	Х	x	0.29	0.44	0.14	0.06	0.04	0.01	0.29
-1.58	-0.46	-1.28	0.63	1.82	1.50	0.23	-0.14	1.64	-1.70	1.12	0.08	0.18	-1.27	-2.07	1.25	0.33	5.53	3.83	1.49	1.74	-0.08	0.10	N/A	-1.58	N/A	0.29	3.45	0.11	N/A	N/A	0.13	0.47	-0.49	-0.04	-0.48	-0.73	-0.65	-0.28
0.56	1.55	0.14	1.80	0.83	0.42	1.28	1.12	0.41	0.53	0.11	1.36	1.60	0.10	0.71	0.04	1.32	3.41	2.80	0.32	0.08	1.47	2.22	N/A	0.04	N/A	1.81	1.94	1.43	N/A	N/A	1.48	1.78	2.11	1.59	2.20	2.43	2.43	1.56
-1.28	-1.30	1.09	1.15	1.32	1.26	1.41	1.37	-1.26	-1.04	1.03	1.60	1.47	1.18	-1.27	1.44	-1.26	1.01	-1.27	-1.12	1.20	1.49	1.06	Х	-1.26	-1.13	1.08	1.18	1.48	1.16	-1.44	-1.23	1.07	1.07	1.58	-1.37	1.34	-1.08	-0.97
0.56	0.16	0.40	0.12	0.09	0.21	0.63	0.03	0.18	0.97	0.93	0.01	0.08	0.28	0.42	0.06	0.01	0.77	0.10	0.65	0.18	0.02	0.66	х	0.62	x	0.57	0.12	0.18	0.36	0.34	0.90	0.31	0.92	0.11	0.18			0.85
1.48	-1.29	-1.42	-1.37	-1.13	1.62	1.13	-1.09	-1.32	1.52	1.31	1.04	-0.97	1.46	-1.18	1.41	-1.55	1.14	-1.85	1.47	1.43	1.41	-1.49	x	-1.22	×	x	-0.99	1.16	x	-1.22	1.79	×	1.30	1.15	1.27	1.90	-1.15	-1.06
0.47	0.79		0.33	3 0.56	0.12	0.90	0.67	0.20	×	0.34	0.86	0.69	0.43	3 0.81	0.21	5 0.01	0.31	5 0.04	0.10	0.01	0.02	9 0.10	Х	0.84	x	х	0.45	0.33	X	0.68	×	x	0.67					0.94
0.10		-			1.44	1.27	0.14	-1.29	0.24	4 1.17	5 1.32	0.25	3 1.32	-1.23	1.43	-1.41			0.18	1.32	2 1.45		N/A	-1.24	N/A	N/A	0.10	3 1.32	N/A	-1.33	0.28	N/A	7 1.19		-0.05		3 -1.12	
1.38	0.01		1 1.26		0.18		1.23	0.03	1.28	0.14	0.28		0.14	3 0.05	0.02			0.29	1.30	0.12	0.04		N/A	4 0.02	N/A	N/A	1.09	0.16		0.11	1.51	N/A	0.12	_				0.05

SPO1681	SPO1680	SPO1679	SPO1678	SPO1677	SPO1676	SPO1675	SPO1674	SPO1673	SPO1672	SPO1671	SPO1670	SPO1669	SPO1668	SPO1667	SPO1666	SPO1665	SPO1664	SPO1663	SPO1662	SPO 1661	SPO1660	SPO1659	SPO1658	SPO1657	SPO1656	SPO1655	SPO1652	SPO1651	SPO1650	SPO1649	SPO1648	SPO1647	SPO1646	SPO1645	SPO1644	SPO1643
recG	ligA	ctrA		mnmA		lpxB		lpxA	fabZ					dxr	cdsA	Sddm	frr		pyrH	miaA																
DNA helicase RecG (EC:3.6.1)	DNA ligase (EC:6.5.1.2)	response regulator CtrA	hypothetical protein	2-thiouridylase MnmA (EC:2.1.1.61)	acetyltransferase	synthase (EC:2.4.1.182)	hypothetical protein	acyltransferase (EC:2.3.1.129)	dehydratase (EC:4.2.1)	hypothetical protein	OMP85 family outer membrane protein	hypothetical protein	zinc metalloprotease	5-phosphate reductoisomerase (EC:1.1.1.267)	cytidylyltransferase (EC:2.7.7.41)	diphosphate synthase (EC:2.5.1.31)	recycling factor	hypothetical protein	kinase (EC:2.7.4)	есни(27-заорениену футорногримсе и шизтегане (ЕС:2.5.1.75)	AraC family transcriptional regulator	binding protein	oligopeptide/dipeptide ABC transporter permease	oligopeptide/dipeptide ABC transporter permease	binding protein	hypothetical protein	transposase, truncation	hypothetical protein	hypothetical protein	invasion protein IbeA	glycine cleavage system protein T	binding protein	oligopeptide/dipeptide ABC transporter permease	oligopeptide/dipeptide ABC transporter permease	ollgopeptide/dipeptide ABC transporter ATP- binding protein	selenium-binding protein
1.12	1.01	-1.40	-1.41	-1.13	-2.19	1.35	1.22	1.38	1.40	1.27	2.12	2.41	1.49	1.40	-1.63	-1.09	1.00	1.51	1.01	1.34	-1.23	-1.60	-1.17	-1.07	-1.34	х	х	х	1.06	х	4.31	4.98	6.88	8.33	8.89	-1.41
0.19	0.99	0.02	0.04	0.48	0.27	0.14	0.95	0.01	0.22	0.55	0.00	0.20	0.91	0.04	0.01	0.22	0.03	0.11	0.14	×	х	0.02	0.89	0.06	0.19	x	х	×	0.62	х	0.03	0.22	0.04	0.04	0.03	0.33
1.12	1.20	2.22	-1.22	1.06	-1.53	1.33	1.29	1.36	1.09	1.37	1.38	1.59	-1.07	-1.09	1.37	1.13	-1.20	1.34	0.96	1.03	1.05	-1.82	-1.88	-1.63	-1.67	-1.26	-1.18	1.03	1.12	1.29	1.25	-1.02	-1.04	х	1.06	-1.78
0.62	0.55	0.09	0.44	0.77	0.46	0.34	0.59	0.02	0.78	0.14	0.02	0.18	0.92	0.56	0.16	0.34	0.37	0.42	0.71	0.88	0.86	0.08	0.12	0.02	0.02	0.43	0.47	0.83	0.37	0.40	×	0.99	0.98	×	0.81	0.27
1.12	1.11	0.41	-1.32	-0.03	-1.86	1.34	1.26	1.37	1.25	1.32	1.75	2.00	0.21	0.16	-0.13	0.02	-0.10	1.43	0.99	1.19	-0.09	-1.71	-1.53	-1.35	-1.51	N/A	N/A	N/A	1.09	N/A	2.78	1.98	2.92	N/A	4.98	-1.60
0.00	0.10	1.81	0.10	1.10	0.33	0.01	0.04	0.01	0.15	0.05	0.37	0.41	1.28	1.25	1.50	1.11	1.10	0.09	0.02	0.16	1.14	0.11	0.36	0.28	0.17	N/A	N/A	N/A	0.03	N/A	1.53	3.00	3.96	N/A	3.92	0.18
1.38	1.29	-4.64	1.63	1.73	-1.46	2.17	1.41	2.26	1.75	1.12	2.29	0.80	1.67	1.94	1.24	1.93	1.59	1.17	1.27	1.73	1.08	-3.05	-2.49	-2.19	-2.30	-2.57	-2.24	-1.05	-2.19	-1.13	-0.90	-2.15	-1.28	x	-1.43	-3.40
0.42	0.82	0.01	0.40	0.02	0.06	0.02	0.86	0.01	0.07	0.67	0.01	0.17	0.83	0.04	0.07	0.00	0.00	0.70	0.59	0.55	Х	0.00	0.25	0.00	0.03	Х	0.03	0.05	0.10	0.57	0.29	0.45	1.00	×	0.87	0.07
-1.01	-1.09	4.54	1.41	-1.13	-1.12	1.00	-1.05	-1.05	1.04	1.15	1.20	-1.09	-1.23	-1.08	-1.16	1.07	1.23	-1.24	1.28	1.18	1.21	2.29	1.40	1.25	1.62	-1.07	-0.96	1.26	-1.08	-1.14	-1.24	1.35	-1.02	x	1.21	2.09
0.92	0.63	0.02	0.17	0.12	0.65	0.64	0.76	0.19	0.86	0.38	0.20	0.46	0.75	0.34	0.14	0.63	0.36	0.54	0.14	0.54	0.71	0.01	0.38	0.09	0.05	X	0.67	0.27	0.86	х	х	0.79	1.00	×	×	0.16
0.19	0.10	-0.05	1.52	0.30	-1.29	1.59	0.18	0.61	1.40	1.14	1.75	-0.14	0.22	0.43	0.04	1.50	1.41	-0.04	1.28	1.46	1.15	-0.38	-0.55	-0.47	-0.34	-1.82	-1.60	0.11	-1.64	-1.14	-1.07	-0.40	-1.15	N/A	-0.11	-0.66
1.20	1.19	4.59	0.11	1.43	0.17	0.59	1.23	1.66	0.36	0.01	0.55	0.95	1.45	1.51	1.20	0.43	0.18	1.21	0.01	0.27	0.06	2.67	1.95	1.72	1.96	0.75	0.64	1.16	0.56	0.01	0.17	1.75	0.13	N/A	1.32	2.75
-1.21	1.45	-1.22	-1.08	1.59	1.23	1.00	1.44	1.06	1.06	1.00	1.08	1.85	1.31	-1.25	1.32	1.07	-1.19	-1.25	1.16	-1.02	-1.03	1.20	1.29	1.38	1.27	-1.09	-1.33	-0.97	1.33	-1.23	-1.42	1.07	-0.95	x	-1.14	1.45
0.65	0.16	0.33	0.45	0.03	0.02	0.59	0.37	0.78	0.90	0.85	0.41	0.08	0.71	0.26	0.10	0.77	0.02	0.50	0.60	0.92	0.98	0.25	0.40	0.08	0.17	0.94	0.24	0.50	0.15	0.50	х	0.94	0.90	x	0.64	0.26
-1.39	1.60	1.55	-1.07	-1.12	1.05	-1.35	1.31	-1.33	1.06	-1.07	-1.06	-1.43	1.22	-1.44	-1.22	-1.21	0.90	-1.71	1.89	-1.40	-1.30	3.02	1.70	1.89	1.49	1.37	-1.06	Х	2.74	х	Х	1.21	-0.99	x	х	1.97
0.25	0.60	0.21	0.70	0.31	0.29	0.08	0.40	0.07	0.75	0.16	0.30	0.13	0.88	0.09	0.05	0.12	0.12	0.14	0.01	0.40	0.70	0.01	0.17	0.02	0.03	X	x	x	x	x	х	0.90	0.74	×	×	0.16
-1.30	1.53	0.17	-1.08	0.24	1.14	-0.18	1.38	-0.14	1.06	-0.04	0.01	0.21	1.27	-1.35	0.05	-0.07	-0.14	-1.48	1.53	-1.21	-1.17	2.11	1.50	1.64	1.38	0.14	-1.20	N/A	2.04	N/A	N/A	1.14	-0.97	N/A	N/A	1.71
0.09	0.08	1.39	0.01	1.36	0.09	1.17	0.06	1.20	0.00	1.04	1.07	1.64	0.05	0.10	1.27	1.14	1.05	0.23	0.37	0.19	0.14	0.91	0.20	0.26	0.11	1.23	0.14	N/A	0.71	N/A	N/A	0.07	0.02	N/A	N/A	0.26

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### Chapter 10: Appendix

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		ureG	ureF	ureE	ureC	ureB	ureA	ureD																		rpsI	rplM					gid		hisI		
TRAP dicarboxylate transporter subunit DctM	hypothetical protein	accessory protein UreG	accessory protein UreF	accessory protein ureE	subunit alpha (EC:3.5.1.5)	subunit beta	subunit gamma (EC:3.5.1.5)	accessory protein UreD	substrate-binding protein	branched chain amino acid ABC transporter	permease branched-chain amino acid ABC transnorter	binding protein branched-chain amino acid ABC transporter	protein protein heanchad chain amino acid ABC transportar ATD	AraC family transcriptional regulator	CaiB/BaiF family protein	hypothetical protein	dihydrodipicolinate synthase	D-isomer specific 2-hydroxyacid dehydrogenase	MmgE/PrpD family protein	Asp/Glu/Hydantoin racemase	class I and II aminotransferase	hydantoinase/oxoprolinase	hydantoin utilization protein	carbohydrate kinase	hypothetical protein	ribosomal protein S9	ribosomal protein L13	GntR family transcriptional regulator	hypothetical protein	thioesterase	enoyl-CoA hydratase	(uracil-5-)-methyltransferase Gid	tRNA synthetase, class I family protein	cyclohydrolase (EC:3.5.4.19)	hypothetical protein	hypothetical protein
Х	-1.65	Х	-1.06	2.44	-2.30	1.05	-1.33	Х	x	-1.51	-3.78	-2.16	1.06	-1.44	-1.03	-1.01	1.13	1.24	3.10	2.95	1.19	1.78	1.95	1.19	1.51	-1.63	-2.40	-2.11	-2.33	1.86	2.54	1.34	-1.15	-1.12	-1.18	0.99
х	0.13	×	0.93	x	х	0.16	0.54	×	Х	0.84	0.02	0.38	1.00	0.49	1.00	0.96	0.88	0.66	0.00	0.01	0.81	0.01	0.01	0.71	0.47	0.19	0.00	0.01	0.00	0.02	0.01	0.63	0.53	0.14	0.55	0.87
1.17	1.14	-1.04	-1.16	-1.09	-1.14	-1.26	1.08	х	х	-1.04	-1.46	-1.12	-1.08	1.07	-1.40	-1.35	1.13	1.14	1.34	1.40	1.53	1.41	1.46	-1.18	1.19	1.00	-1.35	-1.13	1.06	1.13	1.83	-1.10	1.07	1.35	1.01	-1.10
0.22	0.50	0.88	0.20	0.93	0.87	0.73	0.68	×	Х	0.99	0.10	0.57	0.93	0.71	0.33	0.45	0.37	0.70	0.11	0.12	0.01	0.30	0.03	0.39	0.16	0.77	0.16	0.27	0.41	0.26	0.08	0.54	0.92	0.25	0.99	0.59
N/A	-0.26	N/A	-1.11	0.68	-1.72	-0.11	-0.13	N/A	N/A	-1.28	-2.62	-1.64	-0.01	-0.19	-1.22	-1.18	1.13	1.19	2.22	2.18	1.36	1.60	1.71	0.01	1.35	-0.32	-1.88	-1.62	-0.64	1.50	2.19	0.12	-0.04	0.12	-0.09	-0.05
N/A	1.40	N/A	0.05	1.77	0.58	1.16	1.21	N/A	N/A	0.24	1.16	0.52	1.07	1.26	0.19	0.17	0.00	0.05	0.88	0.78	0.17	0.18	0.24	1.19	0.16	1.31	0.53	0.49	1.70	0.37	0.35	1.22	1.11	1.24	1.10	1.05
-1.47	1.34	x	-1.96	-1.51	-2.79	0.76	-0.94	х	Х	-1.87	-4.70	-3.35	-1.76	-1.25	-1.21	-1.86	-1.33	-1.55	1.47	1.38	-1.41	-0.97	-0.99	1.20	1.03	13.10	8.47	1.20	2.16	-1.19	1.96	4.51	1.72	-1.19	-1.46	-1.28
0.68	0.03	×	0.05	0.79	0.49	0.00	0.57	×	х	0.81	0.00	0.22	0.74	0.94	0.98	0.22	0.52	0.36	0.03	0.03	0.13	0.02	0.38	0.86	0.72	0.02	0.00	0.52	0.01	0.51	0.02	0.10	0.10	0.26	0.39	0.03
1.13	1.16	1.08	1.10	1.21	-1.26	-1.17	-1.25	X	Х	-1.04	-1.21	1.04	1.35	1.18	1.07	1.31	1.78	1.49	1.20	-1.02	1.02	-1.12	-1.10	1.62	1.67	-2.49	-2.04	-1.43	1.05	1.36	1.68	-1.52	-1.21	-1.18	-1.19	-1.34
0.45	0.11	x	0.36	0.61	0.75	0.83	0.31	×	х	х	0.14	0.85	0.78	0.74	0.85	0.24	0.07	0.35	0.11	0.74	0.93	0.41	0.15	0.09	0.06	0.06	0.00	0.06	0.87	0.06	0.00	0.34	0.56	0.48	0.65	0.11
-0.17	1.25	N/A	-0.43	-0.15	-2.03	-0.20	-1.09	N/A	N/A	-1.46	-2.96	-1.16	-0.21	-0.04	-0.07	-0.28	0.23	-0.03	1.34	0.18	-0.20	-1.05	-1.04	1.41	1.35	5.31	3.22	-0.12	1.61	0.09	1.82	1.50	0.26	-1.19	-1.33	-1.31
1.30	0.09	N/A	1.53	1.36	0.77	0.97	0.16	N/A	N/A	0.42	1.75	2.20	1.56	1.22	1.14	1.59	1.56	1.52	0.14	1.20	1.22	0.07	0.06	0.21	0.32	7.80	5.26	1.32	0.56	1.28	0.14	3.02	1.47	0.01	0.14	0.03
-1.45	-1.33	-1.01	-1.81	-1.32	-1.68	-1.38	-1.34	-1.35	Х	-1.44	-1.42	-1.06	-1.80	-1.39	-1.57	1.04	-1.37	-1.44	-1.41	-1.19	-1.15	-1.12	-1.49	-1.06	1.14	1.05	-1.67	1.07	-1.08	1.25	1.30	-1.22	-1.17	1.25	1.75	3.43
0.13	0.13	0.67	0.04	0.45	0.38	0.46	0.09	X	×	0.89	0.03	0.84	0.52	0.49	0.47	0.75	0.13	0.27	0.09	0.06	0.35	0.27	0.06	0.78	0.17	0.98	0.00	0.53	0.09	0.04	0.01	0.66	0.64	0.12	0.25	0.01
Х	-1.70	×	-1.81	1.25	-1.27	-1.15	Х	Х	x	1.96	1.26	1.74	-1.48	-1.15	1.14	-0.91	-1.20	-1.23	-1.50	-1.26	-1.18	-1.37	-1.90	1.08	1.20	-2.69	-3.26	-1.14	1.68	1.33	1.50	-1.54	-1.27	-0.99	1.98	6.47
Х	0.08	×	0.03	х	0.78	0.93	Х	Х	Х	Х	0.12	0.09	0.69	0.97	0.68	0.58	0.53	0.51	0.15	0.17	0.19	0.13	0.06	0.92	0.24	0.06	0.00	0.55	0.05	0.04	0.04	0.37	0.47	0.93	0.26	0.00
N/A	-1.52	N/A	-1.81	-0.04	-1.48	-1.27	N/A	N/A	N/A	0.26	-0.08	0.34	-1.64	-1.27	-0.22	0.06	-1.29	-1.34	-1.46	-1.23	-1.17	-1.25	-1.70	0.01	1.17	-0.82	-2.47	-0.03	0.30	1.29	1.40	-1.38	-1.22	0.13	1.87	4.95
N//	0.18	N/A	0.00	1.29	0.20	0.12	N/A	N/A	N/A	1.70	1.34	1.40	0.16	0.12	1.36	0.98	0.09	0.11	0.05	0.04	0.02	0.13	0.21	1.07	0.03	1.87	0.80	1.11	1.38	0.04	0.10	0.16	0.05	1.12	0.12	1.52

SPO1690 SPO1691 SPO1692 SPO1693 SPO1694 SPO1695 SPO1696 SPO1697 SPO1698 SPO1698 SPO1700 SPO1700 SPO1702 SPO1702 SPO1704

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SP01710 SP01711 SP01712 SP01713 SP01714 SP01714 SP01716 SP01716 SP01717 SP01718 SP01718 SPO1705 SPO1707 SPO1708 SPO1709

SPO1761	SPO1760	SPO1759	SPO1758	SPO1757	SPO1756	SPO1755	SPO1754	SPO1753	SPO1752	SPO1751	SPO1750	SPO1749	SPO1748	SPO1747	SPO1746	SPO1745	SPO1744	SPO1743	SPO1742	SPO1741	SPO1740	SPO1739	SPO1738	SPO1737	SPO1736	SPO1735	SPO1734	SPO1733	SPO1732	SPO1729	SPO1728	SPO1727	SPO1725	SPO1724	SPO1723	SPO1722	SPO1721	SPO1720
		ribE		kpsS		kpsC	ribD	nrdR			rpoD	dnaG															hom	glpX	recJ	metA				uxuB	uxuA			
3,4-dihydroxy-2-butanone 4-phosphate synthase	hypothetical protein	riboflavin synthase subunit alpha (EC:2.5.1.9)	hypothetical protein	capsular polysaccharide export protein	polysaccharide biosynthesis/export protein	capsular polysaccharide export protein	1.0014700 otosyntuesis protein Nioz (EC.1.1.1.193	transcriptional regulator NrdR	hypothetical protein	hypothetical protein	RNA polymerase sigma factor RpoD	DNA primase (EC:2.7.7)	hypothetical protein	sarcosine oxidase subunit gamma	sarcosine oxidase subunit alpha	sarcosine oxidase subunit delta	sarcosine oxidase subunit beta	glutamate dehydrogenase (EC:1.4.1.3)	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	dehalogenase	pirin	TetR family transcriptional regulator	hypothetical protein	dehydrogenase (EC:1.1.1.3)	1,6-bisphosphatase II (EC:3.1.3.11)	exonuclease RecJ (EC:3.1)	O-succinyltransferase (EC:2.3.1.46)	esterase	polyphosphate kinase	hypothetical protein	oxidoreductase (EC:1.1.1.57)	dehydratase (EC:4.2.1.8)	GntR family transcriptional regulator	TRAP dicarboxylate transporter subunit DctP	TRAP dicarboxylate transporter subunit DctQ
1.50	1.05	1.37	х	1.10	0.87	-1.18	1.48	1.41	-1.13	1.22	1.96	1.02	1.68	5.88	5.11	6.86	7.26	-3.93	2.59	3.05	2.02	1.10	-1.17	1.18	-3.05	1.11	1.26	1.63	1.10	-1.25	1.25	-1.07	-1.09	-2.13	-1.45	-2.64	-2.67	1.01
0.62	0.89	0.06	×	0.25	0.05	0.13	0.79	0.76	0.76	0.38	0.04	0.13	0.01	0.25	0.32	0.03	0.00	0.00	0.04	0.17	0.03	0.11	0.69	0.57	0.00	0.24	0.76	0.63	0.50	0.03	0.07	0.04	0.05	0.69	0.05	0.17	0.01	0.99
1.25	1.36	1.23	1.22	1.25	-1.38	-1.59	1.25	-1.08	-1.23	0.99	-1.56	-1.82	1.19	1.76	1.62	2.18	1.76	-3.50	-1.11	1.28	1.55	1.19	-1.05	-1.12	-1.33	-1.51	1.20	1.69	1.06	-1.58	-1.21	-1.43	1.09	-1.29	-1.11	-1.06	-1.16	1.04
0.28	0.17	0.36	x	0.04	0.27	0.07	0.75	0.57	0.42	0.47	0.18	0.01	0.62	0.10	0.26	0.11	0.06	0.03	0.46	0.24	0.25	0.54	0.89	0.21	0.10	0.15	0.28	0.01	0.88	0.04	0.44	0.11	0.77	0.88	0.75	0.53	0.62	0.90
1.38	1.21	1.30	N/A	1.18	-0.25	-1.39	1.37	0.17	-1.18	1.11	0.20	-0.40	1.44	3.82	3.37	4.52	4.51	-3.72	0.74	2.17	1.79	1.15	-1.11	0.03	-2.19	-0.20	1.23	1.66	1.08	-1.42	0.02	-1.25	0.00	-1.71	-1.28	-1.85	-1.92	1.03
0.13	0.16	0.07	N/A	0.08	1.13	0.21	0.12	1.25	0.05	0.11	1.76	1.42	0.24	2.06	1.75	2.34	2.75	0.22	1.85	0.89	0.24	0.04	0.06	1.15	0.86	1.31	0.03	0.03	0.02	0.17	1.23	0.18	1.09	0.42	0.17	0.79	0.75	0.02
2.12	1.93	2.32	-1.69	-1.75	-1.42	-2.00	1.36	-1.56	1.01	1.13	1.25	1.46	1.85	-1.38	-1.55	-1.18	-1.51	-1.51	-3.42	-2.10	-2.32	-2.09	1.34	1.19	1.05	1.01	3.08	1.59	2.13	1.76	3.02	-1.39	0.93	-2.58	4.25	-3.78	-3.67	1.25
0.47	0.07	0.05	х	0.02	0.01	0.03	0.83	0.05	0.94	0.78	0.92	0.01	0.00	0.31	0.39	0.79	0.25	0.02	0.00	0.18	0.03	0.00	0.64	0.31	0.75	0.16	0.06	0.61	0.04	0.05	0.01	0.10	0.11	0.56	0.01	0.01	0.01	0.91
-1.07	-1.28	1.06	х	1.08	1.25	-1.36	-1.15	1.07	1.02	1.00	1.24	1.16	1.08	-1.10	1.35	1.26	1.06	1.09	1.03	1.04	1.33	-1.15	2.03	1.03	-1.15	1.09	-1.08	-1.13	-1.23	-1.33	-1.27	1.15	1.43	-1.24	-1.48	-1.18	0.98	1.06
0.43	0.18	0.98	×	0.52	0.12	0.02	0.72	0.96	0.99	0.70	0.19	0.40	0.85	0.71	0.26	0.51	0.90	0.94	0.91	0.96	0.09	0.04	0.19	0.84	0.13	0.83	0.09	0.11	0.45	0.01	0.04	0.26	0.11	0.89	0.05	0.14	0.87	0.90
0.53	0.33	1.69	N/A	-0.34	-0.09	-1.68	0.11	-0.25	1.02	1.07	1.25	1.31	1.47	-1.24	-0.10	0.04	-0.23	-0.21	-1.20	-0.53	-0.50	-1.62	1.69	1.11	-0.05	1.05	1.00	0.23	0.45	0.22	0.88	-0.12	1.18	-1.91	1.39	-2.48	-1.34	1.16
1.60	1.61	0.63	N/A	1.42	1.34	0.32	1.26	1.32	0.01	0.06	0.01	0.15	0.39	0.14	1.45	1.22	1.29	1.30	2.23	1.57	1.83	0.47	0.34	0.08	1.10	0.04	2.08	1.36	1.68	1.55	2.15	1.27	0.25	0.67	2.87	1.30	2.33	0.10
-1.54	1.15	-1.30	-1.46	-1.23	-1.05	-1.35	1.57	1.13	-1.12	1.24	-1.09	-1.19	1.20	-1.02	-1.04	1.17	-1.01	1.50	-2.68	-1.51	-1.74	1.22	1.02	1.32	1.01	-1.79	-1.34	1.11	1.13	-1.59	-1.56	-1.24	1.09	-1.21	-1.07	-1.24	-1.13	-1.08
	0.07		×	0.00	0.05	0.16	0.33	0.71	0.43	0.26	0.27	0.36	0.05	0.98	0.87	0.61		0.09	0.03	0.35	0.00	0.13	0.96	0.14	0.99	0.05	0.20	0.76		0.11	0.05	0.04	0.74	0.92	0.85			0.79
-1.2:	-1.28	-1.20	Х	-1.56	1.24	-1.77	1.20	1.14	1.35	1.72	1.20	0.97	1.11	1.37	1.19	-1.03	1.09	0.90	-1.86	-1.20	-1.52	1.02	-1.20	-0.96	-1.02	-1.40	-1.70	-1.50	-1.18	-1.78	-1.39	-1.13	1.11	1.66	1.39	-2.09	-1.21	-0.99
0.10		0.20	x	5 0.02	0.07	0.22	0.72	0.63		0.04		0.46	0.81	0.60	0.63	0.94		0.23		) 0.56	0.13	0.94	0.71	5 0.41	0.99	0.10	0.02	0.03		3 0.04	0.11	0.37	0.95	×				0.87
0 -1.4	7 -0.07	0 -1.25	N/A	2 -1.40	7 0.10	2 -1.56	2 1.39	3 1.14	4 0.12	4 1.48	9 0.05	6 -0.11	1 1.16	0 0.18	3 0.08	4 0.07		3 1.20	1 -2.27	6 -1.36		4 1.12	1 -0.09	1 0.18	9 -0.01	0 -1.63	2 -1.52	3 -0.20		4 -1.69	1 -1.48	7 -1.19	5 1.10	0.23	4 0.16		3 -1.17	7 -1.03
0 0.14	7 1.22	5 0.05	A N/A	0 0.17	0 1.15	6 0.21	9 0.19	4 0.01	2 1.24	8 0.24	5 1.15	1 1.08	6 0.04	8 1.20	8 1.12	7 1.10	4 1.05	0 0.30	7 0.41	6 0.16	3 0.11	2 0.10	9 1.11	8 1.14	1 1.02	3 0.16	2 0.18	0 1.31	3 1.16	9 0.10	8 0.09	9 0.06	0 0.01	3 1.44	6 1.23	7 0.43		3 0.05

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chaperone IorD		SPO1/98
sequence domain-containing protein		SPO1709
formate dehydrogenase subunit alpha twin-arginine translocation pathway signal		SPO1796
formate dehydrogenase, iron-sulfur subunit		SPO1795
EC:1.2.1.2)		SPO1794
hypothetical protein		SPO1793
excisionase		SPO1792
sulfate/tungstate ABC transporter substrate-binding protein		SPO1791
sulfate/tungstate ABC transporter permease sulfate/tungstate ABC transporter ATP-binding protein		SPO1789 SPO1790
ABC transporter substrate-binding protein		SPO1788
ABC transporter permease		SPO1787
ABC transporter permease		SPO1786
ABC transporter ATP-binding protein		SPO1785
hypothetical protein		SPO1784
phenylhydantoinase (EC:3.5.2.2)	hydA	SPO1783
hypothetical protein		SPO1782
allantoate amidohydrolase		SPO1781
TetR family transcriptional regulator		SPO1780
major facilitator superfamily transporter		SPO1779
hypothetical protein		SPO1778
dihydropyrimidine dehydrogenase (EC:1.3.1.1)		SPO1777
oxidoreductase		SPO1776
hypothetical protein		SPO1775
TRAP dicarboxylate transporter subunit DctP 3-hydroxyanthranilate 3,4-dioxygenase (EC:1.13.11.6)		SPO1773 SPO1774
TRAP dicarboxylate transporter subunit DctQ		SPO1772
TRAP dicarboxylate transporter subunit DctM		SPO1771
hypothetical protein		SPO1770
hypothetical protein		SPO1769
acetyltransferase		SPO1767
hypothetical protein		SPO1766
hypothetical protein		SPO1765
LuxR family transcriptional regulator		SPO1764
transcription antitermination factor NusB	nusB	SPO1763
6,7-dimethyl-8-ribityllumazine synthase (EC:2.5.1.9)	ribH	SPO1762

-1.58	-1.17	-1.77	-3.16	-1.75	1.35	-1.16	1.47	1.92	2.10	1.15	-2.36	-2.01	-2.03	-3.07	-1.06	-4.78	-3.03	-2.03	-1.93	1.92	х	-5.10	-7.21	1.02	0.86	-2.16	-1.51	-1.01	-1.87	-1.82	-1.06	-2.33	2.20	1.41	1.40	1.52
0.01	0.09	0.00	0.00	0.01	0.26	0.84	0.26	0.03	0.70	0.65	0.09	0.11	0.00	0.00	0.33	0.00	0.04	0.04	0.03	0.01	x	0.02	0.00	0.59	0.14	0.02	0.58	0.93	0.01	0.36	0.29	0.01	0.59	0.02	0.02	0.01
-1.50	-1.35	-2.33	-1.73	-1.32	1.08	1.13	1.25	2.15	1.46	1.34	1.07	-1.44	-1.00	1.01	1.09	-1.83	-1.58	-1.15	1.18	1.45	×	-2.83	-2.44	-1.18	-1.62	-2.71	-1.56	-1.04	-1.22	-1.15	1.00	-1.31	2.11	-1.04	1.38	1.36
0.02	0.21	0.04	0.01	0.02	0.34	0.16	0.51	0.01	0.19	0.07	0.80	0.04	0.76	0.91	0.32	0.01	0.10	0.67	0.65	0.17	×	0.05	0.01	0.17	0.25	0.02	0.07	0.67	0.43	0.16	0.94	0.30	0.12	0.59	0.27	0.70
-1.54	-1.26	-2.05	-2.45	-1.54	1.22	-0.02	1.36	2.04	1.78	1.25	-0.65	-1.73	-1.51	-1.03	0.02	-3.31	-2.31	-1.59	-0.38	1.69	N/A	-3.97	-4.83	-0.08	-0.38	-2.44	-1.54	-1.03	-1.55	-1.49	-0.03	-1.82	2.16	0.19	1.39	1.44
0.04	0.09	0.28	0.72	0.21	0.14	1.15	0.11	0.12	0.32	0.10	1.72	0.29	0.52	2.04	1.08	1.48	0.73	0.44	1.56	0.23	N/A	1.14	2.39	1.10	1.24	0.27	0.03	0.02	0.33	0.34	1.03	0.51	0.05	1.23	0.01	0.08
-10.90	-10.40	-15.70	-27.60	-13.90	-6.34	-7.51	-0.99	1.68	1.89	-1.36	-4.90	-3.35	-4.56	-7.01	-2.50	-17.40	-12.90	-9.75	-1.23	1.72	Х	-32.60	-38.20	-2.69	0.76	-4.89	-1.24	-0.98	0.93	-1.69	-1.37	-2.93	-1.11	1.30	5.02	1.98
0.00	0.00	0.00	0.00	0.00	0.02	0.01	0.76	0.01	0.76	0.07	0.01	0.01	0.00	0.00	0.00	0.00	0.00	0.01	0.91	0.01	×	0.01	0.00	0.00	0.00	0.01	0.25	0.25	0.25	0.10	0.08	0.00	0.51	0.19	0.00	0.00
5.29	3.88	4.86	6.77	6.57	6.25	6.06	1.08	-6.44	-3.65	-3.18	-1.27	1.03	-1.19	-1.83	1.60	-1.66	-1.71	-1.59	-1.65	-1.33	х	-3.74	-3.31	1.18	- 1.05	1.11	1.04	1.21	-1.02	1.10	1.59	-1.09	1.49	1.30	-1.19	-1.07
0.00	0.02	0.01	0.00	0.00	0.02	0.00	0.82	0.00	0.06	0.00	0.36	0.84	0.54	0.04	0.12	0.03	0.06	0.01	0.31	0.12	×	0.02	0.01	0.28	0.69	0.67	0.99	0.23	0.67	0.75	0.02	0.86	0.11	0.19	0.02	0.07
-2.81	-3.26	-5.42	-10.42	-3.67	-0.04	-0.73	0.05	-2.38	-0.88	-2.27	-3.09	-1.16	-2.88	-4.42	-0.45	-9.53	-7.31	-5.67	-1.44	0.20	N/A	-18.17	-20.76	-0.76	-0.15	-1.89	-0.10	0.12	-0.05	-0.30	0.11	-2.01	0.19	1.30	1.92	0.46
8.10	7.14	10.28	17.19	10.24	6.30	6.79	1.03	4.06	2.77	0.91	1.82	2.19	1.69	2.59	2.05	7.87	5.60	4.08	0.21	1.53	N/A	14.43	17.45	1.94	0.90	3.00	1.14	1.09	0.97	1.40	1.48	0.92	1.30	0.00	3.11	1.53
1.75	1.91	1.76	2.79	2.97	2.38	1.27	1.67	1.62	1.52	1.74	1.99	-1.23	1.18	1.42	-1.00	1.23	1.14	1.64	-1.06	1.13	х	1.35	1.33	1.36	-1.06	-1.28	1.01	1.16	1.33	-1.13	-1.23	1.14	1.51	-1.03	-1.29	-2.04
0.02	0.06	0.00	0.01	0.00	0.00	0.12	0.34	0.02	0.39	0.06	0.02	0.07	0.04	0.06	0.95	0.11	0.40	0.01	0.49	0.14	×	0.41	0.27	0.18	0.80	0.15	0.97	0.03	0.08	0.60	0.08	0.75	0.13	0.87	0.05	0.00
1.98	2.03	3.25	3.27	2.94	2.18	1.20	1.45	1.19	1.55	2.03	2.24	1.19	-0.98	-1.08	-0.98	1.55	1.28	1.38	-1.56	-1.44	×	1.15	1.47	1.15	1.19	1.22	1.49	-0.93	1.55	-1.20	-1.09	2.00	1.96	-1.00	-1.88	-1.88
0.01	0.04	0.02	0.00	0.00	0.06	0.18	0.30	0.60	0.15	0.01	0.00	0.21	0.17	0.91	0.51	0.01	0.22	0.19	0.20	0.14	×	0.51	0.03	0.47	0.50	0.54	0.08	0.21	0.03	0.26	0.84	0.16	0.04	0.53	0.01	0.02
1.87	1.97	2.51	3.03	2.96	2.28	1.24	1.56	1.41	1.54	1.89	2.12	-0.02	0.10	0.17	-0.99	1.39	1.21	1.51	-1.31	-0.16	N/A	1.25	1.40	1.26	0.06	-0.03	1.25	0.12	1.44	-1.17	-1.16	1.57	1.74	-1.01	-1.59	-1.96
0.12	0.06	0.75	0.24	0.02	0.10	0.04	0.11	0.22	0.02	0.15	0.13	1.21	1.08	1.25	0.01	0.16	0.07	0.13	0.25	1.29	N/A	0.10	0.07	0.11	1.13	1.25	0.24	1.05	0.11	0.04	0.07	0.43	0.23	0.02	0.30	0.08

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SPO1799

hypothetical protein

SPO1802 SPO1803 SPO1804 SPO1805 SPO1806

SPO1800 SPO1801

SPO1809

SPO1807

SPO 1808

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					livF-1																		acs	adk-2												
Gfo/Idh/MocA family oxidoreductase	sugar ABC transporter ATP-binding protein	acetamidase/formamidase	permease	oranched-chain amino acid ABC transporter permease	binding protein	substrate-binding protein branched-chain amino acid ABC transporter ATP-	binding protein branched chain amino acid ABC transporter	branched-chain amino acid ABC transporter ATP-	isochorismatase	RpiR family transcriptional regulator	glutamine synthetase	LysR family transcriptional regulator	sugar ABC transporter ATP-binding protein	sugar ABC transporter permease	sugar ABC transporter permease	sugar ABC transporter substrate-binding protein	phosphodiesterase	hypothetical protein	hypothetical protein	TRAP dicarboxylate transporter subunit DctM	TRAP dicarboxylate transporter subunit DctQ	TRAP dicarboxylate transporter subunit DctP	acetyl-coenzyme A synthetase (EC:6.2.1.1)	adenylate kinase (EC:2.7.4.3)	hypothetical protein	sodium/solute symporter family protein	hucleoudylitansierase/CBS/cyclic nucleoude- binding domain-containing protein	hypothetical protein	DNA polymerase III subunit epsilon	flavin reductase domain-containing protein	hypothetical protein	hypothetical protein	Mrp/NBP35 family protein	hypothetical protein	hypothetical protein	(Fe-S)-binding protein
-1.51	-1.22	-2.29	-1.33	-3.05	-1.77	-4.20	-3.25		-2.64	-2.44	-3.22	1.37	-2.16	-1.05	х	-1.73	1.62	1.00	1.09	-1.30	1.01	-1.45	7.34	7.69	25.20	15.40	25.30	7.42	2.54	-1.26	1.06	-1.23	1.14	1.19	-1.91	-1.35
0.31	0.46	0.00	0.12	0.30	0.01	0.00	0.00		0.02	0.00	0.32	0.18	0.01	0.97	x	0.08	0.17	0.55	0.70	0.09	0.08	0.03	0.01	0.10	0.01	0.01	0.01	0.00	0.05	0.51	0.93	0.17	0.76	0.77	0.02	0.01
-1.94	-1.40	-1.14	-1.01	-1.01	-1.10	-1.47	-1.14	1.01	A 1.07	-1.28	-1.14	-1.34	-1.26	-1.18	-1.53	-1.44	-1.22	1.01	1.00	-1.48	-1.88	-1.90	-1.61	-1.19	-1.50	-1.27	1.23	1.10	-1.15	1.03	1.05	1.03	1.04	-1.30	-1.32	1.23
0.01	0.14	0.59	0.95	0.96	0.70	0.06	0.23		0.39	0.44	0.55	0.17	0.54	0.72	0.20	0.61	0.83	0.98	0.98	0.13	0.01	0.02	0.13	0.70	0.10	0.10	0.07	0.20	0.41	0.97	0.85	0.97	0.93	0.01	0.04	0.18
-1.73	-1.31	-1.72	-1.17	-2.03	-1.44	-2.84	-2.20		-0.79	-1.86	-2.18	0.02	-1.71	-1.12	N/A	-1.59	0.20	1.00	1.04	-1.39	-0.44	-1.68	2.87	3.25	11.85	7.07	13.27	4.26	0.70	-0.12	1.06	-0.10	1.09	-0.06	-1.62	-0.06
0.21	0.09	0.58	0.16	1.02	0.34	1.37	1.06		N/A	0.58	1.04	1.36	0.45	0.06	N/A	0.15	1.42	0.01	0.05	0.09	1.45	0.23	4.48	4.44	13.35	8.34	12.04	3.16	1.85	1.15	0.01	1.13	0.05	1.25	0.30	1.29
-3.66	-2.83	-2.07	-1.14	-2.17	-1.25	-4.26	-3.52		-2.19	-2.25	-3.91	1.41	5.77	-1.30	Х	-1.63	1.60	1.10	1.37	-1.69	1.13	-4.89	2.61	3.17	7.91	5.37	11.70	3.82	2.26	-1.51	1.10	1.22	1.59	1.35	-1.54	-2.83
0.03	0.03	0.05	0.93	0.09	0.29	0.00	0.00		0.04	0.00	0.17	0.09	0.01	0.94	×	0.55	0.17	0.98	0.01	0.04	0.59	0.00	0.05	0.20	0.02	0.03	0.00	0.00	0.18	0.30	0.98	0.58	0.05	0.09	0.02	0.00
1.26	-1.10	1.31	-1.04	-1.24	-1.00	-1.02	1.16		-1.17	-1.29	-1.11	1.08	×	-1.23	x	1.05	1.23	1.03	-1.38	1.26	1.97	2.07	7.80	9.84	17.50	15.50	17.80	9.73	8.85	-1.22	1.08	-1.04	1.20	1.27	1.23	1.09
0.14	0.49	0.11	0.89	0.45	0.97	0.79	0.06	0 00	0.36	0.08	0.56	0.82	×	0.17	×	0.75	×	0.96	0.21	0.34	0.03	0.01	0.00	0.01	0.00	0.00	0.00	0.00	0.00	0.46	0.74	0.39	0.42	0.56	0.19	0.65
-1.20	-1.97	-0.38	-1.09	-1.71	-1.13	-2.64	-1.18		-1.68	-1.77	-2.51	1.25	N/A	-1.27	N/A	-0.29	1.42	1.07	0.00	-0.22	1.55	-1.41	5.21	6.51	12.71	10.44	14.75	6.78	5.56	-1.37	1.09	0.09	1.40	1.31	-0.16	-0.87
2.46	0.87	1.69	0.05	0.47	0.13	1.62	2.34		0.51	0.48	1.40	0.16	N/A	0.04	N/A	1.34	0.19	0.04	1.38	1.48	0.42	3.48	2.60	3.34	4.80	5.07	3.05	2.96	3.30	0.15	0.01	1.13	0.20	0.04	1.39	1.96
-1.29	-0.99	1.09	1.44	1.44	1.17	1.33	-1.13		A	-1.12	1.06	-1.13	-1.08	-1.14	-1.13	-1.74	-1.00	-0.99	1.14	1.40	1.69	1.69	1.85	2.05	1.35	2.29	2.37	1.19	1.22	-1.59	1.76	1.63	1.94	1.06	1.27	1.23
0.03	0.98	0.24	0.15	0.32	0.11	0.02	0.62		0.01 A	0.19	0.79	0.63	0.91	0.81	0.56	0.18	0.97	0.93	0.40	0.06	0.03	0.03	0.09	0.12	0.31	0.01	0.03	0.08	0.30	0.20	0.06	0.02	0.02	0.91	0.05	0.09
-1.26	-1.05	-0.94	-0.99	-1.38	1.31	1.47	1.36		-1.24	-1.62	2.19	1.90	1.19	1.15	x	1.25	1.78	1.62	2.12	1.21	2.05	2.58	2.32	2.13	1.75	2.29	1.66	1.37	1.25	-1.07	1.46	1.86	2.18	1.52	1.09	2.15
0.29	0.92	0.50	0.51	Х	0.13	0.10	0.18		0.58	0.06	0.05	0.40	х	0.69	x	0.69	х	0.15	0.03	0.15	0.05	0.01	0.02	0.24	0.12	0.01	0.17	0.23	0.23	0.90	0.16	0.06	0.07	0.03	0.26	0.03
-1.28	-1.02	0.08	0.23	0.03	1.24	1.40	0.12		0.03	-1.37	1.63	0.39	0.05	0.01	N/A	-0.25	0.39	0.32	1.63	1.31	1.87	2.14	2.09	2.09	1.55	2.29	2.02	1.28	1.24	-1.33	1.61	1.75	2.06	1.29	1.18	1.69
0.02	0.03	1.01	1.21	1.41	0.07	0.07	1.25		N/A	0.25	0.57	1.52	1.14	1.15	N/A	1.50	1.39	1.30	0.49	0.10	0.18	0.45	0.24	0.04	0.20	0.00	0.35	0.09	0.02	0.26	0.15	0.12	0.12	0.23	0.09	0.46

SPO1810 SPO1811 SPO1812 SPO1813 SPO1814 SPO1814 SPO1815 SPO1816 SPO1817 SPO1817 SPO1820 SPO1822 SPO1822 SPO1823 SPO1823 SPO1825 SPO1827 SPO1827

SPO1833 SPO1834 SPO1835 SPO1836 SPO1831

SPO1829 SPO1830

SPO1832

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SPO1871	SPO1870	SPO1869	SPO1868	SPO1867	SPO1866	SPO1865	SPO1864	SPO1863	SPO1862	SPO1861	SPO1860	SPO1859	SPO1858	SPO1857	SPO1856	SPO1855	SPO1854	SPO1853	SPO1852	SPO1851	SPO1850	SPO1849	SPO1848	SPO1847	SPO1846	SPO1845	SPO1844	SPO1843	SPO1842	SPO1841	SPO1840	SPO1839	SPO1838	SPO1837
	purL					tkt							rrmJ														nagA				murQ			
mechanosensitive ion channel protein MscS	(EC:6.3.5.3)	BolA family protein	glutaredoxin-like protein	hypothetical protein	hypothetical protein	transketolase (EC:2.2.1.1)	hypothetical protein	mentyter any oronace.common/mon-summ protein methyltransferase	MetF-like protein	hypothetical protein	twin-argume iransiocation partiway signat sequence domain-containing protein	Ppx/GppA phosphatase	EC:2.1.1)	hypothetical protein	ribonuclease BN	hypothetical protein	hypothetical protein	TetR family transcriptional regulator	zinc-binding dehydrogenase oxidoreductase	permease	permease branched.chain amino acid ABC transnorter	binding protein branched-chain amino acid ABC transnorter	binding protein	AMP-binding protein branched-chain amino acid ABC transporter ATP-	substrate-binding protein	molybdopterin-binding oxidoreductase branched-chain amino acid ABC transporter	N-acetyigiucosamine-o-pnospnate deacetyiase (EC:3.5.1.25)	SIS domain-containing protein	GntR family transcriptional regulator	BadF/BadG/BcrA/BcrD ATPase	N-acetylmuramic acid 6-phosphate etherase	sugar ABC transporter substrate-binding protein	sugar ABC transporter permease	sugar ABC transporter permease
0.96	1.28	3.16	2.43	1.11	0.92	1.52	-1.49	0.98	1.31	1.18	1.80	-1.07	-1.15	х	-1.19	1.41	-1.82	-2.76	-2.51	-3.27	-2.87	-2.00	-2.17	-2.39	-1.82	-1.49	1.04	-1.45	-2.05	-2.49	-2.59	-1.81	-1.40	-1.20
0.21	0.31	0.00	0.38	0.46	0.03	0.33	0.01	0.83	0.08	0.76	0.11	0.86	0.71	х	0.49	0.02	0.28	0.10	0.00	0.00	0.02	0.50	0.09	0.00	0.01	0.09	0.98	0.32	0.08	0.01	0.00	0.03	0.16	0.21
1.03	1.39	0.99	1.13	1.21	1.24	1.39	1.29	-1.07	-1.18	1.13	1.05	1.01	-1.03	Х	-1.05	-1.56	1.04	-1.71	-1.57	-2.11	-2.70	-3.22	-3.47	-4.66	-2.30	-1.53	-1.04	-1.39	-1.50	-1.40	-2.57	-2.88	-2.42	-1.57
0.92	0.22	0.84	0.44	0.42	0.45	0.57	0.38	0.43	0.16	0.63	0.77	0.99	0.79	х	0.75	0.01	0.95	0.38	0.11	0.11	0.11	0.02	0.01	0.00	0.03	0.16	0.96	0.12	0.15	0.02	0.01	0.02	0.00	0.01
1.00	1.34	2.08	1.78	1.16	1.08	1.46	-0.10	-0.04	0.07	1.16	1.43	-0.03	-1.09	N/A	-1.12	-0.08	-0.39	-2.24	-2.04	-2.69	-2.79	-2.61	-2.82	-3.53	-2.06	-1.51	0.00	-1.42	-1.78	-1.95	-2.58	-2.35	-1.91	-1.39
0.03	0.05	1.08	0.65	0.05	0.16	0.07	1.39	1.03	1.25	0.03	0.38	1.04	0.06	N/A	0.07	1.49	1.43	0.53	0.47	0.58	0.09	0.61	0.65	1.14	0.24	0.02	1.04	0.03	0.28	0.55	0.01	0.54	0.51	0.19
-1.43	6.42	1.66	1.79	-1.33	1.00	5.63	1.60	3.58	2.98	3.19	-2.55	1.28	3.07	Х	-1.62	2.22	-2.11	-7.08	-3.38	-14.10	-13.30	-11.40	-16.00	-26.50	-9.76	-4.07	-1.23	-2.58	-4.03	-3.18	-8.22	-7.70	-4.66	-3.22
0.01	0.00	0.01	0.24	0.08	0.12	0.05	0.04	0.01	0.00	0.00	0.00	0.62	0.01	х	0.01	0.01	0.34	0.05	0.00	0.00	0.03	0.14	0.00	0.00	0.01	0.01	0.89	0.25	0.02	0.00	0.00	0.00	0.00	0.01
1.87	-1.23	-1.19	1.00	1.20	1.49	-1.33	-1.38	1.23	1.77	1.39	1.37	1.04	-1.41	Х	-1.41	-1.21	1.51	1.51	1.29	1.04	-1.03	1.01	1.17	1.03	1.46	1.47	-1.11	-1.31	-1.11	-1.40	1.21	1.05	1.02	-1.39
0.01	0.08	0.14	0.57	0.41	0.05	0.46	0.03	0.26	0.02	0.21	0.29	0.98	0.02	х	0.18	0.13	0.69	0.48	0.07	0.87	0.88	0.97	0.41	0.98	0.08	0.30	0.85	0.25	0.58	0.03	0.11	0.99	0.92	0.16
0.22	2.60	0.24	1.40	-0.07	1.25	2.15	0.11	2.41	2.38	2.29	-0.59	1.16	0.83	N/A	-1.52	0.51	-0.30	-2.79	-1.05	-6.53	-7.17	-5.20	-7.42	-12.74	-4.15	-1.30	-1.17	-1.95	-2.57	-2.29	-3.51	-3.33	-1.82	-2.31
1.65	3.83	1.43	0.40	1.27	0.24	3.48	1.49	1.18	0.61	0.90	1.96	0.12	2.24	N/A	0.11	1.72	1.81	4.30	2.34	7.57	6.14	6.21	8.59	13.77	5.61	2.77	0.06	0.64	1.46	0.89	4.72	4.38	2.84	0.92
1.14	-1.14	-1.80	0.97	1.70	1.54	0.98	-1.30	-1.02	-1.62	-1.30	1.30	1.65	-1.30	Х	1.28	2.78	1.59	-1.13	1.94	1.90	1.20	1.24	-1.01	-1.56	1.50	1.20	1.50	-1.05	-1.41	1.61	-1.58	1.17	-1.18	1.12
0.56	0.33	0.01	0.48	0.01	0.07	0.90	0.06	0.39	0.01	0.05	0.51	0.32	0.20	x	0.08	0.02	0.49	0.61	0.01	0.01	0.44	0.48	0.72	0.05	0.04	0.60	0.45	0.74	0.03	0.03	0.07	0.30	0.15	0.26
1.61	-1.50	-1.32	0.99	1.18	1.66	-1.20	-2.27	-1.10	-1.22	-1.51	1.50	1.39	-1.51	Х	1.49	2.59	1.22	1.41	2.26	2.88	2.37	2.64	2.31	1.79	3.14	2.60	1.22	-1.64	1.07	-1.02	0.97	1.34	1.19	-1.12
0.03	0.10	0.02	0.27	0.16	0.01	0.55	0.02	0.11	0.25	0.15	0.13	0.55	0.22	x	0.01	0.00	0.70	0.51	0.01	0.01	0.10	0.01	0.00	0.02	0.00	0.05	0.68	0.02	0.88	0.76	0.54	0.21	0.24	0.78
1.38	-1.32	-1.56	0.98	1.44	1.60	-0.11	-1.79	-1.06	-1.42	-1.41	1.40	1.52	-1.41	N/A	1.39	2.69	1.41	0.14	2.10	2.39	1.79	1.94	0.65	0.12	2.32	1.90	1.36	-1.35	-0.17	0.30	-0.31	1.26	0.01	0.00
0.24	0.18	0.24	0.01	0.26	0.06	1.09	0.49	0.04	0.20	0.11	0.10	0.13	0.11	N/A	0.11	0.10	0.19	1.27	0.16	0.49	0.59	0.70	1.66	1.68	0.82	0.70	0.14	0.30	1.24	1.32	1.28	0.09	1.19	1.12

### Chapter 10: Appendix

SPO 1909	SPO1908	SPO 1907	SPO 1906	SPO1905	SPO 1904	SPO 1903	SPO1902	SPO1901	SPO1900	SPO1899	SPO1898	SPO1897	SPO1896	SPO1895	SPO1893	SPO1892	SPO1891	SPO1890	SPO1889	SPO1888	SPO1887	SPO1886	SPO1885	SPO1884	SPO 1883	SPO1882	SPO1881	SPO 1880	SPO1879	SPO1878	SPO1877	SPO1876	SPO1875	SPO1874	SPO1873	SPO1872
				fumC					tag	fda			ccdA			dctD-1		purQ		purS	purC							ccmF			ccmE	argC	murl			
hypothetical protein	chromate transporter	hypothetical protein	hypothetical protein	fumarate hydratase (EC:4.2.1.2)	hypothetical protein	hypothetical protein	TetR family transcriptional regulator	hypothetical protein	DNA-3-methyladenine glycosylase I (EC:3.2.2.20)	fructose-1,6-bisphosphate aldolase (EC:4.1.2.13)	cytochrome P450 family protein	hypothetical protein	cytochrome c-type biogenesis protein CcdA	SirA family protein	ribonuclease (EC:3.4.24)	C4-uicarooxyraic naiisport naiiscriptionar regulatory protein DctD	C4-dicarboxylate transport sensor protein DctB	риоэриотоозупониу дусшашалы зушазе т (EC:6.3.5.3)	alcohol dehydrogenase (EC:1.1.1.1)	subunit PurS	phosphoribosylaminoimidazolesuccinocarboxamide synthase (EC:6.3.2.6) phosphoribosylformulalycinamidina synthase	hypothetical protein	hypothetical protein	methionine synthase I (EC:2.1.1.13)	hypothetical protein	enoyl-CoA hydratase	cytochrome c biogenesis family protein	cytochrome c-type biogenesis protein CcmF	hypothetical protein	hypothetical protein	cytochrome c-type biogenesis protein CcmE	19-ассіу- запша-зпиаттут-риозрпате геоистазе (ЕС:1.2.1.38)	glutamate racemase (EC:5.1.1.3)	acyltransferase domain-containing protein	пнолерутичае тепечоли одноленненая (ЕС:1.2.7.8)	LysR family transcriptional regulator
-1.26	1.05	-1.23	-1.35	-1.45	1.02	1.08	1.07	-1.12	-1.16	-1.36	-1.39	-1.23	1.60	-1.15	-1.34	0.96	-1.28	1.60	-1.10	1.09	1.16	3.16	х	-6.01	-1.86	1.19	2.35	2.57	1.01	х	1.96	1.33	1.91	1.42	-0.99	1.08
0.26	0.44	0.12	0.00	0.08	0.95	0.90	0.89	0.07	0.67	0.06	0.00	0.08	0.43	0.32	0.03	0.03	0.63	0.08	0.52	0.47	0.53	0.01	х	0.00	0.02	0.72	0.01	0.41	0.55	×	0.00	0.18	0.01	0.11	0.99	0.32
1.73	1.08	1.33	-1.04	-1.99	-1.59	-1.52	-1.43	-1.19	-1.17	1.30	1.18	-1.06	1.37	1.41	-1.05	-1.23	-2.42	-1.18	1.42	1.57	1.42	1.45	Х	-1.93	-1.00	1.32	2.19	1.82	-1.00	1.98	1.66	1.05	1.36	1.38	-1.47	-1.01
0.16	0.44	0.06	0.60	0.05	0.15	0.04	0.10	0.43	0.42	0.10	0.42	0.31	0.39	0.01	0.28	0.03	0.02	0.18	0.08	0.10	0.16	0.03	Х	0.02	0.92	0.47	0.01	0.07	0.97	0.00	0.03	0.81	0.03	0.03	0.06	0.96
0.24	1.07	0.05	-1.20	-1.72	-0.29	-0.22	-0.18	-1.16	-1.17	-0.03	-0.11	-1.15	1.49	0.13	-1.20	-0.14	-1.85	0.21	0.16	1.33	1.29	2.31	N/A	-3.97	-1.43	1.26	2.27	2.20	0.01	N/A	1.81	1.19	1.64	1.40	-1.23	0.04
1.50	0.02	1.28	0.16	0.27	1.31	1.30	1.25	0.03	0.01	1.33	1.29	0.09	0.12	1.28	0.15	1.09	0.57	1.39	1.26	0.24	0.13	0.86	N/A	2.04	0.43	0.07	0.08	0.38	1.00	N/A	0.15	0.14	0.28	0.02	0.24	1.05
-8.67	-1.11	-1.40	-1.85	-1.62	1.03	1.18	1.93	1.35	1.60	2.16	-1.44	-1.79	1.98	1.48	1.48	2.52	-1.34	2.38	3.10	4.77	2.31	2.19	Х	2.56	-2.20	1.21	2.80	3.27	1.20	1.16	2.31	1.49	4.87	2.84	-1.54	2.27
0.01	0.48	0.01	0.00	0.13	1.00	0.35	0.09	0.24	0.10	0.01	0.07	0.02	0.11	0.09	0.03	0.00	0.16	0.03	0.01	0.00	0.03	0.01	х	0.01	0.02	0.47	0.01	0.03	0.92	Х	0.03	0.02	0.00	0.03	0.17	0.03
3.19	1.18	1.50	1.46	1.29	1.14	1.25	-1.43	-1.08	-1.31	-1.84	1.12	1.13	-1.53	1.08	-1.10	1.34	1.14	0.96	0.97	-1.32	0.97	1.16	Х	1.32	1.34	1.40	1.30	1.04	1.09	4.55	1.02	-1.27	-1.21	-1.38	-1.29	-1.33
0.02	0.11	0.06	0.02	0.56	0.73	0.08	0.04	0.28	0.06	0.01	0.34	0.31	0.26	0.73	0.11	0.08	0.50	0.54	0.16	0.01	0.15	0.45	х	0.04	0.34	0.39	0.09	0.96	0.87	0.03	0.84	0.09	0.10	0.04	0.20	0.17
-2.74	0.03	0.05	-0.20	-0.17	1.09	1.22	0.25	0.14	0.15	0.16	-0.16	-0.33	0.23	1.28	0.19	1.93	-0.10	1.67	2.04	1.73	1.64	1.68	N/A	1.94	-0.43	1.31	2.05	2.16	1.15	2.86	1.67	0.11	1.83	0.73	-1.42	0.47
5.93	1.15	1.45	1.66	1.46	0.05	0.04	1.68	1.22	1.46	2.00	1.28	1.46	1.76	0.20	1.29	0.59	1.24	0.71	1.07	3.05	0.67	0.52	N/A	0.62	1.77	0.10	0.75	1.12	0.05	1.70	0.65	1.38	3.04	2.11	0.13	1.80
-1.27	-1.28	1.46	1.18	-1.38	1.07	1.27	-1.52	1.17	-1.02	1.05	-1.02	-1.06	1.24	-1.58	1.25	1.71	-1.16	-1.87	-1.60	1.00	-1.07	1.30	Х	-1.12	1.46	-1.02	1.27	1.59	-1.13	1.42	1.62	-1.11	1.05	1.27	-0.99	-1.40
0.23	0.09	0.11	0.08	0.46	0.57	0.37	0.01	0.25	0.62	0.88	0.91	0.51	0.23	0.74	0.08	0.08	0.05	0.02	0.01	0.77	0.55	0.04	Х	0.12	0.11		0.12	0.02	0.87	0.07	0.17	0.21	0.88	0.28	0.96	0.07
1.42	-1.26	1.65	1.64	1.18	1.04	1.22	-1.61	1.02	0.98	-1.13	-1.39	-0.98	1.25	-1.98	1.21	1.88	1.05	-1.78	-2.02	-1.80	-1.20	1.27	Х	-1.11	1.29	1.19	1.10	1.25	1.39	X	1.74	1.53	-1.49	-1.64	-1.18	-1.70
0.41	0.21	0.02	0.05	0.72	0.94	0.11	0.04	0.83	0.57	0.27	0.18	0.16	0.70	0.04	0.22	0.04	0.59	0.03	0.01	0.04	0.21	0.30	x	0.06	0.04	0.56	0.58	0.16	0.17	x	0.03	0.09	0.08	0.04	0.23	0.11
0.08	-1.27	1.56	1.41	-0.10		1.25	-1.57	1.10	-0.02	-0.04	-1.21	-1.02	1.25	-1.78	1.23	1.80	-0.05	-1.83	-1.81	-0.40	-1.14		N/A	-1.12	1.38		3 1.19	1.42	0.13	N/A	1.68	0.21		-0.19	-1.09	-1.55
1.35	0.01	0.10	0.23	1.28	0.02	0.03	0.05	0.08	1.00	1.09	0.18	0.04	0.01	0.20	0.02	0.09	1.11	0.05	0.21	1.40	0.06	0.02	N/A		0.09	1.11	0.09	0.17	1.26	N/A	0.06	1.32	1.27	1.46	0.10	0.15

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SPO1946	SPO1945	SPO1944	SPO1943	SPO1942	SPO1941	SPO1940	SPO1939	SPO1938	SPO1937	SPO1936	SPO1935	SPO1934	SPO1933	SPO1932	SPO1931	SPO1930	SPO1929	SPO1928	SPO1927	SPO1926	SPO1925	SPO1923	SPO1922	SPO1921	SPO1920	SPO1919	SPO1918	SPO1917	SPO1916	SPO1915	SPO1914	SPO1913	SPO1912	SPO1911
	gmk															lpxD									trgB	trgA	cfa	phrB			acul	dmdA		
transferase	guanylate kinase (EC:2.7.4.8)	hypothetical protein	hypothetical protein	3-GeoXy-7-Phosphoneprulonate synthase (EC:2.5.1.54)	AraC family transcriptional regulator	hypothetical protein	substrate-binding protein	oranched-chain amino acid ABC transporter binding protein branched-chain amino acid ABC transporter	binding protein	branched-chain amino acid ABC transporter permease branched-chain amino acid ABC transporter ATP	branched-chain amino acid ABC transporter permease	hypothetical protein	invasion associated family protein	beta-ketoacyl synthase	acyl carrier protein	UDP-3-U-3-hydroxymyristoyi glucosamine N- acyltransferase (EC:2.3.1)	peptidoglycan binding protein	Tat pathway signal sequence domain-containing protein	phage integrase site specific recombinase	hypothetical protein	hypothetical protein	hypothetical protein	mechanosensitive ion channel protein MscS	cysteine synthase A	tellurite resistance protein	tellurite resistance protein	cyclopropane-ratty-acyt-phospholipid synthase (EC:2.1.1.79)	deoxyribodipyrimidine photolyase (EC:4.1.99.3)	class V aminotransferase	glyoxalase	acrylate utilisation protein	dimethyl sulfoniopropionate demethylase	GntR family transcriptional regulator	hypothetical protein
1.01	1.20	1.09	-2.07	1.38	-1.70	1.29	-2.15	-1.77	-1.20	-1.22	Х	-1.11	1.70	2.09	2.11	1.25	0.93	-2.66	1.80	1.71	1.94	1.58	-1.37	2.47	-1.22	1.36	-1.12	-1.33	1.95	1.29	16.30	29.60	3.24	1.30
0.94	0.75	0.59	0.23	0.07	0.00	0.39	0.02	0.05	0.64	0.74	x	0.24	0.01	0.07	0.09	0.13	0.08	0.02	0.29	0.73	0.01	0.26	0.44	0.33	0.60	0.80	0.30	0.71	0.02	0.02	0.01	0.00	0.04	0.74
1.35	1.10	1.04	2.28	1.14	-1.35	1.12	-1.84	-1.89	-1.56	-1.36	-1.10	1.05	1.35	1.53	1.67	-1.34	-1.34	1.52	-1.51	-1.38	-1.20	1.11	1.25	1.21	-1.03	1.04	-1.61	-0.99	1.06	1.98	16.40	17.00	3.86	3.87
0.18	0.73	0.92	0.03	0.72	0.25	0.60	0.01	0.05	0.11	0.13	0.59	0.87	0.08	0.19	0.01	0.27	0.23	0.19	0.25	0.73	0.23	0.74	0.14	0.50	0.84	0.87	0.23	0.95	0.46	0.01	0.00	0.00	0.05	0.12
1.18	1.15	1.07	0.11	1.26	-1.53	1.21	-2.00	-1.83	-1.38	-1.29	N/A	-0.03	1.53	1.81	1.89	-0.05	-0.21	-0.57	0.15	0.17	0.37	1.35	-0.06	1.84	-1.13	1.20	-1.37	-1.16	1.51	1.64	16.35	23.30	3.55	2.59
0.17	0.05	0.03	2.18	0.12	0.18	0.09	0.16	0.06	0.18	0.07	N/A	1.08	0.18	0.28	0.22	1.30	1.13	2.09	1.66	1.55	1.57	0.23	1.31	0.63	0.10	0.16	0.24	0.17	0.45	0.35	0.05	6.30	0.31	1.29
1.84	1.75	1.77	-1.87	3.21	1.24	-1.61	0.98	-1.03	1.13	1.29	2.15	-1.10	2.70	5.24	5.64	1.67	-2.97	2.16	1.22	1.00	1.27	2.18	1.08	3.30	2.63	1.46	-2.07	-1.64	2.97	-1.10	8.23	12.50	4.64	1.35
0.03	0.22	0.01	0.02	0.01	0.04	0.24	0.10	0.70	0.65	0.32	0.00	0.69	0.00	0.07	0.01	0.04	0.01	0.06	0.74	0.92	0.22	0.10	0.85	0.02	0.15	0.64	0.02	0.60	0.04	0.70	0.00	0.00	0.01	0.60
1.14	1.26	1.19	2.86	1.28	1.44	1.36	1.15	-1.68	-1.70	-1.56	-1.87	-1.18	1.14	-1.07	-1.32	-1.18	-1.15	1.33	-1.50	1.36	1.49	1.18	1.22	1.02	-1.33	1.08	1.44	1.06	-1.18	2.78	26.30	35.00	7.45	8.62
0.47	0.47	0.65	0.02	0.34	0.07	0.02	0.84	0.03	0.06	0.03	0.01	0.26	0.45	0.45	0.06	0.20	0.14	0.12	0.24	0.75	0.09	0.28	0.20	0.77	0.41	0.70	0.42	0.70	0.12	0.07	0.00	0.00	0.02	0.08
1.49	1.51	1.48	0.50	2.25	1.34	-0.13	1.07	-1.36	-0.29	-0.14	0.14	-1.14	1.92	2.09	2.16	0.25	-2.06	1.75	-0.14	1.18	1.38	1.68	1.15	2.16	0.65	1.27	-0.32	-0.29	0.90	0.84	17.27	23.75	6.05	4.99
0.35	0.25	0.29	2.37	0.97	0.10	1.49	0.08	0.32	1.42	1.43	2.01	0.04	0.78	3.16	3.48	1.43	0.91	0.42	1.36	0.18	0.11	0.50	0.07	1.14	1.98	0.19	1.76	1.35	2.08	1.94	9.04	11.25	1.41	3.64
-1.06	-1.68	-1.61	1.43	1.17	-1.75	1.53	1.65	-1.34	1.15	-1.17	1.38	1.11	-1.15	-1.39	1.13	-1.13	-1.90	2.24	1.16	-1.66	1.19	1.14	1.32	1.09	1.34	1.28	1.59	-1.02	-1.32	1.13	1.23	-1.23	1.14	-0.99
0.27	0.14	0.02	0.12	0.53	0.04	0.09	0.09	0.06	0.56	0.35	0.06	0.36	0.06	0.27	0.64	0.48	0.01	0.00	0.62	0.11	0.16	0.10	0.01	0.38	0.51	0.17	0.09	0.98	0.02	0.37	0.18	0.17	0.50	0.96
-1.20	-1.88	-1.45	2.04	1.16	-1.41	1.46	2.80	-1.06	-1.04	-1.17	-1.04	-1.08	-1.24	-1.80	-1.69	-1.10	0.94	1.66	-1.05	-1.40	1.26	-1.02	1.11	1.02	1.04	1.30	1.88	1.37	-1.30	-1.30	1.21	-1.60	1.33	1.26
0.18	0.23	0.00	0.07	0.94	0.09	0.29	0.01	0.54	0.68	0.24	0.82	0.20	0.05	0.03	0.07	0.15	0.59	0.03	0.82	0.56	0.40	0.48	0.34	0.26	0.99	0.22	0.20	0.21	0.06	0.61	0.49	0.08	0.56	0.34
-1.13	-1.78	-1.53	1.74	1.17	-1.58	1.50	2.23	-1.20	0.05	-1.17	0.17	0.02	-1.20	-1.60	-0.28	-1.12	-0.48	1.95	0.05	-1.53	1.23	0.06	1.22	1.06	1.19	1.29	1.74	0.18	-1.31	-0.09	1.22	-1.42	1.24	0.13
0.07	0.10	0.08	0.31	0.01	0.17	0.04	0.58	0.14	1.10	0.00	1.21	1.10	0.05	0.21	1.41	0.01	1.42	0.29	1.11	0.13	0.04	1.08	0.11	0.04	0.15	0.01	0.15	1.20	0.01	1.22	0.01	0.19	0.10	1.13

SPO 1982	SPO1981	SPO 1980	SPO 1979	SPO1978	SPO1977	SPO1976	SPO1975	SPO1974	SPO1973	SPO1972	SPO1971	SPO1970	SPO 1969	SPO 1968	SPO1967	SPO 1966	SPO 1965	SPO1964	SPO 1963	SPO1962	SPO1961	SPO 1960	SPO1959	SPO1958	SPO1957	SPO1956	SPO1955	SPO1954	SPO1953	SPO 1952	SPO1951	SPO 1950	SPO1949	SPO1948	SPO 1947
						rpsB	tsf		aroQ																	sul P	gcdH		phoB	phoU	pstB	pstA	pstC	pstS	
hypothetical protein	hypothetical protein	hypothetical protein	acyltransferase	hypothetical protein	hypothetical protein	30S ribosomal protein S2	elongation factor Ts	LuxR family transcriptional regulator	3-dehydroquinate dehydratase (EC:4.2.1.10)	nodulation protein N	enoyl-CoA hydratase (EC:4.2.1.17)	phosphoglycerate mutase	zinc-binding dehydrogenase oxidoreductase	acyl-CoA dehydrogenase	acyl-CoA dehydrogenase	oxidoreductase	onori chuin achrideananaa/radachaa OXidoredaluctase	hypothetical protein	phosphoglycerate mutase	phosphotransferase	IclR family transcriptional regulator	beta-lactamase	acetoacetyl-CoA synthetase (EC:6.2.1.1)	acyl-CoA thioesterase	oxidoreductase	sulfate permease	glutaryl-CoA dehydrogenase (EC:1.3.99.7)	LysR family transcriptional regulator	phosphate regulon transcriptional regulatory protein PhoB	phosphate transport system regulatory protein PhoU	EC:3.6.3.27)	phosphate ABC transporter permease	phosphate ABC transporter permease	prospiate ADC transporter substrate-billoring	phosphate regulon sensor histidine kinase
1.18	1.34	Х	-3.83	-2.92	-2.74	-2.23	-1.15	-2.43	-2.50	-1.09	-1.12	1.10	-1.21	1.10	1.57	1.33	1.62	1.80	1.67	1.39	-1.62	-1.15	-1.98	-0.95	1.36	Х	1.97	-3.34	2.08	2.72	0.97	1.83	2.47	5.57	-1.68
0.19	0.12	×	0.01	0.00	0.00	0.26	0.43	0.00	0.04	0.69	0.46	0.99	0.04	0.98	0.80	0.54	0.00	0.02	0.00	0.00	0.14	0.01	0.52	0.63	0.17	×	0.53	0.00	0.00	0.01	0.66	0.39	0.28	0.02	0.08
1.01	1.45	1.12	1.16	-1.15	-1.15	-1.34	1.64	-1.02	-1.04	-1.53	-1.81	1.01	-1.47	-1.45	-1.60	-1.33	-1.59	-1.27	-1.28	-1.16	-1.18	-1.24	1.11	1.22	1.07	1.16	1.18	-1.79	1.43	1.28	-1.16	-1.09	1.09	1.37	1.45
0.97	0.07	0.22	0.40	0.41	0.23	0.12	0.07	0.84	0.80	0.50	0.08	1.00	0.36	0.08	0.11	0.11	0.13	0.18	0.41	0.19	0.05	0.17	0.11	0.05	0.32	0.38	0.27	0.01	0.21	0.05	0.65	0.60	0.69	0.22	0.17
1.10	1.40	N/A	-1.34	-2.04	-1.95	-1.79	0.25	-1.73	-1.77	-1.31	-1.47	1.06	-1.34	-0.18	-0.02	0.00	0.02	0.27	0.20	0.12	-1.40	-1.20	-0.44	0.14	1.22	N/A	1.58	-2.57	1.76	2.00	-0.10	0.37	1.78	3.47	-0.12
0.09	0.05	N/A	2.50	0.88	0.80	0.45	1.40	0.71	0.73	0.22	0.35	0.05	0.13	1.28	1.59	1.33	1.61	1.54	1.48	1.28	0.22	0.05	1.55	1.08	0.15	N/A	0.40	0.78	0.33	0.72	1.06	1.46	0.69	2.10	1.57
1.64	1.40	2.26	-2.25	-2.74	-3.13	4.64	1.10	-1.21	-1.35	-2.74	-4.12	1.17	-2.90	-3.24	-2.80	-2.35	-2.33	-1.74	-2.68	-1.96	-5.48	-1.46	0.94	1.60	-1.57	-1.17	1.71	-1.46	2.18	2.36	-1.24	1.17	1.40	3.35	-1.25
0.03	0.13	0.01	0.03	0.00	0.01	0.04	0.83	0.08	0.24	0.16	0.04	0.97	0.02	0.01	0.07	0.21	0.01	0.05	0.01	0.03	0.06	0.01	0.37	0.05	0.09	0.89	0.42	0.03	0.00	0.01	0.36	0.86	0.20	0.01	0.55
1.02	1.33	-1.87	-1.67	1.52	1.34	-2.22	-3.81	-1.19	-1.32	-1.01	-1.04	1.13	-1.50	-1.05	1.01	-1.07	-1.12	1.04	-1.08	1.22	1.07	1.97	1.03	-1.19	1.11	x	1.37	1.29	1.18	1.32	1.20	1.27	1.68	1.88	-1.20
0.99	0.05	0.02	0.10	0.03	0.22	0.01	0.01	0.20	0.42	0.95	0.60	0.71	0.45	0.27	0.91	0.62	0.07	0.85	0.30	0.19	0.43	0.05	0.92	0.31	0.35	x	0.09	0.19	0.18	0.14	0.53	0.55	0.10	0.02	0.51
1.33	1.37	0.20	-1.96	-0.61	-0.90	1.21	-1.36	-1.20	-1.34	-1.88	-2.58	1.15	-2.20	-2.15	-0.90	-1.71	-1.73	-0.35	-1.88	-0.37	-2.21	0.26	0.98	0.21	-0.23	N/A	1.54	-0.09	1.68	1.84	-0.02	1.22	1.54	2.62	-1.23
0.31	0.03	2.07	0.29	2.13	2.24	3.43	2.46	0.01	0.02	0.87	1.54	0.02	0.70	1.10	1.91	0.64	0.61	1.39	0.80	1.59	3.28	1.72	0.05	1.40	1.34	N/A	0.17	1.38	0.50	0.52	1.22	0.05	0.14	0.73	0.03
-2.02	-1.20	-1.22	-1.04	1.21	1.08	-1.51	1.69	-1.23	-1.08	1.10	-1.13	1.28	1.60	1.30	1.04	1.01	-1.25	1.09	-1.41	-1.02	1.19	-1.04	-1.12	-1.24	1.08	-1.00	1.28	-1.60	1.05	-1.19	-1.34	-1.62	-1.69	-1.25	-1.02
0.01	0.08	0.26	0.96	0.08	0.05	0.20	0.01	0.30	0.66	0.86	0.47	0.39	0.03	0.03	0.90	0.98	0.02	0.60	0.12	0.61	0.08	0.72	0.61	0.32	0.43	0.84	0.19	0.01	0.51	0.24	0.21	0.08	0.07	0.03	0.92
-1.95	-1.36	-1.91	-1.35	1.52	1.19	-3.15	-2.15	-1.14	1.11	1.74	1.45	1.65	1.89	1.26	1.49	1.22	1.25	1.10	-1.20	1.09	-0.96	1.50	-1.11	-1.54	1.22	Х	1.24	-1.31	-1.32	-1.47	1.08	1.06	-1.19	-1.11	-1.16
0.03	0.10	0.08	x	0.04	0.21	0.00	0.02	0.19	0.93	0.36	0.15	0.15	0.22	0.07	0.10	0.28	0.10	0.23	0.36	0.43	0.31	0.05	0.58	0.08	0.16	×	0.29	0.51	0.08	0.02	0.92	0.96	0.12	0.47	0.81
-1.99	-1.28	-1.57	-1.20	1.37	1.14	-2.33	-0.23	-1.19	0.02	1.42	0.16	1.47	1.75	1.28	1.27	1.12	0.00	1.10	-1.31	0.04	0.12	0.23	-1.12	-1.39	1.15	N/A	1.26	-1.46	-0.14	-1.33	-0.13	-0.28	-1.44	-1.18	-1.09
0.04	0.08	0.35	0.16	0.16	0.05	0.82	1.92	0.05	1.10	0.32	1.29	0.19	0.15	0.02	0.23	0.11	1.25	0.01	0.11	1.06	1.07	1.27	0.01	0.15	0.07	N/A	0.02	0.15	1.19	0.14	1.21	1.34	0.25	0.07	0.07

### Chapter 10: Appendix

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sufB					sufC				SufS																					rplU	rpmA			obgE	proB	proA	
cysteine desulfurase	hypothetical protein	hypothetical protein	FkbM family methyltransferase	exoV domain-containing protein	FeS assembly ATPase SufC	FeS assembly protein SufD	hypothetical protein	hypothetical protein	cysteine desulfurase	hypothetical protein	LysR family transcriptional regulator	spermidine/putrescine ABC transporter permease	spermidine/putrescine ABC transporter permease	sperimume/purescine ABC transporter substrate- binding protein	spermidine/puttescine ABC transporter A1P- binding protein	class III aminotransferase	hypothetical protein	acetyltransferase	acetylpolyamine aminohydrolase	oxidoreductase, FAD-binding	endoribonuclease L-PSP	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	ABC transporter ATP-binding protein/permease	HlyD family type I secretion memorane fusion protein	hypothetical protein	hypothetical protein	50S ribosomal protein L21	50S ribosomal protein L27	amino acid transporter LysE	acetyltransferase	GTPase ObgE	gamma-glutamyl kinase (EC:2.7.2.11)	gannia-gunaniyi prospnate recuctase (EC:1.2.1.41)	hypothetical protein
-1.11	0.99	1.03	1.05	1.11	1.10	1.33	1.38	1.41	1.48	1.78	-2.12	-1.47	-2.06	-2.26	-2.11	-1.31	Х	-1.18	-1.56	-1.79	0.92	х	-1.06	-1.93	-3.06	0.98	-1.75	-1.32	1.01	-2.76	-1.28	-1.46	-2.09	-1.20	-1.20	2.59	1.28
0.89	0.39	0.54	0.99	0.35	0.93	0.64	0.07	0.64	0.31	0.05	0.00	х	0.01	0.00	0.03	х	x	0.74	0.69	0.40	0.37	х	0.79	0.08	0.00	0.88	0.29	0.00	0.16	0.00	0.02	0.49	0.02	0.08	0.02	0.06	0.77
1.04	1.02	1.02	1.20	-1.12	-1.13	-1.24	-1.32	1.06	1.40	1.13	-1.36	-1.40	-1.30	-1.40	-1.43	-0.99	-1.08	1.04	1.03	-1.37	1.10	1.06	-1.47	-1.41	-1.48	-1.18	-1.12	-1.15	-1.00	1.13	1.11	-2.09	-2.72	0.99	1.02	1.02	2.34
0.93	0.82	0.99	0.49	0.21	0.22	0.51	0.01	0.79	0.03	0.45	0.07	0.61	0.42	0.41	0.39	0.62	0.68	0.75	0.95	0.44	0.84	0.77	0.16	0.05	0.08	0.12	0.74	0.06	1.00	0.46	0.70	0.12	0.02	0.88	0.98	0.96	0.05
-0.04	1.00	1.03	1.13	-0.01	-0.01	0.05	0.03	1.24	1.44	1.46	-1.74	-1.44	-1.68	-1.83	-1.77	-1.15	N/A	-0.07	-0.27	-1.58	1.01	N/A	-1.27	-1.67	-2.27	-0.10	-1.44	-1.24	0.01	-0.82	-0.09	-1.78	-2.41	-0.10	-0.09	1.81	1.81
1.08	0.02	0.01	0.08	1.12	1.12	1.29	1.35	0.18	0.04	0.32	0.38	0.04	0.38	0.43	0.34	0.16	N/A	1.11	1.30	0.21	0.09	N/A	0.20	0.26	0.79	1.08	0.32	0.09	1.01	1.95	1.20	0.31	0.31	1.10	1.11	0.79	0.53
1.06	1.31	1.06	-1.13	1.75	1.06	1.75	1.93	2.18	2.44	1.50	-1.75	-2.58	-2.31	-3.55	-2.73	-1.43	-1.43	-1.50	-1.65	-2.42	0.82	-3.40	1.14	-2.71	-1.82	-1.34	-1.44	-1.22	1.09	4.80	5.98	1.02	-1.52	1.27	1.95	2.73	-1.48
0.76	0.07	0.49	0.85	0.07	0.98	0.34	0.02	0.04	0.01	0.10	0.04	0.19	0.05	0.03	0.03	0.02	0.55	0.43	0.77	0.04	0.40	Х	0.19	0.03	0.02	0.64	0.50	0.09	0.31	0.00	0.00	0.65	0.04	0.13	0.00	0.06	0.03
-1.41	-1.16	0.99	-1.13	-1.06	-1.22	1.04	-1.20	-1.43	-1.12	1.44	-1.15	Х	-1.02	1.01	1.10	1.65	-1.04	-0.98	-1.65	-1.11	1.53	-1.07	1.82	1.41	1.85	1.46	1.61	1.30	1.22	-1.62	-1.77	-1.07	-1.12	-1.34	-1.12	0.98	2.46
0.10	0.29	0.50	0.52	0.75	0.10	0.88	0.09	0.06	0.12	0.07	0.27	×	0.92	0.96	0.75	0.14	0.97	0.83	0.32	0.76	0.23	×	0.03	0.13	0.04	0.16	0.07	0.10	0.13	0.06	0.02	0.78	0.46	0.06	0.11	0.13	0.05
-0.18	0.08	1.02	-1.13	0.35	-0.08	1.40	0.37	0.38	0.66	1.47	-1.45	N/A	-1.67	-1.27	-0.82	0.11	-1.24	-1.24	-1.65	-1.77	1.17	-2.24	1.48	-0.65	0.02	0.06	0.09	0.04	1.16	1.59	2.11	-0.03	-1.32	-0.04	0.42	1.86	0.49
1.24	1.24	0.04	0.00	1.41	1.14	0.36	1.57	1.81	1.78	0.03	0.30	N/A	0.65	2.28	1.92	1.54	0.20	0.26	0.00	0.65	0.36	1.17	0.34	2.06	1.84	1.40	1.53	1.26	0.06	3.21	3.88	1.05	0.20	1.31	1.54	0.87	1.97
1.12	-1.18	1.14	-1.20	-1.35	-1.43	1.10	1.04	-1.25	1.29	-1.07	-1.44	-1.09	1.09	-1.07	-1.06	1.13	1.07	1.23	1.14	-1.24	1.62	-1.15	-1.14	-1.24	1.00	1.36	1.22	1.57	1.96	-1.49	1.20	1.77	1.71	1.43	-1.09	1.16	1.27
0.64	0.31	0.03	0.23	0.04	0.15	0.60	0.89	0.03	0.22	0.52	0.06	0.83	0.36	0.83	0.94	0.30	0.75	0.48	0.77	0.59	0.10	0.77	0.34	0.29	0.59	0.05	0.46	0.21	0.00	0.85	0.45	0.13	0.17	0.08	0.40	0.45	0.20
-1.12	-1.13	1.32	-1.47	-1.18	-1.34	1.00	1.25	-1.16	-1.48	1.20	-1.66	х	-1.10	1.15	1.29	Х	Х	1.64	1.18	1.47	1.48	х	-1.32	1.66	1.54	1.69	1.28	1.47	1.14	-1.39	-1.79	2.54	2.44	1.26	0.94	1.22	1.17
0.49	0.24	0.15	0.02	0.18	0.07	0.61	0.30	0.04	0.09	0.72	0.04	х	0.75	0.73	0.66	x	х	х	0.75	0.62	0.12	×	0.32	0.22	0.05	0.03	0.25	0.03	0.14	0.06	0.01	0.09	0.05	0.29	0.51	0.18	0.58
0.00	-1.16	1.23	-1.34	-1.27	-1.39	1.05	1.15	-1.21	-0.10	0.06	-1.55	N/A	-0.01	0.04	0.12	N/A	N/A	1.44	1.16	0.12	1.55	N/A	-1.23	0.21	1.27	1.53	1.25	1.52	1.55	-1.44	-0.30	2.16	2.08	1.35	-0.08	1.19	1.22
1.12	0.03	0.09	0.14	0.09	0.04	0.05	0.11	0.05	1.39	1.14	0.11	N/A	1.10	1.11	1.18	N/A	N/A	0.20	0.02	1.36	0.07	N/A	0.09	1.45	0.27	0.17	0.03	0.05	0.41	0.05	1.50	0.38	0.36	0.09	1.01	0.03	0.05

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SPO2010 SPO2013 SPO2014 SPO2016 SPO2016 SPO2017 SPO2017 SPO2017 SPO2020 SPO2020 SPO2022 SPO2022 SPO2022

SPO2007 SPO2008 SPO2009 SPO2006

SPO2001 SPO2002 SPO2003 SPO2004 SPO2005 SPO 1996 SPO 1997 SPO 1998 SPO 1999

SPO1994 SPO1995 SPO 1991 SPO 1992

SPO2000

SPO 1986 SPO 1987 SPO 1988 SPO 1989 SPO 1990

SPO1984 SPO1985 SPO 1983

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SPO2062	SPO2061	SPO2060	SPO2059	SPO2058	SPO2057	SPO2056	SPO2055	SPO2054	SPO2053	SPO2052	SPO2051	SPO2050	SPO2049	SPO2048	SPO2047	SPO2046	SPO2045	SPO2044	SPO2042	SPO2041	SPO2040	SPO2039	SPO2038	SPO2037	SPO2036	SPO2035	SPO2034	SPO2033	SPO2032	SPO2031	SPO2030	SPO2029	SPO2028	SPO2027	SPO2026	SPO2025
											gyrA			zwf-1	pgl	pgi	dmdB					guaB					recA	alaS		typA						
hypothetical protein	indigoidine synthase A family protein	PfkB family kinase	glutathione S-transferase	hypothetical protein	hypothetical protein	thiamine-phosphate pyrophosphorylase	RNA methyltransferase	cytochrome c oxidase assembly protein	thermostable carboxypeptidase (EC:3.4.17.19)	hypothetical protein	DNA gyrase subunit A (EC:5.99.1.3)	Usg	radical SAM domain-containing protein	giucose-o-phosphale 1-denydrogenase (EC:1.1.1.49)	6-phosphogluconolactonase (EC:3.1.1.31)	glucose-6-phosphate isomerase (EC:5.3.1.9)	3-methylmercaptopropionyl-CoA ligase (EC:b.2.1 )	lipoprotein	UbiE/COQ5 family methlytransferase	(Fe-S)-binding protein	CaiB/BaiF family protein	mosine-5 -monopnosphate denydrogenase (EC:1.1.1.205)	metallo-beta-lactamase	NOL1/NOP2/sun family protein	sensory box sensor histidine ktanse/response regulator (EC:2.7.3)	hypothetical protein	recombinase A	alanyl-tRNA synthetase (EC:6.1.1.7)	hypothetical protein	GTP-binding protein TypA	SMR family multidrug efflux pump	glutamine amidotransferase	HD domain-containing protein	hypothetical protein	hypothetical protein	iron-sulfur cluster assembly transcription factor IscR
1.18	-1.12	-0.98	-1.67	-1.69	1.10	1.36	-1.16	1.77	2.03	1.58	-1.33	-1.66	-1.55	-1.34	1.02	1.22	1.80	-1.52	-1.50	2.54	1.14	1.62	1.13	Х	-1.51	1.20	1.68	1.18	1.70	1.36	1.62	-1.29	0.98	1.14	1.11	1.16
0.12	0.80	0.70	0.03	0.15	0.83	0.93	0.66	0.03	0.05	0.01	0.02	0.11	0.02	0.63	0.38	0.30	0.51	0.24	0.14	0.06	Х	0.00	0.85	х	0.67	0.41	0.02	0.67	0.18	0.33	0.53	0.50	0.98	0.54	0.93	0.67
-1.68	1.03	-1.17	-1.18	-1.06	1.28	1.43	1.45	1.33	1.53	1.58	1.01	1.28	1.00	1.15	1.38	1.68	-1.03	1.34	-1.11	1.35	1.23	1.33	1.48	1.19	1.78	-1.02	1.15	-1.14	1.28	1.25	1.55	-1.42	-1.04	1.09	-1.36	1.21
0.01	0.93	0.43	0.29	0.81	0.41	0.15	0.15	0.07	0.04	0.03	0.96	0.26	0.93	0.67	0.18	0.03	0.94	0.22	0.68	0.09	0.08	0.24	0.29	0.11	0.05	0.95	0.75	0.54	0.41	0.69	0.18	0.19	0.78	0.43	0.05	0.45
-0.25	-0.05	-1.08	-1.43	-1.38	1.19	1.40	0.15	1.55	1.78	1.58	-0.16	-0.19	-0.28	-0.10	1.20	1.45	0.39	-0.09	-1.31	1.95	1.19	1.48	1.31	N/A	0.14	0.09	1.42	0.02	1.49	1.31	1.59	-1.36	-0.03	1.12	-0.13	1.19
1.43	1.08	0.09	0.25	0.32	0.09	0.03	1.31	0.22	0.25	0.00	1.17	1.47	1.27	1.25	0.18	0.23	1.42	1.43	0.19	0.60	0.05	0.15	0.18	N/A	1.65	1.11	0.27	1.16	0.21	0.06	0.04	0.06	1.01	0.02	1.24	0.03
2.13	2.55	2.70	1.37	-1.86	-1.05	3.46	3.50	2.28	3.42	-5.83	1.07	-11.30	2.48	1.53	1.31	1.44	0.98	-1.85	0.97	4.06	4.57	7.13	4.66	3.12	-2.51	1.01	-1.86	2.08	2.83	6.38	1.60	1.31	1.79	3.46	2.52	-1.99
0.02	0.05	0.01	0.17	0.02	0.59	0.56	0.02	0.00	0.01	0.00	0.74	0.02	0.00	0.22	0.24	0.06	0.84	0.01	0.97	0.03	0.00	0.00	0.13	0.00	0.02	0.74	0.01	0.02	0.08	0.00	0.61	0.55	0.81	0.01	0.01	0.07
-1.35	-1.27	1.01	1.45	1.29	1.08	-1.50	-1.64	-1.12	-1.46	2.07	-1.07	1.27	-1.09	1.57	2.07	1.53	1.60	1.25	1.57	-1.44	-1.44	-1.34	-1.37	-1.26	1.16	1.05	0.97	1.01	-1.07	-1.47	-1.19	0.99	-1.06	1.07	-1.10	-1.37
0.03	0.13	0.66	0.15	0.23	0.51	0.10	0.03	0.14	0.08	0.04	0.36	0.36	0.62	0.22	0.01	0.04	0.43	0.02	0.03	0.03	0.15	0.10	0.44	0.08	0.58	0.97	0.77	0.46	0.60	0.02	0.34	0.82	0.62	0.96	0.35	0.02
0.39	0.64	1.86	1.41	-0.29	0.02	0.98	0.93	0.58	0.98	-1.88	0.00	-5.02	0.70	1.55	1.69	1.49	1.29	-0.30	1.27	1.31	1.57	2.90	1.65	0.93	-0.68	1.03	-0.44	1.55	0.88	2.46	0.21	1.15	0.37	2.27	0.71	-1.68
1.74	1.91	0.85	0.04	1.58	1.07	2.48	2.57	1.70	2.44	3.95	1.07	6.29	1.79	0.02	0.38	0.05	0.31	1.55	0.30	2.75	3.01	4.24	3.02	2.19	1.84	0.02	1.42	0.54	1.95	3.93	1.40	0.16	1.43	1.20	1.81	0.31
1.04	-1.73	-1.98	1.13	-1.02	2.12	1.14	-1.11	-1.12	1.14	-1.31	1.00	1.10	-1.50	-1.39	-1.03	-1.04	1.46	-1.18	-1.32	-1.31	-1.00	1.23	1.04	-1.52	1.07	-1.10	1.18	1.09	1.04	-1.57	-1.03	-1.11	1.37	1.49	-1.47	-1.06
0.96	0.04	0.01	0.65	0.58	0.02	0.67	0.40	0.35	0.34	0.05	0.77	0.32	0.02	0.38	0.76	0.60	0.31	0.12	0.17	0.22	0.94	0.03	0.96	0.02	0.64	0.82	0.50	0.48	0.98	0.02	0.72	0.55	0.34	0.11	0.04	0.14
1.09	-2.33	-1.79	1.21	1.88	-1.04	-1.28	-2.13	1.14	-1.47	-1.51	-1.16	-1.47	-1.28	-2.01	-1.42	-1.18	2.45	1.67	-1.31	-2.03	-1.52	-1.18	-2.25	-2.81	-1.10	-1.50	1.56	1.44	-1.16	-1.84	-1.19	-1.31	1.04	1.19	0.99	-1.34
0.89	0.02	0.03	0.13	0.04	0.72	0.27	0.04	0.71	0.01	0.18	0.09	0.49	0.10	0.12	0.06	0.28	0.18	0.04	0.14	0.03	0.15	0.16	0.24	0.01	1.00	0.51	0.27	0.06	0.27	0.01	0.36	0.29	0.93	0.39	0.58	0.42
1.07	-2.03	-1.89	1.17	0.43	0.54	-0.07	-1.62	0.01	-0.17	-1.41	-0.08	-0.19	-1.39	-1.70	-1.23	-1.11	1.96	0.25	-1.32	-1.67	-1.26	0.03	-0.61	-2.17	-0.02	-1.30	1.37	1.27	-0.06	-1.71	-1.11	-1.21	1.21	1.34	-0.24	-1.20
0.03	0.30	0.10	0.04	1.45	1.58	1.21	0.51	1.13	1.31	0.10	1.08	1.29	0.11	0.31	0.19	0.07	0.50	1.43	0.01	0.36	0.26	1.21	1.65	0.65	1.09	0.20	0.19	0.18	1.10	0.14	0.08	0.10	0.17	0.15	1.23	0.14

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SPO2099	SPO2098	SPO2097	SPO2096	SPO2095	SPO2094	SPO2093	SPO2092	SPO2091	SPO2090	SPO2089	SPO2088	SPO2087	SPO2086	SPO2085	SPO2084	SPO2083	SPO2082	SPO2081	SPO2080	SPO2079	SPO2078	SPO2077	SPO2076	SPO2075	SPO2074	SPO2073	SPO2072	SPO2071	SPO2070	SPO2069	SPO2068	SPO2067	SPO2066	SPO2065	SPO2064	SPO2063
cycF		hpt				amt-1			ispDF		ntrB	ntrC	ntrY	ntrX		trkA	trkH-2	hfq	hflX				hem B		mfd									bdh-1		
cytochrome c-554	LysR family transcriptional regulator	nypoxantnine prosproribosyltransterase (EC:2.4.2.8)	aromatic-rich family protein	MmgE/PrpD family protein	ACT domain-containing protein	ammonium transporter	competence/damage inducible protein Cin A	phosphatidylglycerophosphatase	2-C-metnyi-D-erythritoi 4-phosphate cytidylyltransferase (EC:2.7.7.60 4.6.1.12)	tRNA-dihydrouridine synthase	nitrogen regulation protein NtrB (EC:2.7.3)	nitrogen regulation protein NtrC	nitrogen regulation protein ntrY (EC:2.7.3)	nitrogen assimilation regulatory protein NtrX	hypothetical protein	potassium transporter peripheral membrane protein	Trk system potassium uptake protein TrkH	RNA-binding protein Hfq	GTP-binding protein HflX	Na(+)/H(+) antiporter-like protein	penicillin amidase	twin-arginine translocation pathway signal sequence domain-containing protein	(EC:4.2.1.24)	hypothetical protein delta aminole vulturicacid debudratase	transcription-repair coupling factor	hypothetical protein	hypothetical protein	Bcr/CfIA subfamily drug resistance transporter	DSBA-like thioredoxin family protein	benzoate-coenzyme A ligase	DNA-binding protein	hypothetical protein	ABC transporter substrate-binding protein	D-осиа-пушохуошугас испушоденаse (EC:1.1.1.30)	patatin family phospholipase	acetyltransferase
1.03	-1.12	-1.34	-1.56	-1.03	1.21	х	2.34	1.49	-1.53	-1.17	х	1.55	-2.03	-1.91	1.32	1.02	1.26	0.91	-1.16	х	-1.08	-2.35	3.51	1.14	1.57	-1.20	1.08	-1.81	-1.95	-1.06	×	3.75	-1.92	2.03	1.58	-1.34
0.67	0.85	0.07	х	0.82	0.49	×	0.01	0.75	0.24	0.78	x	0.10	0.00	0.04	0.32	0.96	0.68	0.31	0.03	x	0.82	0.00	0.01	0.81	0.09	0.03	0.77	0.00	0.05	0.89	×	0.01	0.14	0.42	0.65	0.17
1.16	1.25	1.34	-1.01	1.01	1.25	1.89	1.36	1.35	1.33	-1.14	-1.04	-1.05	-1.48	-1.41	1.70	1.05	1.28	-1.06	-1.04	1.08	1.05	-1.90	1.45	-1.09	1.19	1.02	-1.12	-1.59	-1.43	-1.28	1.09	3.42	-1.36	1.08	1.07	-1.10
0.66	0.12	0.01	0.97	0.89	0.04	0.27	0.03	0.18	0.05	0.58	0.44	0.89	0.18	0.25	0.02	0.71	0.10	0.07	0.80	0.19	0.87	0.01	0.03	0.58	0.13	0.83	0.76	0.02	0.15	0.25	0.76	0.00	0.20	0.63	0.50	0.19
1.10	0.06	0.00	-1.29	-0.01	1.23	N/A	1.85	1.42	-0.10	-1.16	N/A	0.25	-1.76	-1.66	1.51	1.04	1.27	-0.08	-1.10	N/A	-0.02	-2.13	2.48	0.02	1.38	-0.09	-0.02	-1.70	-1.69	-1.17	N/A	3.59	-1.64	1.56	1.33	-1.22
0.06	1.19	1.34	0.28	1.02	0.02	N/A	0.49	0.07	1.43	0.02	N/A	1.30	0.28	0.25	0.19	0.02	0.01	0.98	0.06	N/A	1.07	0.23	1.03	1.12	0.19	1.11	1.10	0.11	0.26	0.11	N/A	0.17	0.28	0.48	0.25	0.12
-1.69	1.08	2.57	2.03	1.94	2.34	5.56	2.77	1.52	-1.37	1.48	1.81	2.33	-1.77	-1.52	-1.47	1.35	1.48	1.32	2.87	2.02	-1.83	1.18	3.52	3.21	4.22	-1.44	-1.54	1.84	-2.47	1.69	4.30	10.20	1.00	4.05	2.81	1.67
0.24	0.90	0.01	0.02	0.00	0.02	0.11	0.00	0.72	0.35	0.33	0.01	0.02	0.03	0.04	0.22	0.10	0.02	0.92	0.01	0.01	0.44	0.21	0.02	0.09	0.01	0.06	0.09	0.00	0.01	0.45	0.01	0.00	0.93	0.03	0.32	0.02
2.31	1.10	1.02	- 1.37	1.20	1.44	1.34	1.03	-1.31	-1.07	-1.10	-1.22	-1.16	1.47	1.48	-0.98	-1.24	1.53	1.37	1.17	-1.23	-1.60	-1.13	1.35	- 1.44	-1.51	1.61	1.56	1.18	-1.08	1.46	-1.31	1.56	-1.29	1.43	1.03	1.15
0.05	0.37	0.77	0.37	0.37	0.12	x	0.96	0.44	0.43	0.61	0.50	0.33	0.08	0.15	0.77	0.16	0.05	0.19	0.38	0.05	0.06	0.07	0.17	0.14	0.05	0.04	0.13	0.21	0.41	0.13	0.26	0.02	0.11	0.08	0.96	0.31
0.31	1.09	1.80	0.33	1.57	1.89	3.45	1.90	0.11	-1.22	0.19	0.30	0.59		-0.02		0.06	1.51	1.35	2.02	0.40	-1.72	0.03	2.44	0.89	1.36	0.09	0.01	1.51	-1.78	1.58	1.50	5.88	-0.15	2.74		
2.00	0.01	0.78		0.37	0.45	2.11	0.87	1.42	0.15	1.29	1.52	1.75	5 1.62	2 1.50	2 0.25		0.03	0.03	0.85	1.63	2 0.12	1.16	1.09	2.33	2.87	1.53		0.33	3 0.70	0.12	2.81	4.32	5 1.15	1.31	0.89	
0 2.14	1 1.71	8 1.20					7 -1.30	-		9 -1.20		5 -1.15	2 1.04	0 1.41	5 1.33	0 1.24	3 1.05	3 1.28	5 1.25	3 1.09	2 1.24	6 1.47	9 1.11		7 -1.18			3 -1.57		2 -1.63		2 1.39	5 1.12	1 1.11	9 1.39	
4 0.11	0.12	0 0.16	0.30	4 0.15	0.01	15 0.62	30 0.01	38 0.23		20 0.28		15 0.55	0.74	0.24	0.06	0.41	0.76	0.26	0.07	0.12	0.13	0.17	.1 0.50	50 0.32	18 0.20	.5 0.33	9 0.60	57 0.14	30 0.09	53 0.15	0.89	9 0.10	2 0.21	.1 0.65	9 0.28	
			-	-1.13					)4 -1.62			-1.80			)6 -1.09		<sup>7</sup> 6 -1.14				-1.37		50 1.05				50 1.38			-1.49		-		55 1.73	1.45	)6 -1.07
2.27 0.	1.38 0.	-1.15 0.			1.19 0.		-1.98 0.			-1.27 0.			1.58 0.					1.85 0.	1.04 0.			1.12 0.							1.33 0.							
0.06 2	0.16 1	0.13 0		0.42 0	0.21 1	0 X	0.00 -1	0.12 -1	0.07 -1	0.45 -1	0.59 -1	0.02 -1	0.31 1	0.12 1	0.69 0	0.39 0	0.70 -0	0.13 1	0.89 1	0.17 -0	0.30 -0	0.27 1	0.91 1	0.02 -1	0.10 -1		0.50 1	0.00 -1	0.12 0	0.06 -1	0.06 -1	0.23 1	0.29 -0	0.12 1	0.33 1	0.97 0
2.21	1.55	0.03				0.11	-1.64	-1.46	-1.42		-1.33	-1.48					-0.04	1.57	1.15	-0.21	-0.07	1.30	1.08			0.05		-1.89		-1.56						0.16
0.06	0.16	1.18	0.01	1.14	0.42	1.26	0.34	0.08	0.20	0.04	0.06	0.33	0.27	0.08	1.21	1.23	1.10	0.29	0.11	1.30	1.31	0.18	0.03	0.01	0.21	1.10	0.10	0.32	1.32	0.07	0.29	0.16	1.18	0.31	0.03	1.23

SPO2140	SPO2139	SPO2137	SPO2136	SPO2135	SPO2134	SPO2133	SPO2132	SPO2131	SPO2130	SPO2129	SPO2128	SPO2127	SPO2126	SPO2125	SPO2124	SPO2123	SPO2122	SPO2121	SPO2120	SPO2119	SPO2118	SPO2117	SPO2116	SPO2113	SPO2112	SPO2111	SPO2110	SPO2109	SPO2108	SPO2107	SPO2106	SPO2105	SPO2104	SPO2103	SPO2102	SPO2101	SPO2100
			nrdJ				aspC-3	cysS									phnA	arcB		rocF								guaA							lipA		cycG
protein	hypothetical protein	lipoprotein	(EC:1.17.4.1)	hypothetical protein	trimethylamine methyltransferase	hypothetical protein	aspartate aminotransferase (EC:2.6.1.1)	cysteinyl-tRNA synthetase (EC:6.1.1.16)	alpha-isopropylmalate synthase	hypothetical protein	transporter	DNA-3-methyladenine glycosylase II	phospholipase/carboxylesterase	acetyltransferase	RNA pseudouridylate synthase	HNH endonuclease	alkylphosphonate utilization protein PhnA	ornithine cyclodeaminase (EC:4.3.1.12)	amidinotransferase	arginase (EC:3.5.3.1)	AsnC family transcriptional regulator	DsbB family disulfide bond formation protein	DedA family protein	hypothetical protein	protein	transporter transmembrane drug/metabolite transporter family	hypothetical protein OmpP1/FadL/TodX family outer membrane	GMP synthase (EC:6.3.5.2)	trimethylamine methyltransferase	hypothetical protein	hypothetical protein	LysR family transcriptional regulator	hypothetical protein	anti-oxidant AhpCTSA family protein	lipoyl synthase	amidohydrolase	diheme cytochrome c-type
-1.03	1.37	1.16	2.54	-1.62	82.40	-2.55	х	1.23	1.12	1.07	1.64	-1.36	2.11	-1.62	x	1.02	-1.10	-4.73	-7.66	x	1.17	0.93	2.23	-1.22	1.39	Х	-2.08	1.46	3.64	-1.78	-1.35	-1.81	-1.37	1.92	2.87	2.13	-1.50
0.95	0.32	0.87	0.02	0.02	0.00	0.01	×	0.42	0.94	0.99	0.18	0.31	0.03	0.02	х	0.83	0.75	0.00	0.00	х	0.63	0.89	0.02	0.77	0.58	X	0.59	0.12	0.01	0.04	0.05	0.02	0.06	0.41	0.01	0.01	0.49
-1.13	-1.20	1.26	1.38	-1.66	15.60	-1.43	1.55	1.32	1.53	1.41	1.16	-1.03	1.76	1.85	-1.23	1.15	1.30	1.51	1.47	1.08	1.08	1.24	1.41	1.12	-1.40	-1.14	1.14	1.03	2.41	1.74	-1.08	-1.91	-1.04	1.85	1.85	1.90	-1.28
0.24	0.04	0.24	0.04	0.09	0.00	0.08	0.14	0.12	0.42	0.49	0.59	0.84	0.06	0.02	0.21	0.72	0.25	0.10	0.01	x	0.81	0.17	0.01	0.39	0.32	0.87	0.69	0.95	0.02	0.03	0.93	0.09	0.59	0.14	0.03	0.00	0.04
-1.08	0.09	1.21	1.96	-1.64	49.00	-1.99	N/A	1.28	1.33	1.24	1.40	-1.20	1.94	0.12	N/A	1.09	0.10	-1.61	-3.10	N/A	1.13	1.09	1.82	-0.05	-0.01	N/A	-0.47	1.25	3.03	-0.02	-1.22	-1.86	-1.21	1.89	2.36	2.02	-1.39
0.05	1.29	0.05	0.58	0.02	33.40	0.56	N/A	0.05	0.20	0.17	0.24	0.17	0.18	1.74	N/A	0.06	1.20	3.12	4.57	N/A	0.04	0.15	0.41	1.17	1.40	N/A	1.61	0.21	0.61	1.76	0.14	0.05	0.17	0.03	0.51	0.12	0.11
2.01	2.32	1.05	1.92	1.39	-1.64	2.16	3.18	1.87	2.13	1.38	2.12	-1.01	4.53	-3.16	1.90	-1.69	1.39	-8.00	-9.99	Х	1.03	1.07	3.98	1.36	1.57	1.28	2.27	3.38	-1.17	-6.19	-6.97	-1.86	-1.81	10.70	5.27	2.04	-2.05
0.05	0.05	0.65	0.04	0.78	0.02	0.01	0.04	0.16	0.55	0.86	0.09	0.48	0.00	0.02	0.03	0.18	0.49	0.00	0.00	х	0.96	0.95	0.01	0.23	0.26	Х	0.61	0.01	0.40	0.01	0.02	0.04	0.01	0.03	0.00	0.01	0.04
-1.07	1.14	1.30	2.05	-1.95	-1.37	-2.10	-1.10	-1.27	-1.60	-1.46	-1.01	-1.19	1.07	-1.30	-1.27	1.76	-1.43	-0.98	-1.71	х	-1.04	-1.16	-1.26	-1.16	-1.14	1.02	-1.33	-1.19	-1.51	6.68	2.41	1.32	1.26	-1.28	4.01	2.07	2.48
0.15	0.22	0.07	0.00	0.02	0.46	0.00	0.82	0.19	0.31	0.45	0.92	0.36	0.89	0.16	0.08	0.32	0.21	0.71	0.04	х	0.88	0.12	0.16	0.23	0.52	×	0.31	0.01	0.10	0.00	0.16	0.03	0.41	0.08	0.00	0.10	0.02
0.47	1.73	1.18	1.99	-0.28	-1.51	0.03	1.04	0.30	0.27	-0.04	0.56	-1.10	2.80	-2.23	0.32	0.04	-0.02	-4.49	-5.85	N/A	-0.01	-0.04	1.36	0.10	0.22	1.15	0.47	1.10	-1.34	0.25	-2.28	-0.27	-0.28	4.71	4.64	2.06	0.22
1.54	0.59	0.13	0.06	1.67	0.14	2.13	2.14	1.57	1.87	1.42	1.57	0.09	1.73	0.93	1.59	1.73	1.41	3.51	4.14	N/A	1.04	1.12	2.62	1.26	1.36	0.13	1.80	2.29	0.17	6.44	4.69	1.59	1.54	5.99	0.63	0.01	2.27
1.03	-1.02	-1.18	1.11	3.12	-1.34	3.39	1.75	-1.25	1.03	-1.03	1.16	1.03	1.34	1.11	-1.90	1.54	-1.36	-1.04	-1.69	-2.37	1.11	1.58	1.45	-1.37	1.01	-1.19	-1.30	-2.24	-1.23	-1.18	1.16	-1.37	-1.14	1.09	1.14	1.14	1.04
0.97	0.76	0.15	0.55	0.03	0.16	0.01	0.19	0.25	0.96	0.93	0.15	0.83	0.21	0.31	0.01	0.40	0.27	0.81	0.03	0.09	0.75	0.21	0.06	0.24	0.92	0.40	0.30	0.01	0.11	0.44	0.70	0.03	0.17	0.67	0.41	0.38	0.88
1.03	-1.13	-1.18	1.28	3.12	-2.12	1.53	x	-1.25	-1.21	-1.10	1.06	1.25	1.03	-2.11	-2.35	1.42	-1.44	-1.26	-1.46	х	-1.12	2.05	0.97	-1.48	2.33	-1.32	-1.10	-2.44	-1.65	х	1.68	-1.18	-1.01	-1.38	1.12	1.20	1.49
0.75	0.40	0.05	0.13	0.01	0.09	0.48	×	0.21	0.67	0.89	0.65	0.05	0.98	0.00	0.00	0.31	0.47	0.23	0.06	x	0.79	0.02	0.15	0.06	0.02	×	0.88	0.02	0.07	×	0.40	0.39	0.89	0.18	0.91	0.45	0.02
1.03	-1.08	-1.18	1.20	3.12	-1.73	2.46	N/A	-1.25	-0.09	-1.07	1.11	1.14	1.19	-0.50	-2.13	1.48	-1.40	-1.15	-1.58	N/A	-0.01	1.82	1.21	-1.43	1.67	-1.26	-1.20	-2.34	-1.44	N/A	1.42	-1.28	-1.08	-0.15	1.13	1.17	1.27
0.00	0.05	0.00	0.09	0.00	0.39	0.93	N/A	0.00	1.12	0.04	0.05	0.11	0.16	1.61	0.23	0.06	0.04	0.11	0.12	N/A	1.12	0.24	0.24	0.05	0.66	0.07	0.10	0.10	0.21	N/A	0.26	0.10	0.06	1.24	0.01	0.03	0.23

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-1.(	ABC transporter ATP-binding protein		SPO2181
-2.4	hypothetical protein		SPO2180
1.0	hypothetical protein		SPO2179
-1.6	synthetase (EC:2.7.8)	cls	SPO2178
1.1	utilization protein AcuC	acuC	SPO2177
-1.4	hypothetical protein		SPO2176
-1.0	metallo-beta-lactamase		SPO2174
-1.2	sensor histidine kinase (EC:2.7.3)		SPO2173
-1.1	LysR family transcriptional regulator		SPO2171
-2.3	metallo-beta-lactamase		SPO2170
-1.2	synthetase (EC:6.3.3.1)	purM	SPO2169
-1.0	formyltransferase (EC:2.1.2.2)	purN	SPO2168
-1.5	D (EC:3.1.26.3)	rnd	SPO2167
1.6	lipoprotein		SPO2166
-1.2	Fe-S metabolism associated family protein		SPO2165
x	hypothetical protein		SPO2164
1.5	hypothetical protein		SPO2163
1.9	hypothetical protein		SPO2162
2.2	hypothetical protein		SPO2161
3.1	corrinoid methyltransferase		SPO2160
-1.2	radical SAM domain-containing protein		SPO2159
3.2	hypothetical protein		SPO2158
4.4	synthase I (EC:2.3.3.1)	gltA	SPO2157
3.6	synthetase (EC:6.1.1.17)	gltX	SPO2156
0.9	competence protein		SPO2155
-1.1	repressor (EC:3.4.21.88)	lexA	SPO2154
1.3	cofactor biosynthesis protein A	moaA	SPO2153
1.0	cofactor biosynthesis protein MoaC	moaC	SPO2152
1.0	phosphate synthase (EC:4.1.1.48)	trpC	SPO2151
-1.0	phosphoribosyltransferase (EC:2.4.2.18)	trpD	SPO2150
1.0	synthase component II (EC:4.1.3.27)	trpG	SPO2149
-1.1	hypothetical protein		SPO2148
X	hypothetical protein		SPO2147
-1.1	synthase component I (EC:4.1.3.27)	trpE	SPO2146
1.8	hypothetical protein		SPO2145
1.0	hypothetical protein		SPO2144
1.1	xanthine-guanine phosphoribosyltransterase (EC:2.4.2.22)	gpt	SPO2143
0.9	enoyl-ACP reductase (EC:1.3.1.10)	fabI-2	SPO2142
0.9	pyridoxamine 5"-phosphate oxidase		SPO2141

-1.05	-2.42	1.03	-1.63	1.10	-1.47	-1.02	-1.24	-1.15	-2.36	-1.26	-1.09	-1.57	1.67	-1.21	х	1.54	1.96	2.25	3.14	-1.23	3.28	4.40	3.64	0.98	-1.13	1.30	1.03	1.04	-1.08	1.06	-1.14	х	-1.17	1.86	1.06	1.11	0.93	0.94
0.65	0.24	0.90	0.55	0.99	0.04	0.64	0.25	0.82	0.00	0.03	0.49	0.38	0.02	0.06	×	×	0.30	0.39	0.10	x	0.11	0.10	0.33	0.51	0.77	0.04	0.89	0.44	0.18	0.72	0.17	×	0.10	0.25	0.40	0.96	0.45	0.05
1.57	-2.17	1.22	1.02	1.18	-1.15	1.09	-1.22	-1.69	-2.50	1.49	1.50	-1.42	1.38	1.16	-1.17	1.11	1.13	-1.60	1.90	-1.18	3.68	1.43	1.04	1.10	1.04	1.11	1.36	-1.20	1.31	1.07	-1.12	1.13	0.98	1.06	1.03	-1.07	-1.28	1.04
0.05	0.01	0.47	0.69	0.79	0.04	0.35	0.56	0.01	0.06	0.07	0.12	0.14	0.11	0.32	0.30	0.59	0.45	0.02	0.10	0.48	0.00	0.26	0.98	0.72	0.89	0.49	0.40	0.13	0.10	0.87	0.44	0.01	0.77	0.86	0.91	0.65	0.14	0.93
0.26	-2.30	1.13	-0.31	1.14	-1.31	0.04	-1.23	-1.42	-2.43	0.12	0.21	-1.50	1.53	-0.03	N/A	1.33	1.55	0.33	2.52	-1.21	3.48	2.92	2.34	1.04	-0.04	1.21	1.20	-0.08	0.12	1.07	-1.13	N/A	-0.09	1.46	1.05	0.02	-0.17	0.99
1.31	0.13	0.10	1.33	0.04	0.16	1.06	0.01	0.27	0.07	1.38	1.30	0.08	0.15	1.19	N/A	0.21	0.42	1.93	0.62	0.03	0.20	1.49	1.30	0.06	1.09	0.10	0.17	1.12	1.20	0.01	0.01	N/A	1.08	0.40	0.02	1.09	1.11	0.05
1.32	-1.42	1.42	-1.27	1.90	-1.41	1.04	1.07	2.47	1.02	4.86	2.63	1.26	1.84	1.60	-0.88	-0.97	-1.67	1.44	4.27	1.37	-1.08	3.14	5.24	-1.21	-2.43	2.64	0.88	1.29	1.33	1.78	-1.27	3.42	2.14	3.38	3.62	3.28	2.77	-1.24
0.18	0.09	0.23	0.94	0.76	0.20	0.39	0.43	0.01	0.68	0.00	0.01	0.84	0.00	0.02	0.12	0.30	0.01	0.22	0.05	0.01	0.95	0.04	0.20	0.34	0.34	0.03	0.53	0.07	0.27	0.01	0.00	0.01	0.02	0.05	0.00	0.04	0.00	0.02
1.34	1.14	1.50	-0.97	1.21	1.25	1.01	1.04	-1.23	-2.15	-1.20	-1.35	-1.33	-1.45	-1.17	1.33	1.19	1.07	0.99	1.50	-1.06	1.70	1.27	-1.14	1.34	-1.63	-1.16	1.01	1.10	1.05	1.09	1.04	-1.41	-1.15	-1.05	1.05	1.56	1.04	1.40
0.08	0.72	0.03	0.75	0.40	0.13	0.87	0.57	0.39	0.02	0.22	0.26	0.12	0.04	0.04	0.06	0.16	0.61	0.56	0.08	0.94	0.01	0.19	0.84	0.43	0.09	0.21	0.86	0.70	0.99	0.66	0.91	0.35	0.24	0.41	0.92	0.03	0.84	0.08
1.33	-0.14	1.46	-1.12	1.56	-0.08	1.03	1.06	0.62	-0.57	1.83	0.64	-0.04	0.20	0.22	0.23	0.11	-0.30	1.22	2.89	0.16	0.31	2.21	2.05	0.07	-2.03	0.74	0.94	1.20	1.19	1.44	-0.12	1.01	0.50	1.17	2.34	2.42	1.91	0.08
0.01	1.28	0.04	0.15	0.35	1.33	0.02	0.02	1.85	1.59	3.03	1.99	1.30	1.65	1.39	1.10	1.08	1.37	0.22	1.39	1.22	1.39	0.94	3.19	1.28	0.40	1.90	0.07	0.10	0.14	0.35	1.16	2.42	1.65	2.22	1.29	0.86	0.87	1.32
1.14	1.02	1.77	-1.12	1.54	1.30	-1.20	-1.18	1.03	2.30	-1.77	-1.59	-1.18	1.35	1.41	-1.48	-1.27	-1.03	-1.62	-1.41	1.36	-1.08	1.20	-1.16	1.28	1.32	-1.16	1.33	-1.77	-1.06	-1.33	1.22	-1.30	-1.45	-1.27	1.01	1.02	-1.38	1.38
0.19	0.96	0.02	0.63	0.41	0.02	0.12	0.26	0.70	0.04	0.04	0.18	0.24	0.05	0.04	0.02	0.23	0.84	0.05	0.10	0.29	0.62	0.40	0.78	0.23	0.48	0.59	0.20	0.02	0.15	0.06	0.33	0.17	0.13	0.04	0.94	0.60	0.22	0.03
1.48	1.54	1.99	-1.56	1.24	1.13	-1.45	-0.99	1.42	6.78	-2.08	-2.15	-1.42	-1.06	1.09	х	1.12	-1.00	-1.51	-1.94	1.61	-1.59	1.39	-1.65	1.49	2.03	-2.32	1.26	-1.75	-1.24	-1.31	1.26	-1.62	-1.40	0.94	-1.23	1.08	-1.21	1.70
0.16	0.30	0.09	0.10	0.35	0.13	0.13	0.79	0.04	0.00	0.01	0.05	0.04	0.58	0.87	×	0.49	0.75	0.05	0.02	0.06	0.16	0.17	0.57	0.20	0.06	0.03	0.66	0.03	0.26	0.07	0.38	0.12	0.31	0.05	0.23	0.27	0.31	0.01
1.31	1.28	1.88	-1.34	1.39	1.22	-1.33	-1.08	1.23	4.54	-1.93	-1.87	-1.30	0.15	1.25	N/A	-0.08	-1.01	-1.57	-1.68	1.49	-1.34	1.30	-1.41	1.39	1.68	-1.74	1.30	-1.76	-1.15	-1.32	1.24	-1.46	-1.43	-0.17	-0.11	1.05	-1.30	1.54
0.17	0.26	0.11	0.22	0.15	0.09	0.13	0.10	0.19	2.24	0.16	0.28	0.12	1.21	0.16	N/A	1.20	0.02	0.06	0.27	0.13	0.26	0.10	0.25	0.11	0.36	0.58	0.04	0.01	0.09	0.01	0.02	0.16	0.03	1.10	1.12	0.03	0.09	0.16

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		uvrA				betC-2	mmsB-1										mmsA			coaD		gap-3				hutF	hutU	hutH	hutl	hutC								
hypothetical protein	rhodanese-like domain-containing protein	ABC subunit A	hypothetical protein	DNA-binding protein	glyoxalase	sulfatase (EC:3.1.6.6)	dehydrogenase (EC:1.1.1.31)	enoyl-CoA hydratase	acyl-CoA dehydrogenase (EC:1.3.99)	BNR/Asp-box repeat-containing protein	carboxylate-amine ligase	hypothetical protein	hypothetical protein	hypothetical protein	ErfK/YbiS/YcfS/YnhG family protein	hypothetical protein	dehydrogenase (EC:1.2.1.27)	LysR family transcriptional regulator	hypothetical protein	adenylyltransferase (EC:2.7.7.3)	hypothetical protein	dehydrogenase, type I (EC:1.2.1)	hypothetical protein	diaminopropionate ammonia-lyase (EC:4.3.1.15)	amidohydrolase	deiminase (EC:3.5.3.13)	hydratase (EC:4.2.1.49)	ammonia-lyase (EC:4.3.1.3)	(EC:3.5.2.7)	utilization repressor	AsnC family transcriptional regulator	acetyltransferase	protein	TRAP transporter, 4TM/12TM fusion protein TRAP transporter solute recentor TAXI family	universal stress protein family protein	hypothetical protein	hypothetical protein	permease
1.16	1.01	1.10	-1.42	-1.05	1.66	2.12	1.02	1.83	1.10	0.91	-1.18	1.13	1.41	2.58	2.40	2.88	10.30	1.27	1.17	1.87	-1.09	1.06	1.79	-1.10	-1.12	-1.10	1.34	1.35	1.13	-1.30	-2.01	-1.35	-2.37	-2.60	-3.84	-1.85	1.60	1.63
0.91	0.96	0.95	0.55	0.75	0.02	0.21	0.59	0.59	0.99	0.15	0.64	0.43	0.19	0.30	0.00	0.01	0.00	0.31	0.92	0.03	0.70	0.96	0.16	0.91	0.42	0.73	0.04	0.88	0.72	0.78	0.00	0.66	0.00	0.00	0.00	0.02	0.83	x
1.30	-1.06	-1.21	-1.31	-1.07	-1.54	-1.26	-1.16	1.12	-1.02	2.62	-1.11	1.03	-1.30	1.01	1.12	1.19	2.39	1.10	1.38	2.14	1.59	-1.15	1.17	1.16	-0.99	1.24	1.00	-1.11	1.07	-1.05	1.10	1.18	-2.23	-1.98	-1.43	-1.13	1.53	2.05
0.20	0.92	0.51	0.38	0.53	0.02	0.55	0.59	0.73	0.90	0.00	0.62	0.90	0.40	0.98	0.37	0.29	0.04	0.32	0.21	0.02	0.05	0.52	0.51	0.77	0.34	0.18	0.97	0.50	0.17	0.93	0.40	0.68	0.02	0.00	0.13	0.09	0.31	0.12
1.23	-0.03	-0.05	-1.37	-1.06	0.06	0.43	-0.07	1.48	0.04	1.76	-1.15	1.08	0.05	1.80	1.76	2.04	6.35	1.19	1.28	2.01	0.25	-0.04	1.48	0.03	-1.06	0.07	1.17	0.12	1.10	-1.18	-0.46	-0.09	-2.30	-2.29	-2.64	-1.49	1.57	1.84
														_	_	~		_	_	_			-		~		_		_									~
0.07	1.04	1.16	0.05	0.01	1.60	1.69	1.09	0.36	1.06	0.86	0.03	0.05	1.36	0.79	0.64	0.85	3.96	0.09	0.11	0.14	1.34	1.11	0.31	1.13	0.06	1.17	0.17	1.23	0.03	0.13	1.56	1.27	0.07	0.31	1.21	0.36	0.04	0.21
0.07 -1.41	1.04 2.47	1.16 1.78	0.05 -1.02	0.01 -1.96	1.60 -1.80	1.69 2.28	1.09 -1.64	0.36 -1.04	1.06 -1.90	0.86 -2.32	0.03 -1.46	0.05 3.27	1.36 -2.09	0.79 -1.22	0.64 1.81	0.85 1.94	3.96 1.96	0.09 -1.15	0.11 -1.86	0.14 1.39	·	1.11 1.10	0.31 1.79	1.13 1.61	0.06 1.24		0.17 -2.55	1.23 0.92	0.03 -2.65	0.13 -2.01	1.56 1.19	1.27 1.13	0.07 1.69	0.31 1.14	1.21 -1.46	0.36 -1.29	0.04 1.85	).21 3.06
	2.47																				·					-1.69												
-1.41	- 2.47 0.54 -	1.78	-1.02 0.92 -	-1.96	-1.80	2.28	-1.64	-1.04	-1.90	-2.32	-1.46	3.27	-2.09	-1.22	1.81	1.94	1.96	-1.15	-1.86	1.39	2.20 0.03	1.10 0.88	1.79 0.10	1.61 0.38	1.24	-1.69 0.40 .	-2.55	0.92	-2.65 0.07 -	-2.01	1.19	1.13	1.69	1.14	-1.46	-1.29	1.85	3.06
-1.41 0.82	2.47 0.54 -1.24	1.78 0.25	-1.02 0.92	-1.96 0.20	-1.80 0.10	2.28 0.11	-1.64 0.03	-1.04 0.97	-1.90 0.08	-2.32 0.00	-1.46 0.50	3.27 0.00 -	-2.09 0.02	-1.22 0.82 -	1.81 0.02	1.94 0.02	1.96 0.02	-1.15 0.64	-1.86 0.01	1.39 0.22 1.83	2.20 0.03 1.06	1.10 0.88 -8.43	1.79 0.10	1.61 0.38	1.24 0.08 .	-1.69 0.40 -1.51	-2.55 0.02 -1.59	0.92 0.86	-2.65 0.07 -1.07	-2.01 0.61 -	1.19 0.15	1.13 0.66 .	1.69 0.02	1.14 0.16 -	-1.46 0.00	-1.29 0.26	1.85 0.76	3.06 0.09
-1.41 0.82 1.51	. 2.47 0.54 -1.24 0.64	1.78 0.25 1.06	-1.02 0.92 -1.02	-1.96 0.20 1.17	-1.80 0.10 2.14	2.28 0.11 1.09	<b>-1.64 0.03</b> 1.21	-1.04 0.97 3.53	-1.90 0.08 4.62	-2.32 0.00 2.83	-1.46 0.50 1.61	3.27 0.00 -1.22	-2.09 0.02 1.38	-1.22 0.82 -1.17	1.81 0.02 1.45	1.94 0.02 1.46	1.96 0.02 6.98	-1.15 0.64 1.46	-1.86 0.01 1.71	1.39 0.22 1.83	2.20 0.03 1.06 0.76	1.10 0.88 -8.43 0.03 -	1.79 0.10 1.01	1.61 0.38 -1.21	1.24 0.08 -1.13	-1.69 0.40 -1.51 0.21	-2.55 0.02 -1.59 0.04	0.92 0.86 -1.25 0.43	-2.65 0.07 -1.07 0.60	-2.01 0.61 -1.26 0.36	1.19 0.15 -1.20	1.13 0.66 -0.98	1.69 0.02 1.34	1.14 0.16 -1.12	-1.46 0.00 1.30 0.02	-1.29 0.26 1.37	1.85 0.76 1.19 0.59	3.06 0.09 1.83
-1.41 0.82 1.51 0.07	2.47 0.54 -1.24 0.64 0.62	1.78 0.25 1.06 0.91	-1.02 0.92 -1.02 0.92 -	-1.96 0.20 1.17 0.58	-1.80 0.10 2.14 0.01	2.28 0.11 1.09 0.60	-1.64 0.03 1.21 0.74	-1.04 0.97 3.53 0.14	<b>-1.90</b> 0.08 4.62 0.00	-2.32 0.00 2.83 0.01	-1.46 0.50 1.61 0.27	3.27 0.00 -1.22 0.32	-2.09 0.02 1.38 0.46	-1.22 0.82 -1.17 0.79 .	1.81         0.02         1.45         0.13	1.94         0.02         1.46         0.04	1.96         0.02         6.98         0.01	-1.15 0.64 1.46 0.02	-1.86 0.01 1.71 0.02	1.39         0.22         1.83         0.03	2.20 0.03 1.06 0.76 1.63	1.10 0.88 -8.43 0.03 -3.67	1.79 0.10 1.01 0.73	1.61 0.38 -1.21 0.74	1.24 0.08 -1.13 0.40	-1.69 0.40 -1.51 0.21 -1.60	-2.55 0.02 -1.59 0.04	0.92 0.86 -1.25 0.43	-2.65 0.07 -1.07 0.60 -1.86	-2.01 0.61 -1.26 0.36	1.19 0.15 -1.20 0.19	1.13 0.66 -0.98 0.95	1.69 0.02 1.34 0.08	1.14 0.16 -1.12 0.20	-1.46 0.00 1.30 0.02	-1.29 0.26 1.37 0.08	1.85 0.76 1.19 0.59 1.52	3.06 0.09 1.83 0.03
-1.41 0.82 1.51 0.07 0.05	2.47 0.54 -1.24 0.64 0.62	1.78 0.25 1.06 0.91 1.42	-1.02 0.92 -1.02 0.92 -1.02	-1.96 0.20 1.17 0.58 -0.40	-1.80 0.10 2.14 0.01 0.17	2.28 0.11 1.09 0.60 1.69	-1.64 0.03 1.21 0.74 -0.22	-1.04 0.97 3.53 0.14 1.25	<b>-1.90</b> 0.08 <b>4.62 0.00 1.36</b>	-2.32 0.00 2.83 0.01 0.26	-1.46 0.50 1.61 0.27 0.08	3.27 0.00 -1.22 0.32 1.03	-2.09 0.02 1.38 0.46 -0.36	-1.22 0.82 -1.17 0.79 -1.20	1.81         0.02         1.45         0.13         1.63	1.94         0.02         1.46         0.04         1.70	1.96         0.02         6.98         0.01         4.47	-1.15 0.64 1.46 0.02 0.16	-1.86 0.01 1.71 0.02 -0.08	1.39         0.22         1.83         0.03         1.61	2.20 0.03 1.06 0.76 1.63	1.10 0.88 -8.43 0.03 -3.67	1.79 0.10 1.01 0.73 1.40	1.61 0.38 -1.21 0.74 0.20	1.24 0.08 -1.13 0.40 0.06	-1.69 0.40 -1.51 0.21 -1.60 0.09	-2.55 0.02 -1.59 0.04 -2.07	0.92 0.86 -1.25 0.43 -0.17	-2.65 0.07 -1.07 0.60 -1.86 0.79	-2.01 0.61 -1.26 0.36 -1.64	1.19 0.15 -1.20 0.19 -0.01	1.13 0.66 -0.98 0.95 0.07	1.69 0.02 1.34 0.08 1.52	1.14 0.16 -1.12 0.20 0.01	-1.46 0.00 1.30 0.02 -0.08	-1.29 0.26 1.37 0.08 0.04	1.85 0.76 1.19 0.59 1.52	3.06 0.09 1.83 0.03 2.45
-1.41 0.82 1.51 0.07 0.05 1.46 1	2.47 0.54 -1.24 0.64 0.62 1.86 1.26	1.78 0.25 1.06 0.91 1.42 0.36	-1.02 0.92 -1.02 0.92 -1.02 0.00	-1.96 0.20 1.17 0.58 -0.40 1.57 -	-1.80 0.10 2.14 0.01 0.17 1.97	2.28 0.11 1.09 0.60 1.69 0.60	<b>-1.64 0.03 1.21 0.74 -0.22 1.43</b>	-1.04 0.97 3.53 0.14 1.25 2.29	-1.90 0.08 4.62 0.00 1.36 3.26	-2.32 0.00 2.83 0.01 0.26 2.58	-1.46 0.50 1.61 0.27 0.08 1.54	3.27 0.00 -1.22 0.32 1.03 2.25 -	-2.09 0.02 1.38 0.46 -0.36 1.74	-1.22 0.82 -1.17 0.79 -1.20 0.03	1.81         0.02         1.45         0.13         1.63         0.18         .	1.94         0.02         1.46         0.04         1.70         0.24	1.96         0.02         6.98         0.01         4.47         2.51	<b>-1.15</b> 0.64 1.46 0.02 0.16 1.31	-1.86 0.01 1.71 0.02 -0.08 1.79	1.39         0.22         1.83         0.03         1.61         0.22         1.16	2.20 0.03 1.06 0.76 1.63 0.57 1.62	1.10         0.88         -8.43         0.03         -3.67         4.77         1.90	1.79 0.10 1.01 0.73 1.40 0.39	1.61         0.38         -1.21         0.74         0.20         1.41	1.24 0.08 -1.13 0.40 0.06 1.19	-1.69 0.40 -1.51 0.21 -1.60 0.09 1.17	-2.55 0.02 -1.59 0.04 -2.07 0.48	0.92 0.86 -1.25 0.43 -0.17 1.08	-2.65 0.07 -1.07 0.60 -1.86 0.79	-2.01 0.61 -1.26 0.36 -1.64 0.38	<b>1.19</b> 0.15 -1.20 0.19 -0.01 1.20	1.13 0.66 -0.98 0.95 0.07 1.06	1.69         0.02         1.34         0.08         1.52         0.17	1.14 0.16 -1.12 0.20 0.01 1.13	-1.46 0.00 1.30 0.02 -0.08 1.38 -	-1.29 0.26 1.37 0.08 0.04 1.33	1.85         0.76         1.19         0.59         1.52         0.33         1.28	3.06 0.09 1.83 0.03 2.45 0.61
-1.41 0.82 1.51 0.07 0.05 1.46 1.29	<u>2.47</u> 0.54 <u>-1.24</u> 0.64 0.62 1.86 1.26 0.54	1.78 0.25 1.06 0.91 1.42 0.36 1.28	-1.02 0.92 -1.02 0.92 -1.02 0.00 1.07	-1.96 0.20 1.17 0.58 -0.40 1.57 -1.03	-1.80 0.10 2.14 0.01 0.17 1.97 1.17	2.28 0.11 1.09 0.60 1.69 0.60 1.09	-1.64         0.03         1.21         0.74         -0.22         1.43         2.17	-1.04 0.97 3.53 0.14 1.25 2.29 1.19	<b>-1.90</b> 0.08 <b>4.62 0.00 1.36 3.26 1.16</b>	-2.32 0.00 2.83 0.01 0.26 2.58 1.64	<b>-1.46</b> 0.50 1.61 0.27 0.08 1.54 1.13	3.27         0.00         -1.22         0.32         1.03         2.25         -1.32	-2.09 0.02 1.38 0.46 -0.36 1.74 1.10	-1.22 0.82 -1.17 0.79 -1.20 0.03 1.16	1.81         0.02         1.45         0.13         1.63         0.18         -1.32	1.94         0.02         1.46         0.04         1.70         0.24         1.11	1.96         0.02         6.98         0.01         4.47         2.51         1.14	<b>-1.15</b> 0.64 1.46 0.02 0.16 1.31 1.07	-1.86         0.01         1.71         0.02         -0.08         1.79         1.55	1.39         0.22         1.83         0.03         1.61         0.22         1.16	2.20 0.03 1.06 0.76 1.63 0.57 1.62 0.15	1.10         0.88         -8.43         0.03         -3.67         4.77         1.90	1.79 0.10 1.01 0.73 1.40 0.39 1.34	1.61         0.38         -1.21         0.74         0.20         1.41         1.13         0.84         .	1.24 0.08 -1.13 0.40 0.06 1.19 -1.28	-1.69 0.40 -1.51 0.21 -1.60 0.09 1.17 0.17 -	-2.55 0.02 -1.59 0.04 -2.07 0.48 -1.21	0.92 0.86 -1.25 0.43 -0.17 1.08 -1.37	-2.65 0.07 -1.07 0.60 -1.86 0.79 -1.40	-2.01 0.61 -1.26 0.36 -1.64 0.38 -1.14	1.19         0.15         -1.20         0.19         -0.01         1.20         -1.07	1.13         0.66         -0.98         0.95         0.07         1.06         1.31	1.69         0.02         1.34         0.08         1.52         0.17         1.39	1.14 0.16 -1.12 0.20 0.01 1.13 1.10	-1.46 0.00 1.30 0.02 -0.08 1.38 -1.44	-1.29 0.26 1.37 0.08 0.04 1.33 1.28	1.85         0.76         1.19         0.59         1.52         0.33         1.28         0.12	3.06         0.09         1.83         0.03         2.45         0.61         1.40
-1.41 0.82 1.51 0.07 0.05 1.46 1.29 0.05	· 2.47 0.54 -1.24 0.64 0.62 1.86 1.26 0.54 1.24	1.78 0.25 1.06 0.91 1.42 0.36 1.28 0.20	-1.02 0.92 -1.02 0.92 -1.02 0.00 1.07 0.92	-1.96 0.20 1.17 0.58 -0.40 1.57 -1.03 0.92 1.35	-1.80         0.10         2.14         0.01         0.17         1.97         1.17         0.05	2.28         0.11         1.09         0.60         1.69         0.60         1.09         0.68	<b>-1.64 0.03 1.21 0.74 -0.22 1.43 2.17 0.02 1.36</b>	-1.04         0.97         3.53         0.14         1.25         2.29         1.19         0.70	<b>-1.90</b> 0.08 <b>4.62 0.00 1.36 3.26 1.16</b> 0.18	-2.32 0.00 2.83 0.01 0.26 2.58 1.64 0.01	-1.46         0.50         1.61         0.27         0.08         1.54         1.13         0.79	3.27 0.00 -1.22 0.32 1.03 2.25 -1.32 0.06 -	-2.09 0.02 1.38 0.46 -0.36 1.74 1.10 0.81	-1.22 0.82 -1.17 0.79 -1.20 0.03 1.16 0.74 -	1.81         0.02         1.45         0.13         1.63         0.18         -1.32         0.13         .	1.94         0.02         1.46         0.04         1.70         0.24         1.11         0.36	1.96         0.02         6.98         0.01         4.47         2.51         1.14         0.37	-1.15 0.64 1.46 0.02 0.16 1.31 1.07 0.75 -	-1.86 0.01 1.71 0.02 -0.08 1.79 1.55 0.01	1.39         0.22         1.83         0.03         1.61         0.22         1.16         0.59	2.20 0.03 1.06 0.76 1.63 0.57 1.62 0.15	1.10         0.88         -8.43         0.03         -3.67         4.77         1.90         0.23	1.79 0.10 1.01 0.73 1.40 0.39 1.34 0.08	1.61         0.38         -1.21         0.74         0.20         1.41         1.13         0.84         .	1.24         0.08         -1.13         0.40         0.06         1.19         -1.28         0.16         .	-1.69 0.40 -1.51 0.21 -1.60 0.09 1.17 0.17 -1.38	-2.55 0.02 -1.59 0.04 -2.07 0.48 -1.21 0.06	0.92 0.86 -1.25 0.43 -0.17 1.08 -1.37 0.25	-2.65 0.07 -1.07 0.60 -1.86 0.79 -1.40 0.05 -	-2.01 0.61 -1.26 0.36 -1.64 0.38 -1.14 0.07	<b>1.19</b> 0.15 -1.20 0.19 -0.01 1.20 -1.07 0.13 -	1.13         0.66         -0.98         0.95         0.07         1.06         1.31         0.31	1.69         0.02         1.34         0.08         1.52         0.17         1.39         0.14	1.14 0.16 -1.12 0.20 0.01 1.13 1.10 0.41	-1.46 0.00 1.30 0.02 -0.08 1.38 -1.44 0.01 -1.31	-1.29 0.26 1.37 0.08 0.04 1.33 1.28 0.09 1.14	1.85         0.76         1.19         0.59         1.52         0.33         1.28         0.12         1.24	<b>3.06</b> 0.09 <b>1.83 0.03 2.45</b> 0.61 <b>1.40</b> 0.13
-1.41 0.82 1.51 0.07 0.05 1.46 1.29 0.05 1.13	· 2.47 0.54 -1.24 0.64 0.62 1.86 1.26 0.54 1.24	1.78 0.25 1.06 0.91 1.42 0.36 1.28 0.20 1.87	-1.02 0.92 -1.02 0.92 -1.02 0.00 1.07 0.92 2.03	-1.96 0.20 1.17 0.58 -0.40 1.57 -1.03 0.92 1.35	-1.80         0.10         2.14         0.01         0.17         1.97         1.17         0.05         1.17	2.28 0.11 1.09 0.60 1.69 0.60 1.09 0.68 1.93	<b>-1.64 0.03 1.21 0.74 -0.22 1.43 2.17 0.02 1.36</b>	-1.04 0.97 3.53 0.14 1.25 2.29 1.19 0.70 1.34	<b>-1.90</b> 0.08 <b>4.62 0.00 1.36 3.26 1.16</b> 0.18 <b>1.13</b>	-2.32 0.00 2.83 0.01 0.26 2.58 1.64 0.01 2.00	<b>-1.46</b> 0.50 1.61 0.27 0.08 1.54 1.13 0.79 1.38	3.27         0.00         -1.22         0.32         1.03         2.25         -1.32         0.06         -1.93	-2.09 0.02 1.38 0.46 -0.36 1.74 1.10 0.81 1.33	-1.22 0.82 -1.17 0.79 -1.20 0.03 1.16 0.74 -1.19	1.81         0.02         1.45         0.13         1.63         0.18         -1.32         0.13         -1.40	1.94         0.02         1.46         0.04         1.70         0.24         1.11         0.36         -1.32	1.96         0.02         6.98         0.01         4.47         2.51         1.14         0.37         1.03	-1.15 0.64 1.46 0.02 0.16 1.31 1.07 0.75 -1.34	<b>-1.86 0.01</b> 1.71 <b>0.02 -0.08</b> 1.79 1.55 <b>0.01</b> 1.63	1.39         0.22         1.83         0.03         1.61         0.22         1.16         0.59         1.01	2.20         0.03         1.06         0.76         1.63         0.57         1.62         0.15         -1.22	1.10         0.88         -8.43         0.03         -3.67         4.77         1.90         0.23         1.63         0.42	1.79         0.10         1.01         0.73         1.40         0.39         1.34         0.08         1.77	1.61         0.38         -1.21         0.74         0.20         1.41         1.13         0.84         -1.16         0.88         .	1.24         0.08         -1.13         0.40         0.06         1.19         -1.28         0.16         -1.44	-1.69 0.40 -1.51 0.21 -1.60 0.09 <b>1.17</b> 0.17 -1.38 0.36	-2.55 0.02 -1.59 0.04 -2.07 0.48 -1.21 0.06 1.20	0.92 0.86 -1.25 0.43 -0.17 1.08 -1.37 0.25 1.19	-2.65 0.07 -1.07 0.60 -1.86 0.79 -1.40 0.05 -1.39 0.24	-2.01 0.61 -1.26 0.36 -1.64 0.38 -1.14 0.07 1.19 0.64	1.19         0.15         -1.20         0.19         -0.01         1.20         -1.07         0.13         -1.26	1.13         0.66         -0.98         0.95         0.07         1.06         1.31         0.31         -1.08	1.69         0.02         1.34         0.08         1.52         0.17         1.39         0.14         1.48	1.14 0.16 -1.12 0.20 0.01 1.13 1.10 0.41 1.05	-1.46 0.00 1.30 0.02 -0.08 1.38 -1.44 0.01 -1.31	-1.29 0.26 1.37 0.08 0.04 1.33 1.28 0.09 1.14	1.85         0.76         1.19         0.59         1.52         0.33         1.28         0.12         1.24         0.58	<b>3.06</b> 0.09 <b>1.83 0.03 2.45</b> 0.61 <b>1.40</b> 0.13 <b>1.26</b>

### Chapter 10: Appendix

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SPO2259	SPO2258	SPO2257	SPO2256	SPO2255	SPO2253	SPO2252	SPO2251	SPO2250	SPO2249	SPO2248	SPO2247	SPO2246	SPO2245	SPO2244	SPO2243	SPO2242	SPO2241	SPO2240	SPO2239	SPO2238	SPO2237	SPO2236	SPO2235	SPO2234	SPO2233	SPO2232	SPO2231	SPO2230	SPO2229	SPO2228	SPO2227	SPO2226	SPO2225	SPO2224	SPO2223	SPO2222	SPO2221
											cysE	cysK				pdhC	pdhB	pdhA					pgk									queA					
head-tail adaptor	hypothetical protein	TP901-1 family major tail protein	hypothetical protein	dnaK suppressor protein	O-acetyltransferase (EC:2.3.1.30)	synthase A (EC:2.5.1.47)	hypothetical protein	hypothetical protein	hypothetical protein	(EC:2.3.1.12)	dehydrogenase subunit beta (EC:1.2.4.1)	dehydrogenase complex, E1 component subuntt alpha (EC:1.2.4.1)	hypothetical protein	LysR family transcriptional regulator	hypothetical protein	hypothetical protein	kinase (EC:2.7.2.3)	cyclophilin type peptidyl-prolyl cis-trans isomerase	cyclophilin type peptidyl-prolyl cis-trans isomerase	hypothetical protein	hypothetical protein	M24/M37 family peptidase	hypothetical protein	AhpC/TSA family protein	hypothetical protein	ribosyltransferase-isomerase	hypothetical protein	hypothetical protein	hypothetical protein	dihydrolipoamide dehydrogenase (EC:1.8.1.4)	von Willebrand factor A						
1.32	1.27	2.23	-1.12	1.61	-1.18	х	1.09	-2.32	x	-1.10	1.52	1.37	2.53	1.29	1.28	3.48	3.69	1.97	-1.71	х	1.39	-1.70	-1.31	0.98	0.95	-2.30	0.96	1.21	1.25	-1.36	1.11	х	3.11	2.81	1.81	2.26	х
0.83	0.95	0.16	0.14	0.13	0.60	×	×	0.26	×	0.24	0.34	0.78	0.33	0.21	0.14	0.00	0.16	0.06	0.01	×	0.07	0.43	0.12	0.67	0.55	0.00	0.62	0.92	0.14	0.75	0.79	×	0.02	0.00	0.05	0.10	×
1.22	1.09	1.15	1.14	1.41	1.27	1.10	1.35	-1.21	1.14	-1.00	-1.90	-1.70	1.84	1.10	1.18	1.08	-1.12	-1.28	-1.36	-1.64	-1.67	-1.40	1.12	-1.39	1.70	1.53	0.99	-1.35	-1.48	-1.26	-1.19	1.35	1.75	1.27	1.95	1.70	1.05
0.68	0.87	0.26	0.53	0.13	0.27	0.22	0.29	0.19	0.47	0.93	0.12	0.19	0.02	0.60	0.01	0.80	0.17	0.49	0.05	0.02	0.30	0.07	0.38	0.06	0.03	0.01	0.55	0.19	0.05	0.54	0.11	0.68	0.04	0.63	0.01	0.17	0.38
1.27	1.18	1.69	0.01	1.51	0.05	N/A	1.22	-1.77	N/A	-1.05	-0.19	-0.17	2.19	1.20	1.23	2.28	1.29	0.35	-1.54	N/A	-0.14	-1.55	-0.10	-0.21	1.33	-0.39	0.97	-0.07	-0.12	-1.31	-0.04	N/A	2.43	2.04	1.88	1.98	N/A
0.05	0.09	0.54	1.13	0.10	1.23	N/A	0.13	0.56	N/A	0.05	1.71	1.54	0.35	0.10	0.05	1.20	2.41	1.63	0.17	N/A	1.53	0.15	1.22	1.18	0.37	1.92	0.01	1.28	1.37	0.05	1.15	N/A	0.68	0.77	0.07	0.28	N/A
1.24	1.44	1.65	-1.25	2.65	-2.10	Х	-1.01	-3.11	1.51	-0.96	1.89	5.11	-1.79	-1.01	2.38	4.30	5.51	2.02	-1.63	1.84	-1.05	-1.68	-1.81	2.82	2.83	-1.98	1.60	1.68	2.15	1.60	1.07	3.31	2.05	-2.08	2.26	4.17	3.03
0.80	0.89	0.02	0.06	0.04	0.06	х	0.75	0.06	x	0.23	0.14	0.04	0.58	0.38	0.02	0.00	0.00	0.01	0.01	0.01	0.56	0.70	0.03	0.12	0.01	0.04	0.14	0.14	0.08	0.30	0.39	0.40	0.02	0.01	0.05	0.02	0.01
2.61	3.56	2.97	2.41	4.39	2.63	2.78	3.88	1.96	1.49	1.32	1.61	2.40	1.42	1.49	1.58	1.13	1.36	0.96	-1.27	-1.11	-1.40	-1.20	1.09	-1.50	-1.10	1.81	1.34	1.40	1.09	1.38	1.26	-1.73	-1.56	1.21	1.16	1.37	-1.16
0.03	0.14	0.00	0.02	0.00	0.01	0.04	0.03	0.16	0.31	0.03	0.04	0.02	0.37	0.27	0.01	0.43	0.38	0.53	0.25	0.71	0.30	0.55	0.83	0.09	0.23	0.11	0.08	0.10	0.60	0.34	0.44	0.58	0.10	0.25	0.27	0.03	0.58
1.93	2.50	2.31	0.58	3.52	0.27	N/A	1.44	-0.58	1.50	0.18	1.75	3.76	-0.19	0.24	1.98	2.72	3.44	1.49	-1.45	0.37		-1.44	-0.36	0.66	0.87	-0.09	1.47	1.54	1.62	1.49	1.17	0.79	0.25	-0.44	1.71	2.77	0.94
0.69	1.06	0.66	1.83	0.87	2.37	N/A	2.45	2.54	0.01	1.14	0.14	1.36	1.61	1.25	0.40	1.59	2.08	0.53	0.18	1.48	0.17	0.24	1.45	2.16	1.97		0.13	0.14	0.53	0.11	0.10	2.52	1.81	. 1.65	0.55	1.40	2.10
1.32	-1.03	1.16	1.07	1.04	1.24	1.15	1.17	1.04	1.29	1.19	1.53	1.14	-1.41	-1.05	1.17	-1.24	-1.24	-1.64	-1.05	-0.98	1.18	1.50	-1.27	-1.16	1.79	-	1.23	-1.74	-1.25	-1.13	1.36		_		1.31	1.21	1.13
0.24	3 0.96	0.19	0.25	0.75	0.14	0.45	0.25	0.88	0.37	0.22	0.04	0.71	0.48	5 0.95	0.11	4 0.07	4 0.11	4 0.08	5 0.65	3 0.45	0.31	0.02	-	5 0.30	0.01		0.35		5 0.57	3 0.50	0.15	0.59		0.58		0.70	0.03
1.06	-1.47	-	-1.01	-1.28	-1.16	1.34	-1.05	3 1.59	×	-1.15	2.09	1.59	-1.10	-1.47	1.62	1.17	1.53	1.06	1.12	-1.00	1.08	x	1.90	-1.34	1.42	x	1.42		-1.58			-2.45		0.92		1.69	-0.89
0.86	0.74	5 0.01	0.84	3 0.67	5 0.57	×	0.86	0.05	x	5 0.85	0.04	0.10	0.88	0.52	0.06	0.79	0.12	0.71	0.64	0.81	0.45	x	0.02	4 0.01	0.04	x	0.52		3 0.04	0.70	4 0.93	0.44					0.22
6 1.19	4 -1.25		4 0.03		7 0.04	1.25	6 0.06	5 1.32	N/A	5 0.02	4 1.81	0 1.37	-1.26		6 1.40	9 -0.04	2 0.15	-0.29	4 0.04	-0.99	5 1.13	N/A	2 0.32	-1.25	4 1.61		2 1.33	4 -1.54	4 -1.42	0 -0.07				-	-		2 0.12
0.13	0.22	1.96	1.04	1.16	1.20	0.10	1.11	0.28	N/A	1.17	0.28	0.23	0.16	5 0.21	0.23	4 1.21	1.39	) 1.35	1.09	0.01	0.05	. N/A	1.59	5 0.09	0.18				0.17	7 1.06	1.20	3 0.47	0.18	0.09	0.11	0.24	1.01

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SPO2300	SPO2299 d	SPO2298	SPO2297	SPO2296	SPO2295 g	SPO2294 g	SPO2293	SPO2292	SPO2290	SPO2289	SPO2288 tig	SPO2287	SPO2286 lt	SPO2285	SPO2284	SPO2283 r	SPO2282 r	SPO2281 r	SPO2280	SPO2279	SPO2278	SPO2277	SPO2276 fc	SPO2275 fc	SPO2274 a	SPO2273	SPO2272	SPO2271 fc	SPO2270	SPO2269	SPO2268	SPO2267	SPO2266	SPO2264	SPO2263	SPO2262	SPO2261	0102200
	dddP				glnA	glnB-1					00		luxR-2			rplI	rpsR	rpsF					fabD	fabG	acpP			fabF										
hypothetical protein	DMSP lyase	hypothetical protein	hypothetical protein	hypothetical protein	synthetase, type I (EC:6.3.1.2)	regulatory protein P-II	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	factor (EC:5.2.1.8)	autoinducer synthesis protein	family transcriptional regulator	DNA-binding protein	transglycosylase	ribosomal protein L9	ribosomal protein S18	ribosomal protein S6	hypothetical protein	hypothetical protein	hypothetical protein	cytochrome	S-malonyltransferase (EC:2.3.1.39)	reductase (EC:1.1.1.100)	carrier protein	acetyltransferase	phosphoribosylformimino-5-aminoimidazole carboxamide ribotide isomerase	synthase (EC:2.3.1.41)	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	terminase large subunit	HK97 family portal protein	hypothetical protein	HK97 family phage prohead protease	HK97 family major capsid protein	пурошенсат ргонян

-1.04	13.30	-1.36	2.56	-1.75	1.49	2.29	1.37	-1.11	x	1.00	1.43	0.88	-1.29	1.62	2.22	1.09	-1.48	-2.02	1.38	-1.39	-1.14	1.09	1.57	1.49	1.49	-1.08	5.00	2.09	1.42	-1.48	-1.20	х	1.73	х	-1.13	1.97	-1.54	1.20
0.98	0.02	0.01	0.03	0.17	0.74	0.01	0.06	0.25	×	0.82	0.41	0.01	0.54	0.01	0.49	1.00	0.00	0.01	0.19	0.28	0.92	0.58	0.07	0.02	0.03	0.20	0.00	0.00	0.01	0.01	0.70	×	0.37	×	0.79	0.11	0.03	0.77
-1.07	4.01	-1.10	1.30	-1.45	-1.27	-1.53	-1.27	1.52	x	-4.01	1.48	-1.19	-1.26	1.04	1.01	1.49	-1.27	-1.72	1.28	-2.03	-1.22	-1.27	-1.20	1.14	1.04	1.12	1.45	-1.07	-1.08	-1.06	-1.39	x	-1.23	1.05	-1.14	1.14	1.09	1.29
0.96	0.06	0.72	0.06	0.07	0.16	0.08	0.49	0.05	x	0.26	0.29	0.47	0.62	0.92	0.83	0.35	0.12	0.03	0.20	0.09	0.90	0.05	0.37	0.31	1.00	0.69	0.06	0.69	0.29	0.71	0.20	x	0.67	0.80	0.81	0.51	0.75	0.39
-1.06	8.66	-1.23	1.93	-1.60	0.11	0.38	0.05	0.21	N/A	-1.51	1.46	-0.16	-1.28	1.33	1.62	1.29	-1.38	-1.87	1.33	-1.71	-1.18	-0.09	0.19	1.32	1.27	0.02	3.23	0.51	0.17	-1.27	-1.30	N/A	0.25	N/A	-1.14	1.56	-0.23	1.25
0.02	4.65	0.13	0.63	0.15	1.38	1.91	1.32	1.32	N/A	2.50	0.03	1.03	0.02	0.29	0.61	0.20	0.11	0.15	0.05	0.32	0.04	1.18	1.39	0.18	0.23	1.10	1.78	1.58	1.25	0.21	0.10	N/A	1.48	N/A	0.01	0.42	1.32	0.05
4.40	-1.62	-2.71	-2.16	-1.61	4.33	6.56	1.88	-0.98	x	1.45	5.87	-1.83	-1.35	0.89	3.56	5.00	8.93	7.75	-2.43	1.01	1.29	-1.43	2.62	2.91	3.92	1.30	-1.72	4.16	2.44	-1.67	1.06	х	1.65	-0.95	-1.44	-1.06	-4.29	-1.17
0.07	0.37	0.00	0.01	0.12	0.07	0.00	0.05	0.22	х	0.48	0.02	0.00	0.47	0.06	0.02	0.18	0.00	0.00	0.01	0.76	0.85	0.19	0.01	0.00	0.02	0.04	0.09	0.00	0.02	0.04	0.48	x	0.32	0.15	0.20	0.52	0.01	0.50
1.54	1.34	1.16	1.46	1.65	1.51	1.63	-1.26	1.22	х	-2.93	-1.67	2.42	1.65	1.45	1.20	-7.10	-5.14	-5.03	1.28	-1.51	1.30	-1.24	0.99	-1.90	-1.73	-1.11	-1.08	-1.06	-1.29	-0.97	-1.10	х	-1.11	1.90	3.54	4.35	3.87	2.99
0.17	0.40	0.29	0.04	0.06	0.14	0.01	0.08	0.41	x	0.31	0.15	0.01	0.47	0.07	0.59	0.00	0.00	0.00	0.09	0.29	0.84	0.50	0.54	0.10	0.14	0.32	0.33	0.09	0.02	0.67	0.12	x	0.78	0.01	0.02	0.00	0.00	0.02
2.97	-0.14	-0.78	-0.35	0.02	2.92	4.10	0.31	0.12	N/A	-0.74	2.10	0.30	0.15	1.17	2.38	-1.05	1.90	1.36	-0.58	-0.25	1.30	-1.34	1.81	0.51	1.10	0.10	-1.40	1.55	0.58	-1.32	-0.02	N/A	0.27	0.47	1.05	1.65	-0.21	0.91
1.43	1.48	1.94	1.81	1.63	1.41	2.47	1.57	1.10	N/A	2.19	3.77	2.13	1.50	0.28	1.18	6.05	7.04	6.39	1.86	1.26	0.01	0.10	0.81	2.41	2.83	1.21	0.32	2.61	1.87	0.35	1.08	N/A	1.38	1.43	2.49	2.71	4.08	2.08
1.15	1.33	-1.16	-1.28	-1.32	-1.31	-1.72	1.28	-1.40	х	1.24	-1.05	1.54	-1.08	1.35	1.33	1.56	-1.86	-2.06	-1.48	1.42	1.06	-1.17	-1.17	1.21	1.46	2.02	1.82	1.08	-1.11	-1.02	1.22	х	1.26	1.07	-1.25	-1.37	-1.37	1.15
0.44	0.48	0.58	0.13	0.04	0.81	0.00	0.25	0.39	х	0.85	0.83	0.00	0.83	0.02	0.03	0.09	0.00	0.02	0.01	0.11	0.97	0.03	0.27	0.35	0.13	0.00	0.00	0.62	0.01	0.96	0.38	x	0.38	0.58	0.23	0.05	0.00	0.59
1.33	1.16	1.32	1.63	-1.18	-1.53	-1.58	1.53	-1.76	x	1.75	-1.41	1.89	1.11	1.79	1.30	-2.03	-2.94	-4.15	0.98	1.41	1.21	-0.97	-1.14	0.93	0.99	1.52	1.23	0.99	0.97	1.65	1.07	х	1.68	-3.30	-1.18	-2.56	-2.74	-1.43
0.29	0.15	0.06	0.01	0.04	0.23	0.00	0.10	0.09	х	0.69	0.33	0.02	0.99	0.02	0.22	0.08	0.03	0.00	0.24	0.21	0.72	0.38	0.10	0.07	0.20	0.15	0.04	0.50	0.56	0.36	0.38	x	0.04	0.04	0.80	0.01	0.02	0.54
1.24	1.25	0.08	0.18	-1.25	-1.42	-1.65	1.41	-1.58	N/A	1.50	-1.23	1.72	0.02	1.57	1.32	-0.24	-2.40	-3.11	-0.25	1.42	1.14	-1.07	-1.16	1.07	1.22	1.77	1.53	1.04	-0.07	0.32	1.15	N/A	1.47	-1.12	-1.22	-1.97	-2.06	-0.14
0.09	0.09	1.24	1.46	0.07	0.11	0.07	0.13	0.18	N/A	0.26	0.18	0.18	1.10	0.22	0.02	1.80	0.54	1.05	1.23	0.01	0.08	0.10	0.02	0.14	0.24	0.25	0.30	0.04	1.04	1.34	0.08	N/A	0.21	2.19	0.04	0.60	0.69	1.29

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SPO2339	SPO2338	SPO2337	SPO2336	SPO2335	SPO2334	SPO2333	SPO2332	SPO2331	SPO2330	SPO2329	SPO2328	SPO2327	SPO2326	SPO2325	SPO2324	SPO2323	SPO2322	SPO2321	SPO2320	SPO2319	SPO2318	SPO2317	SPO2316	SPO2315	SPO2314	SPO2313	SPO2312	SPO2311	SPO2310	SPO2309	SPO2308	SPO2307	SPO2306	SPO2305	SPO2304	SPO2303	SPO2302	SPO2301
	rarD-1										engA			serS			yajC	secD	secF			ccmA	ccmB	ccmC			acnA				fliG			purB				
enoyl-CoA hydratase	RarD	decarboxylase	lysM domain-containing protein	ABC transporter ATP-binding protein/permease	CAAX amino terminal protease	hypothetical protein	AcrB/AcrD/AcrF family transporter	RND family efflux transporter MFP subunit	hypothetical protein	PQQ repeat-containing protein	protein EngA	fatty acid desaturase	hypothetical protein	synthetase (EC:6.1.1.11)	hypothetical protein	mechanosensitive ion channel protein MscS	translocase subunit YajC	translocase subunit SecD	translocase subunit SecF	hypothetical protein	hypothetical protein	c biogenesis protein CcmA	exporter protein CcmB	exporter protein CcmC	DsbE subtamily periplasmic protein thiol/disulfide oxidoreductase	hypothetical protein	hydratase (EC:4.2.1.3)	hypothetical protein	lipid A biosynthesis lauroyl acyltransferase	hypothetical protein	motor switch protein FliG	hypothetical protein	hypothetical protein	lyase (EC:4.3.2.2)	hypothetical protein	hypothetical protein	bioY family protein	osmC-like family protein
-1.31	1.07	2.21	1.30	1.19	-1.23	-1.12	1.07	0.98	1.02	1.42	-1.67	1.58	1.05	1.79	17.50	-1.36	1.18	1.15	1.28	-1.19	1.25	1.07	1.89	2.62	2.13	-1.21	6.55	x	-1.35	-1.41	1.75	1.15	-1.90	1.60	1.16	1.98	3.66	-1.14
0.66	0.87	0.22	0.06	0.14	0.88	0.75	0.98	0.88	0.74	0.16	0.00	0.66	0.65	0.19	0.00	0.61	0.40	0.04	0.08	0.53	×	1.00	0.68	0.02	0.04	0.39	0.03	х	0.01	x	0.27	0.45	0.20	0.09	0.78	0.01	0.00	0.13
-1.11	1.41	1.03	1.16	1.05	1.20	-1.43	-2.16	-1.52	1.06	1.43	1.04	-1.21	-1.15	1.00	3.07	-1.03	-1.42	-1.13	1.65	-1.03	1.55	1.81	1.48	1.28	1.34	1.15	1.23	х	1.09	-1.10	1.13	1.17	-1.22	1.41	1.77	1.38	1.45	-1.14
0.75	0.11	0.93	0.22	0.83	0.76	0.11	0.03	0.10	0.88	0.22	0.76	0.51	0.03	0.70	0.06	0.95	0.20	0.54	0.01	0.92	0.15	0.23	0.26	0.17	0.15	0.76	0.29	×	0.42	0.92	0.33	0.64	0.04	0.02	0.08	0.23	0.20	0.74
-1.21	1.24	1.62	1.23	1.12	-0.02	-1.28	-0.55	-0.27	1.04	1.43	-0.32	0.19	-0.05	1.39	10.29	-1.20	-0.12	0.01	1.47	-1.11	1.40	1.44	1.69	1.95	1.74	-0.03	3.89	N/A	-0.13	-1.26	1.44	1.16	-1.56	1.51	1.47	1.68	2.56	-1.14
0.10	0.17	0.59	0.07	0.07	1.22	0.16	1.62	1.25	0.02	0.01	1.36	1.40	1.10	0.40	7.22	0.17	1.30	1.14	0.19	0.08	0.15	0.37	0.20	0.67	0.40	1.18	2.66	N/A	1.22	0.16	0.31	0.01	0.34	0.10	0.31	0.30	1.11	0.00
-1.22	1.83	4.07	1.83	2.63	-2.14	2.05	2.79	2.53	2.36	2.42	2.48	1.42	0.95	2.79	-1.66	-2.05	4.38	5.96	6.96	1.01	2.51	1.82	3.25	4.08	2.37	-3.22	4.67	х	1.96	1.23	-1.76	-2.53	-7.32	9.61	1.77	2.49	4.88	-1.28
0.76	0.20	0.01	0.00	0.01	0.69	0.09	0.01	0.06	0.03	0.02	0.00	0.76	0.12	0.11	0.06	0.41	0.00	0.00	0.00	0.81	0.05	0.44	0.44	0.02	0.02	0.01	0.00	x	0.01	0.87	0.38	0.00	0.01	0.00	0.05	0.00	0.00	0.07
1.21	-1.22	-1.19	1.09	-1.22	-1.18	-1.85	-1.62	-1.23	-1.16	-1.20	-1.42	1.49	1.22	-1.24	1.43	1.49	1.20	-1.14	-1.37	-1.12	-1.13	1.03	-1.22	-1.03	0.99	-1.25	1.65	1.13	-1.07	-1.82	1.57	1.89	2.00	-1.28	-1.45	-1.27	1.36	1.18
0.68	0.17	0.17	0.52	0.08	0.82	0.03	0.10	0.42	0.34	0.50	0.01	0.33	0.35	0.04	0.04	0.45	0.11	0.06	0.08	0.65	0.44	0.98	0.42	0.71	0.87	0.78	0.10	x	0.31	0.42	0.03	0.01	0.08	0.10	0.01	0.07	0.36	0.31
-0.01	0.31	1.44	1.46	0.71	-1.66	0.10	0.59	0.65	0.60	0.61	0.53	1.46	1.08	0.78	-0.12	-0.28	2.79	2.41	2.80	-0.06	0.69	1.43	1.02	1.53	1.68	-2.24	3.16	N/A	0.45	-0.30	-0.10	-0.32	-2.66	4.17	0.16	0.61	3.12	-0.05
1.22	1.53	2.63	0.37	1.93	0.48	1.95	2.21	1.88	1.76	1.81	1.95	0.04	0.14	2.02	1.55	1.77	1.59	3.55	4.17	1.07	1.82	0.40	2.24	2.56	0.69	0.98	1.51	N/A	1.52	1.53	1.67	2.21	4.66	5.45	1.61	1.88	1.76	1.23
1.85	1.12	-1.12	-1.20	-1.17	-1.06	1.06	-1.11	1.38	1.04	1.66	-1.54	1.25	1.16	-1.19	1.25	1.23	-1.10	-1.13	1.37	-1.00	-1.47	1.28	-1.23	1.33	1.17	1.05	-1.12	-1.35	1.13	-1.26	1.11	1.47	1.21	-1.49	-1.02	-1.50	1.59	-1.55
0.16	0.43	0.08	0.06	0.20	0.93	0.77	0.33	0.25	0.98	0.07	0.02	0.69	0.50	0.09	0.40	0.63	0.47	0.27	0.02	0.99	0.09	0.49	0.35	0.37	0.72	0.64	0.09	х	0.58	0.80	0.32	0.02	0.40	0.12	0.87	0.07	0.10	0.03
1.73	-1.41	-1.47	-1.12	-1.67	-1.01	1.68	2.69	3.33	-1.30	1.32	-2.09	1.92	1.65	-1.28	1.10	1.78	1.11	0.97	-1.14	1.54	-2.07	1.12	-1.36	0.98	1.26	1.52	-1.62	x	-1.05	1.12	×	1.28	2.24	-1.95	-1.11	-1.79	1.90	-1.17
0.38	0.06	0.16	0.03	0.16	0.96	0.05	0.02	0.02	0.05	0.19	0.01	0.14	0.05	0.21	0.46	0.35	0.08	0.38	0.13	0.08	0.10	0.84	0.32	0.51	0.44	x	0.06	x	0.33	0.92	x	0.19	0.01	0.01	0.04	0.06	0.10	0.87
1.79	-0.15	-1.30	-1.16	-1.42	-1.04	1.37	0.79	2.36	-0.13	1.49	-1.82	1.59	1.41	-1.24	1.18	1.51	0.01	-0.08	0.12	0.27	-1.77	1.20	-1.30	1.16	1.22	1.29	-1.37	N/A	0.04	-0.07	N/A	1.38	1.73	-1.72	-1.07	-1.65	1.75	-1.36
0.06	1.27	0.18	0.04	0.25	0.03	0.31	1.90	0.98	1.17	0.17	0.27	0.34	0.25	0.05	0.08	0.28	1.11	1.05	1.26	1.27	0.30	0.08	0.07	0.17	0.05	0.23	0.25	N/A	1.09	1.19	N/A	0.10	0.52	0.23	0.05	0.15	0.16	0.19

SP02340 SP02341 SP02342 SP02343 SP02344 SP02344 SP02345 SP02346

### Chapter 10: Appendix

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acyl-CoA dehydrogenase (EC:1.3.99.13)	selenium-binding protein	ferredoxin	CoA-binding domain-containing protein	tRNA-dihydrouridine synthase A	hypothetical protein	AsnC family transcriptional regulator	universal stress protein family protein	sodium:alanine symporter family protein	LuxR family transcriptional regulator	Na/Pi-cotransporter family protein	amino acid ABC transporter ATP-binding protein	amino acid ABC transporter permease	amino acid ABC transporter permease	annio acia ABC nansporter suosuate-omaning protein	phosphatase (EC:3.1.3.3)	protein .	(EC:2.7.3) amino acid ABC transporter substrate-binding	dehydrogenase	TRAP transporter	TRAP transporter	TRAP transporter	dissimilation regulator	hypothetical protein	hypothetical protein	type I secretion target repeat-containing protein	hypothetical protein	Holliday junction resolvase-like protein	c-type biogenesis protein CycH	sarcosine oxidase subunit beta	hypothetical protein	sarcosine oxidase subunit delta	sarcosine oxidase subunit alpha	sarcosine oxidase subunit gamma	hypothetical protein	hypothetical protein	FkbM family methyltransferase	dismutase, Fe (EC:1.15.1.1)
-0.97	1.29	1.39	1.40	-1.20	1.25	-1.36	-5.95	-2.34	1.03	х	1.00	Х	х	×	-1.57	-1.93	-1.34	-1.09	-1.42	-1.40	-1.21	1.21	-1.03	х	-1.22	3.01	1.37	1.09	3.12	2.38	2.54	4.21	4.53	2.01	1.98	-1.45	1.57
0.94	0.68	0.22	0.21	0.23	0.14	0.06	0.04	0.43	0.89	×	0.78	×	×	х	0.19	0.01	0.13	0.94	0.83	0.46	0.69	0.27	0.90	×	×	0.01	0.11	0.90	0.38	0.01	0.55	0.01	0.00	0.25	0.06	0.59	0.34
-1.23	-1.22	1.05	1.16	-1.26	1.13	-1.27	-3.26	-1.33	-1.21	х	-1.10	-1.13	x	х	-1.30	-1.04	-1.37	-1.03	-1.65	-1.47	-1.10	1.19	1.24	1.26	1.47	1.89	1.94	1.73	2.85	1.76	2.71	2.24	1.70	1.22	1.29	1.21	-1.18
0.72	0.53	0.12	0.75	0.28	0.16	0.60	0.03	0.26	0.07	×	0.92	х	×	х	0.60	0.54	0.21	0.98	0.67	0.61	0.16	0.03	0.47	0.19	0.09	0.02	0.09	0.03	0.00	0.13	0.01	0.01	0.08	0.52	0.26	0.37	0.76
-1.10	0.04	1.22	1.28	-1.23	1.19	-1.32	-4.61	-1.84	-0.09	N/A	-0.05	N/A	N/A	N/A	-1.44	-1.49	-1.36	-1.06	-1.54	-1.44	-1.16	1.20	0.11	N/A	0.13	2.45	1.66	1.41	2.99	2.07	2.63	3.23	3.12	1.62	1.64	-0.12	0.20
0.13	1.26	0.17	0.12	0.03	0.06	0.05	1.35	0.51	1.12	N/A	1.05	N/A	N/A	N/A	0.14	0.45	0.02	0.03	0.12	0.04	0.05	0.01	1.14	N/A	1.35	0.56	0.29	0.32	0.14	0.31	0.09	0.99	1.42	0.40	0.35	1.33	1.38
-1.51	-1.72	-1.33	-1.22	1.39	1.16	-2.28	-2.92	-1.86	1.05	х	1.40	-1.52	х	Х	-2.52	-1.86	-1.75	1.67	-1.82	-1.47	-0.92	1.90	1.66	1.96	-1.39	2.28	2.77	4.52	1.60	1.41	1.39	1.75	1.85	0.87	1.13	1.85	2.12
0.57	0.48	0.40	0.82	0.04	0.11	0.03	0.00	0.24	0.30	x	0.71	0.60	×	х	0.05	0.31	0.01	0.23	0.76	0.55	0.19	0.01	0.05	0.06	0.93	0.00	0.01	0.00	0.11	0.02	0.28	0.17	0.04	0.57	0.78	0.25	0.18
-1.11	1.50	1.49	1.20	-1.16	-1.57	-1.47	-4.18	-2.41	-1.05	х	-1.02	Х	x	х	-1.11	1.43	-1.14	6.12	4.75	2.16	4.98	1.13	-1.01	-1.03	-1.06	1.27	1.17	1.10	1.51	1.61	1.19	1.09	-1.15	2.17	1.96	-1.04	-1.52
0.83	0.27	0.19	0.53	0.21	0.07	0.30	0.00	0.08	0.66	x	0.93	х	х	x	0.87	0.10	0.64	0.01	0.14	0.27	0.00	0.14	0.93	0.78	х	0.21	0.05	0.63	0.02	0.03	0.36	0.82	0.24	0.04	0.02	0.76	0.41
-1.31	-0.11	0.08	-0.01	0.12	-0.21	-1.88	-3.55	-2.14	0.00	N/A	0.19	N/A	N/A	N/A	-1.82	-0.22	-1.45	3.90	1.47	0.35	2.03	1.52	0.33	0.47	-1.23	1.78	1.97	2.81	1.56	1.51	1.29	1.42	0.35	1.52	1.55	0.41	0.30
0.20	1.61	1.41	1.21	1.28	1.37	0.41	0.63	0.27	1.05	N/A	1.21	N/A	N/A	N/A	0.71	1.65	0.31	2.23	3.29	1.82	2.95	0.39	1.34	1.50	0.16	0.51	0.80	1.71	0.05	0.10	0.10	0.33	1.50	0.65	0.42	1.45	1.82
-1.50	-1.17	-1.45	1.07	1.27	-1.12	-1.08	-1.20	1.20	-1.21	x	-1.16	-1.24	x	х	1.10	-1.00	-1.04	-1.26	1.20	-1.06	1.17	1.29	1.03	-1.02	1.14	1.30	1.43	1.67	1.22	-1.03	1.18	1.21	-1.05	-1.37	1.03	1.14	1.29
0.26	0.25	0.00	0.79	0.35	0.36	0.55	0.05	0.68	0.22	×	0.92	0.66	×	х	0.76	0.98	0.97	0.47	0.83	0.92	0.25	0.05	0.94	0.92	0.56	0.16	0.08	0.01	0.33	0.79	0.15	0.15	0.41	0.25	0.93	0.40	0.76
1.17	-1.11	-1.35	1.28	1.16	-2.17	-1.40	-1.44	-1.38	-1.19	x	1.19	Х	x	х	1.64	-1.01	1.20	-1.05	1.30	1.34	-1.06	-1.30	-1.06	-1.14	x	-1.21	1.16	1.78	1.31	1.22	1.06	1.04	-1.20	1.45	1.84	-1.33	0.93
0.88	0.75	0.16	0.28	0.74	0.00	0.59	0.02	0.32	0.34	×	0.83	x	x	Х	0.21	0.42	0.41	×	0.75	0.67	0.68	0.09	0.90	0.93	×	0.19	0.50	0.03	0.05	0.50	0.99	0.96	0.17	0.37	0.04	0.34	0.80
-0.17	-1.14	-1.40	1.18	1.22	-1.65	-1.24	-1.32	-0.09	-1.20	N/A	0.02	N/A	N/A	N/A	1.37	-1.01	0.08	-1.16	1.25	0.14	0.05	-0.01	-0.02	-1.08	N/A	0.05	1.30	1.73	1.27	0.10	1.12	1.13	-1.13	0.04	1.44	-0.10	1.11
1.34	0.03	0.05	0.11	0.06	0.53	0.16	0.12	1.29	0.01	N/A	1.18	N/A	N/A	N/A	0.27	0.01	1.12	0.11	0.05	1.20	1.12	1.30	1.05	0.06	N/A	1.26	0.14	0.06	0.05	1.13	0.06	0.09	0.08	1.41	0.41	1.24	0.18

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SP02364 SP02366 SP02366 SP02367 SP02369 SP02370 SP02371 SP02371 SP02372 SP02374 SP02374 SP02377 SP02377

SPO2362 SPO2363 SPO2361

SP02347 SP02348 SP02350 SP02351 SP02352 SP02353 SP02355 SP02355 SP02356 SP02356 SP02356 SP02357 SP02358

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SPO2420	SPO2419	SPO2418	SPO2417	SPO2416	SPO2415	SPO2414	SPO2413	SPO2412	SPO2411	SPO2410	SPO2409	SPO2408	SPO2407	SPO2406	SPO2405	SPO2403	SPO2401	SPO2400	SPO2399	SPO2398	SPO2397	SPO2396	SPO2395	SPO2394	SPO2393	SPO2392	SPO2390	SPO2389	SPO2388	SPO2387	SPO2386	SPO2385	SPO2384	SPO2383	SPO2382	SPO2381
kdg K			idnO															coxC	coxM	coxS-2	coxL-2	coxD	coxE	coxF	coxG											
2-dehydro-3-deoxygluconokinase (EC.2.7.1.45)	hypothetical protein	amino acid transporter LysE	gluconate 5-dehydrogenase (EC:1.1.1.69)	3-hydroxyisobutyrate dehydrogenase	phytanoyl-CoA dioxygenase	mandelate racemase	oxidoreductase	aldolase short chain dehydrogenase/reductase	hypothetical protein	dihydroxy-acid dehydratase (EC:4.2.1.9)	snort chain denydrogenase/reductase oxidoreductase	hypothetical protein	ISSpo6, transposase orfB	ISSpo6, transposase orf A	phage integrase site specific recombinase	hypothetical protein	type I secretion target repeat-containing protein	carbon monoxide dehydrogenase operon C protein	carbon monoxide dehydrogenase, medium subunit (EC:1.2.99.2)	carbon monoxide dehydrogenase, small subunit (EC:1.2.99.2)	(EC:1.2.99.2)	carbon monoxide dehydrogenase D protein	carbon monoxide dehydrogenase E protein	carbon monoxide dehydrogenase F protein	carbon monoxide dehydrogenase G protein	hypothetical protein	phage integrase site specific recombinase	molybdopterin-binding oxidoreductase	LysE family translocator protein	oxidoreductase, FAD-binding	hypothetical protein	benzaldehyde lyase	tricarboxylate transporter family protein	hypothetical protein	tricarboxylate transporter family protein	LysR family transcriptional regulator
1.61	1.29	1.21	-1.15	1.62	1.43	1.31	1.22	1.07	1.10	1.95	1.64	-1.77	x	-1.39	Х	x	1.39	-1.40	27.60	2.26	12.90	5.59	-0.98	2.56	2.21	х	х	2.32	х	-1.79	х	-1.02	х	-1.61	-2.49	-1.37
0.12	0.27	0.36	0.73	0.28	0.07	0.01	0.21	0.82	0.20	0.21	0.04	0.78	×	0.01	х	x	0.37	0.64	0.00	0.01	0.08	0.04	0.18	0.11	0.60	х	х	0.12	×	x	x	0.93	Х	Х	0.04	0.76
-1.01	1.05	-1.47	-1.34	-1.48	-1.24	-1.25	-1.47	-1.44	-1.25	1.03	-1.10	-1.09	×	-1.27	х	x	1.51	-1.10	-1.36	-1.35	-1.17	-1.11	-1.46	-1.25	-1.33	х	-1.62	1.02	-1.35	-1.13	-1.23	-1.48	-1.03	-1.61	-1.90	-1.46
0.79	0.73	0.04	0.82	0.02	0.28	0.36	0.08	0.53	0.76	0.75	0.65	0.97	×	0.43	x	×	0.14	0.67	0.72	0.24	0.76	0.67	0.48	0.19	0.84	×	×	0.95	0.71	0.90	0.08	0.43	0.95	0.25	0.03	0.68
0.30	1.17	-0.13	-1.25	0.07	0.10	0.03	-0.13	-0.19	-0.08	1.49	0.27	-1.43	N/A	-1.33	N/A	N/A	1.45	-1.25	13.12	0.46	5.87	2.24	-1.22	0.66	0.44	N/A	N/A	1.67	N/A	-1.46	N/A	-1.25	N/A	-1.61	-2.20	-1.42
1.31	0.12	1.34	0.10	1.55	1.34	1.28	1.35	1.26	1.18	0.46	1.37	0.34	N/A	0.06	N/A	N/A	0.06	0.15	14.48	1.81	7.04	3.35	0.24	1.91	1.77	N/A	N/A	0.65	N/A	0.33	N/A	0.23	N/A	0.00	0.29	0.04
-1.46	-1.83	-1.39	-2.63	-2.33	-2.66	-2.67	-0.98	-1.06	-1.44	-1.48	-1.37	-2.57	х	-1.38	х	x	1.85	-1.23	-1.93	-7.07	-1.19	-1.43	-3.38	-1.22	-1.65	х	-2.51	1.40	-1.34	-1.63	-1.15	-1.17	-2.38	-1.36	-4.62	-2.09
0.07	0.10	0.28	0.26	0.04	0.02	0.01	0.24	0.55	0.45	0.06	0.12	0.55	×	0.09	х	×	0.11	0.67	0.16	0.01	0.98	0.86	0.01	0.49	0.80	x	x	0.17	×	0.67	0.51	0.82	0.05	0.62	0.02	0.55
-0.98	1.78	1.44	1.24	1.54	1.29	-0.98	1.68	1.29	1.19	1.15	1.47	1.16	Х	-1.12	Х	X	1.47	1.43	1.19	1.26	1.28	-1.04	-1.02	Х	1.30	Х	-1.12	1.23	Х	-1.07	-1.04	-1.23	Х	-1.41	-1.22	-1.16
0.59	0.08	0.09	0.87	0.03	0.36	0.90	0.17	0.68	0.71	0.29	0.07	0.59	×	0.06	х	×	0.15	0.30	х	0.54	0.67	1.00	0.98	×	0.86	×	x	0.30	×	0.23	0.99	0.59	x	0.06	0.42	0.83
-1.22	-0.03	0.03	-0.70	-0.40	-0.69	-1.83	0.35	0.12	-0.13	-0.17	0.05	-0.71	N/A	-1.25	N/A	N/A	1.66	0.10	-0.37	-2.91	0.05	-1.24	-2.20	N/A	-0.18	N/A	-1.82	1.32	N/A	-1.35	-1.10	-1.20	N/A	-1.39	-2.92	-1.63
0.24	1.81	1.42	1.94	1.94	1.98	0.84	1.33	1.18	1.32	1.32	1.42	1.87	N/A	0.13	N/A	N/A	0.19	1.33	1.56	4.17	1.24	0.20	1.18	N/A	1.48	N/A	0.70	0.09	N/A	0.28	0.05	0.03	N/A	0.02	1.70	0.47
-1.30	-1.41	-1.53	-1.08	-1.04	-1.04	1.49	-1.46	-1.06	-1.83	1.58	-1.00	-1.24	x	-1.09	x	×	-1.30	1.26	35.20	18.80	21.00	10.50	3.46	7.32	5.79	x	-1.12	-1.25	1.06	-1.21	-1.08	-1.22	-1.35	1.72	1.19	-1.13
0.23	0.12	0.10	0.95	0.94	0.91	0.32	0.03	0.86	0.16	0.01	0.87	0.92	×	0.45	х	×	0.06	0.39	0.02	0.01	0.02	0.02	0.02	0.00	0.13	x	0.89	0.31	×	0.71	0.56	0.60	0.17	0.04	0.34	0.87
-1.48	-1.43	-1.23	1.38	-1.29	1.13	1.44	-1.43	-0.96	-2.59	1.99	-1.21	2.03	х	-1.45	Х	x	-1.70	-0.97	125.00	44.90	21.30	23.30	3.46	8.63	8.82	Х	Х	-1.04	Х	1.02	х	-1.18	Х	-1.14	1.48	1.51
0.42	0.12	0.30	0.81	0.59	0.32	0.24	Х	0.50	x	0.02	0.61	Х	×	0.23	х	×	0.01	0.79	0.01	0.00	0.03	0.00	0.01	0.01	0.12	x	x	0.99	×	x	×	0.88	x	1.00	0.07	0.69
-1.39	-1.42	-1.38	0.15	-1.17	0.04	1.47	-1.45	-1.01	-2.21	1.79	-1.10	0.40	N/A	-1.27	N/A	N/A	-1.50	0.14	80.10	31.85	21.15	16.90	3.46	7.98	7.31	N/A	N/A	-1.15	N/A	-0.10	N/A	-1.20	N/A	0.29	1.34	0.19
0.09	0.01	0.15	1.23	0.13	1.09	0.03	0.02	0.05	0.38	0.20	0.11	1.64	N/A	0.18	N/A	N/A	0.20	1.12	44.90	13.05	0.15	6.40	0.00	0.66	1.52	N/A	N/A	0.11	N/A	1.12	N/A	0.02	N/A	1.43	0.14	1.32

SPO2457	SPO2456	SPO2455	SPO2454	SPO2453	SPO2452	SPO2451	SPO2450	SPO2449	SPO2448	SPO2447	SPO2446	SPO2445	SPO2444	SPO2443	SPO2442	SPO2441	SPO2440	SPO2439	SPO2438	SPO2437	SPO2436	SPO2435	SPO2434	SPO2433	SPO2432	SPO2431	SPO2430	SPO2429	SPO2428	SPO2427	SPO2426	SPO2425	SPO2424	SPO2423	SPO2422	SPO2421
pdxA				pepA									ndk																			aroE	idnD			
4-пуцгохуппеошпе-4-рнозрпате цепуцгоденахе (ЕС:1.1.1.262)	pepudyi-prolyi cis-trans isomerase domain- containing protein	organic solvent tolerance protein	YjgP/YjgQ family permease	leucyl aminopeptidase (EC:3.4.11.1)	DNA polymerase III subunit chi	acyltransferase	Asp/Glu/Hydantoin racemase	hypothetical protein	multiple antibiotic resistance protein MarC	acetyltransferase	ABC transporter ATP-binding protein	Bcr/CfIA subfamily drug resistance transporter	nucleoside diphosphate kinase (EC:2.7.4.6)	giyeine belaine/pronne ABC transporter A1r- binding protein	glycine betaine/proline ABC transporter permease	glycine betaine/L-proline ABC transporter substrate-binding protein	M24 family metallopeptidase	GntR family transcriptional regulator	phytanoyl-CoA dioxygenase	Asp/Glu/Hydantoin racemase	FAD-dependent oxidoreductase	fumarylacetoacetate hydrolase	GntR family transcriptional regulator	TRAP dicarboxylate transporter subunit DctP	TRAP dicarboxylate transporter subunit DctQ	TRAP dicarboxylate transporter subunit DctM	UxaA family hydrolase	Gfo/Idh/MocA family oxidoreductase	6-phosphogluconate dehydrogenase	oxidoreductase	hypothetical protein	shikimate 5-dehydrogenase (EC:1.1.1.25)	L-idonate 5-dehydrogenase (EC:1.1.1.264)	hypothetical protein	D-isomer specific 2-hydroxyacid dehydrogenase	hypothetical protein
1.38	1.33	1.30	-0.99	1.45	1.16	1.92	2.75	-1.69	1.18	-1.04	Х	-1.30	1.04	х	-1.97	-1.95	-1.29	-1.73	2.43	1.17	1.05	1.95	1.21	Х	1.92	1.18	Х	Х	1.40	1.00	х	-1.52	1.09	1.20	-1.24	-1.15
0.40	0.03	0.73	0.97	0.11	0.95	0.08	0.00	0.01	0.98	1.00	×	0.57	0.81	×	0.03	0.03	0.72	0.19	0.05	x	х	0.02	0.03	×	0.41	0.65	×	×	0.83	0.91	×	×	0.48	0.12	0.54	0.18
1.30	-1.11	1.00	1.23	-1.37	1.13	-1.10	1.82	-1.53	1.02	1.15	1.14	1.10	0.99	-1.53	-1.63	-2.87	-1.29	-1.31	-1.19	-1.05	-1.03	-1.42	-1.36	х	-2.10	-1.01	-1.59	-1.46	-1.36	-1.36	-1.08	-1.23	-1.13	-1.33	-1.42	-1.12
0.09	0.46	0.87	0.03	0.09	0.47	0.27	0.08	0.02	1.00	0.63	0.46	0.84	0.69	0.20	0.05	0.01	0.06	0.47	0.17	0.95	0.99	0.13	0.22	×	0.04	0.93	0.76	х	0.47	0.37	0.53	0.62	0.50	0.25	0.33	0.42
1.34	0.11	1.15	0.12	0.04	1.15	0.41	2.29	-1.61	1.10	0.05	N/A	-0.10	1.01	N/A	-1.80	-2.41	-1.29	-1.52	0.62	0.06	0.01	0.27	-0.08	N/A	-0.09	0.09	N/A	N/A	0.02	-0.18	N/A	-1.38	-0.02	-0.07	-1.33	-1.14
0.04	1.22	0.15	1.11	1.41	0.02	1.51	0.47	0.08	0.08	1.10	N/A	1.20	0.03	N/A	0.17	0.46	0.00	0.21	1.81	1.11	1.04	1.69	1.29	N/A	2.01	1.10	N/A	N/A	1.38	1.18	N/A	0.14	1.11	1.27	0.09	0.01
1.35	1.70	1.33	2.56	1.85	1.16	2.71	3.20	-1.21	1.79	4.00	7.06	-1.56	6.08	1.89	3.82	-1.71	-2.08	-1.70	-2.37	-1.92	-1.06	2.45	1.39	-1.10	-1.25	-1.68	Х	-0.92	-1.81	-3.06	-1.15	-1.93	1.90	-1.38	-1.48	-2.85
0.10	0.03	0.67	0.02	0.07	0.92	0.03	0.02	0.17	0.83	0.08	0.00	0.63	0.01	0.04	0.00	0.02	0.08	0.08	0.04	0.74	0.99	0.01	0.02	х	0.77	0.25	х	0.11	0.69	0.24	0.26	0.11	0.02	0.16	0.41	0.00
1.18	0.96	1.21	1.02	1.83	1.14	-1.48	-1.12	-1.11	-1.49	-1.12	-1.45	-1.37	-1.28	-1.24	-1.24	-1.96	-1.20	-1.38	1.09	1.13	-1.44	1.66	1.12	-1.01	1.54	1.58	1.12	1.07	1.45	1.33	1.06	1.07	1.18	1.29	1.04	1.13
0.19	0.17	0.52	0.75	0.04	0.78	0.26	0.71	0.41	0.41	0.62	0.02	0.63	0.10	0.28	0.41	0.02	0.49	0.06	0.65	0.88	0.51	0.03	0.55	Х	0.03	0.17	0.86	Х	0.21	0.24	0.82	0.70	0.33	0.19	0.90	0.67
1.27	1.33	1.27	1.79	1.84	1.15	0.62	1.04	-1.16	0.15	1.44	2.81	-1.47	2.40	0.33	1.29	-1.84	-1.64	-1.54	-0.64	-0.40	-1.25	2.06	1.26	-1.06	0.15	-0.05	N/A	0.08	-0.18	-0.87	-0.04	-0.43	1.54	-0.04	-0.22	-0.86
0.09	0.37	0.06	0.77	0.01	0.01	2.10	2.16	0.05	1.64	2.56	4.26	0.10	3.68	1.57	2.53	0.13	0.44	0.16	1.73	1.53	0.19	0.39	0.14	0.05	1.40	1.63	N/A	0.99	1.63	2.20	1.11	1.50	0.36	1.34	1.26	1.99
-1.15	-1.22	-1.39	1.17	1.44	2.22	-1.25	1.19	1.10	-1.19	-1.55	1.02	-1.20	-1.60	-1.49	-1.21	-1.05	-1.39	-1.53	-1.32	-1.34	1.13	-1.58	-2.22	х	-1.25	-1.12	-1.14	-1.06	1.23	-1.01	-1.18	-1.21	-1.44	-1.73	-1.37	-1.14
0.35	0.02	0.26	0.12	0.16	0.09	0.24	0.05	0.62	0.66	0.14	0.98	0.67	0.00	0.02	0.39	0.77	0.03	0.01	0.38	0.64	0.71	0.01	0.03	×	0.25	0.66	х	0.96	0.40	0.99	0.14	0.51	0.01	0.05	0.34	0.74
-1.42	1.20	-1.17	-1.17	1.74	1.73	1.24	-1.08	1.30	-1.23	-2.27	-2.01	-1.15	-2.16	-1.13	1.13	-1.19	-1.07	-1.55	-1.26	-1.67	-1.05	-1.15	-1.89	x	-1.38	-1.45	1.90	1.24	1.14	1.13	1.29	1.45	-1.25	-2.21	1.26	1.17
0.01	0.65	0.21	0.15	0.01	0.12	0.25	0.90	0.18	0.67	0.08	0.03	0.94	0.01	0.91	0.46	0.20	0.99	0.17	0.49	х	Х	0.16	0.02	х	0.08	0.44	Х	Х	0.60	0.63	0.20	0.36	0.49	0.13	0.35	0.79
-1.29	-0.01	-1.28	0.00	1.59	1.98	-0.01	0.05	1.20	-1.21	-1.91	-0.50	-1.18	-1.88	-1.31	-0.04	-1.12	-1.23	-1.54	-1.29	-1.51	0.04	-1.37	-2.06	N/A	-1.32	-1.29	0.38	0.09	1.19	0.06	0.06	0.12	-1.35	-1.97	-0.06	0.02
0.14	1.21	0.11	1.17	0.15	0.25	1.25	1.14	0.10	0.02	0.36	1.52	0.03	0.28	0.18	1.17	0.07	0.16		0.03	_	1.09	0.22	0.17	N/A	0.06	0.16	1.52	1.15	0.05	1.07	1.24	1.33	0.10	0.24	1.32	1.16

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### Chapter 10: Appendix

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MerR family transcriptional regulator	integration host factor subunit alpha	3-oxoacyl-ACP synthase (EC:2.3.1.41)	glycerol-3-phosphate acyltransferase PlsX	50S ribosomal protein L32	hypothetical protein	SmpA/OmlA family lipoprotein	hypothetical protein	ABC transporter ATP-binding protein	3-hydroxyacyl-CoA dehydrogenase (EC:1.1.1.35)	hypothetical protein	hypothetical protein	tyrosyl-tRNA synthetase (EC:6.1.1.1)	anhydro-N-acetylmuramic acid kinase	hypothetical protein	drug/metabolite exporter family protein	hypothetical protein	hypothetical protein	FUR family transcriptional regulator	lipoprotein	hypothetical protein	phosphopyruvate hydratase (EC:4.2.1.11)	haloacid dehalogenase	hypothetical protein	acetyltransferase	познистилине ристентив пистеозме пучновае (EC:3.2.2.1)	(EC:3.6.1.19) (EC:3.6.1.19)	amidohydrolase	agmatinase (EC:3.5.3.11)	glyoxalase	hypothetical protein	agmatinase (EC:3.5.3.11)	hypothetical protein	hypothetical protein	peptide chain release factor 1	HemK family modification methylase	hypothetical protein	dimethyladenosine transferase (EC:2.1.1)
1.02	-1.43	-1.62	1.04	1.37	-1.06	1.35	-1.25	-1.28	-1.08	-1.01	1.34	-1.60	-1.23	-1.11	-1.61	-1.28	1.15	-1.14	1.07	Х	2.36	-1.16	Х	1.45	1.60	1.83	2.30	2.53	-3.01	-1.70	1.62	x	-1.50	-1.07	Х	-1.29	1.27
0.70	0.01	0.03	0.96	0.55	0.93	0.52	0.68	0.62	0.91	x	0.55	0.64	0.34	0.42	0.73	0.17	0.14	0.67	0.92	x	0.00	0.60	×	0.04	0.04	0.03	0.01	0.25	0.00	0.07	0.19	×	0.08	0.91	×	0.13	0.96
-1.49	-1.11	0.98	-1.06	-1.16	-1.13	0.99	-1.15	1.06	-1.31	1.12	1.11	1.08	1.15	1.33	-1.04	-2.04	1.23	1.43	1.16	1.70	1.30	1.61	-1.67	1.30	1.02	1.00	1.29	1.24	-1.37	-1.10	1.39	1.08	1.08	1.35	1.73	-1.78	1.42
0.11	0.53	0.49	0.33	0.04	0.87	0.90	0.83	0.80	0.68	0.62	0.74	0.84	0.30	0.46	0.92	0.11	0.49	0.17	0.50	0.01	0.42	0.00	х	0.40	1.00	0.97	0.12	0.45	0.01	0.76	0.22	0.67	0.46	0.04	0.02	0.04	0.63
-0.24	-1.27	-0.32	-0.01	0.11	-1.10	1.17	-1.20	-0.11	-1.20	0.06	1.23	-0.26	-0.04	0.11	-1.33	-1.66	1.19	0.15	1.12	N/A	1.83	0.23	N/A	1.38	1.31	1.42	1.80	1.89	-2.19	-1.40	1.51	N/A	-0.21	0.14	N/A	-1.54	1.35
1.26	0.16	1.30	1.05	1.27	0.03	0.18	0.05	1.17	0.12	1.07	0.12	1.34	1.19	1.22	0.28	0.38	0.04	1.29	0.04	N/A	0.53	1.39	N/A	0.08	0.29	0.42	0.51	0.65	0.82	0.30	0.12	N/A	1.29	1.21	N/A	0.25	0.08
-2.02	-2.72	2.79	3.01	6.90	1.54	1.98	0.89	1.25	-1.21	-1.40	1.50	0.95	-1.26	1.54	-1.59	1.57	-0.95	1.15	-3.96	7.51	5.63	1.68	х	1.57	1.71	2.79	1.65	2.22	-1.74	1.19	2.22	2.64	-1.79	1.75	2.64	2.24	2.07
0.13	0.00	0.01	0.12	0.00	0.03	0.13	0.80	0.47	0.79	×	0.46	0.85	0.44	0.03	0.80	0.17	0.24	0.71	0.01	0.00	0.00	0.03	x	0.08	0.04	0.02	0.01	0.04	0.06	0.16	0.04	0.01	0.06	0.13	0.01	0.04	0.78
1.17	1.23	-1.11	-1.53	-1.63	-1.20	1.65	1.32	-1.50	-1.00	1.25	-1.14	-1.19	-1.07	1.21	1.02	1.08	1.38	-1.10	1.64	-1.61	-1.20	-1.49	x	1.24	1.44	-1.06	1.17	1.04	1.10	-1.35	1.10	1.17	-1.10	-1.69	-1.99	-1.16	-1.20
0.74	0.12	0.20	0.26	0.01	0.14	0.00	0.76	0.00	0.99	×	0.81	0.27	0.28	0.06	0.93	0.81	0.19	0.26	0.01	0.02	0.05	0.10	x	0.15	0.24	0.61	0.22	0.96	0.10	0.15	0.56	0.45	0.77	0.02	0.02	0.32	0.75
-0.43	-0.75	0.84	0.74	2.64	0.17	1.82	1.11	-0.13	-1.10	-0.08	0.18	-0.12	-1.17	1.38	-0.29	1.33	0.21	0.02	-1.16	2.95	2.22	0.10	N/A	1.41	1.58	0.87	1.41	1.63	-0.32	-0.08	1.66	1.91	-1.45	0.03	0.33	0.54	0.44
1.60	1.98	1.95	2.27	4.27	1.37	0.17	0.21	1.38	0.11	1.33	1.32	1.07	0.10	0.17	1.31	0.24	1.17	1.13	2.80	4.56	3.42	1.59	N/A	0.17	0.14	1.93	0.24	0.59	1.42	1.27	0.56	0.74	0.35	1.72	2.32	1.70	1.64
-1.12	3 1.21	1.19	0.99	-1.30	-2.45	-1.24	1.81	-1.55	1.59	-1.11	1.05	-1.28	) 1.15		1.28	-1.23	-1.20	3 1.27	) 1.27	1.08	1.03	1.45	-1.50	-1.10	-1.63	-1.66	-1.08	-1.03	1.08	1.39	-1.08	-0.99	-1.00		1.23		1.08
2 0.93	0.24	€ 0.44	9 0.61	0 0.19	5 0.07	4 0.05	0.35	5 0.06	9 0.53	1 0.94	5 0.59	8 0.30	5 0.68	7 0.14	3 0.31	3 0.11	0 0.84	7 0.10	7 0.09	3 0.18	3 0.80	5 0.06	0 0.41	0 0.52	3 0.07	6 0.01	_	3 0.81	3 0.84	€ 0.30	8 0.38	9 0.79	0 0.89		3 0.17	6 0.05	3 0.93
3 1.56	1.56	4 -1.46	-1.25	-1.54	-2.41	0.90	2.36	-2.03	3 1.13	4 -1.16	) 1.54	-1.36	3 1.64	-1.52	1.12	1.20	4 -1.30	-1.25	9 1.09	-1.55	0.96	1.01	×	-1.33	-1.41	-1.74	-1.32	0.97	4 1.04	-1.23	-1.39	-1.33	×		-2.43	-1.35	-1.14
6 0.11	6 0.05	6 0.04	5 0.08	4 0.07	1 0.03	0 0.21	6 0.32	3 0.05	3 0.79	9 9	4 0.13	6 0.30	4 0.16	2 0.03	2 0.82	0 0.57	0 0.37	5 0.29	9 0.22	5 0.04	6 0.25	0.99	x	3 0.13	0.08	4 0.03		7 0.54	4 0.90	3 X	9 0.05	3 0.28	x		3 0.00	5 0.06	4 0.87
1 0.22	5 1.39	4 -0.14	8 -0.13	7 -1.42	3 -2.43	1 -0.17	2 2.09	5 -1.79	9 1.36	-1.14	3 1.30	0 -1.32	6 1.40	3 -1.40	2 1.20	7 -0.02	7 -1.25	9 0.01	2 1.18	4 -0.24	5 0.99	9 1.23	N/A	-1.22	8 -1.52	3 -1.70	5 -1.20	4 -0.03	0 1.06	0.08	5 -1.24	8 -1.16	N/A		0 -0.60	6 -1.66	7 -0.03
2 1.34	9 0.18	4 1.33	3 1.12	.2 0.12	-3 0.02	7 1.07	9 0.27	0.24	6 0.23	4 0.02	0 0.25	0.04	0 0.24	0 0.13	0 0.08	1.22	.5 0.05	1 1.26	8 0.09	4 1.32	9 0.04	3 0.22	A N/A	2 0.12	0.11	0 0.04	0 0.12	1.00	6 0.02	8 1.31	0.16	6 0.17	A N/A		1.83	0.30	1.11
4	8	33	12	12	ŭ	7(	75	34	33	ŭ	ઝ	¥	34	13	8	22	ઝ	36	9	32	¥	22	A	12	1	¥	12	ŏ	ŭ	31	16	17	A	\$7	ü	õ	=

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						nylA				glnE													nahG	xth					argS			scpA	scpB		
branched-chain amino acto ABC transporter A1F- binding protein	binding protein	oranoned-onain amino acid ABC transporter	orancined-chain amino acid ABC transporter	hypothetical protein	AMP-binding protein (EC:2.3.1.86)	6-aminohexanoate-cyclic-dimer hydrolase (EC:3.5.2.12)	hypothetical protein	hypothetical protein	prolyl-tRNA synthetase	glutamate-ammonia ligase adenylyltransferase (EC:2.7.7.42)	hypothetical protein	lipoprotein	hypothetical protein	hypothetical protein	luciferase	M48 family peptidase	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	ATPase AAA	dnaK suppressor protein	salicylate hydroxylase (EC:1.14.13.1)	exodeoxyribonuclease III (EC:3.1.11.2)	hypothetical protein	hypothetical protein	Iron-sulfur cluster assembly family protein	deoxyguanos in etriphosphate triphosphohydrolase	arginyl-tRNA synthetase (EC:6.1.1.19)	hypothetical protein	beta-N-acetylhexosaminidase	segregation and condensation protein A	segregation and condensation protein B	hypothetical protein	2'-deoxycytidine 5'-triphosphate deaminase (EC:3.5.4.13)
-1.17	-1.21	-1.16	-0.99	-0.96	1.29	1.27	х	1.37	-1.32	7.33	9.39	2.40	-1.11	Х	-1.18	-1.63	-5.28	-1.07	-1.22	-1.57	1.80	1.50	1.22	1.03	1.35	1.17	-1.10	-1.20	1.54	-1.49	-1.16	-1.49	1.01	1.12	-1.16
x	0.47	0.44	0.97	0.70	0.19	0.67	х	0.69	0.47	0.04	0.00	0.67	0.26	×	0.53	0.24	0.00	0.66	0.54	0.04	0.05	0.16	0.93	0.60	0.79	0.32	0.38	0.27	0.73	0.01	0.76	0.00	0.94	0.51	0.48
Х	-1.03	-1.40	1.03	-1.14	-1.02	1.22	Х	1.34	-1.08	2.80	2.35	1.24	1.26	-1.14	-1.48	-1.90	-2.43	1.17	1.24	1.22	1.04	1.03	2.70	1.06	4.72	1.68	1.20	-1.34	1.05	-1.54	-1.26	-1.84	-1.60	-1.75	1.34
x	0.97	0.09	0.85	0.86	0.97	0.50	x	0.53	0.79	0.04	0.05	0.10	0.36	0.30	0.06	0.14	0.06	0.59	0.28	0.27	0.96	0.99	0.03	0.89	0.02	0.09	0.56	0.40	0.85	0.07	0.52	0.02	0.46	0.12	0.03
N/A	-1.12	-1.28	0.02	-1.05	0.14	1.25	N/A	1.36	-1.20	5.07	5.87	1.82	0.08	N/A	-1.33	-1.77	-3.86	0.05	0.01	-0.18	1.42	1.27	1.96	1.05	3.04	1.43	0.05	-1.27	1.30	-1.52	-1.21	-1.67	-0.30	-0.32	0.09
N/A	0.09	0.12	1.01	0.09	1.16	0.03	N/A	0.02	0.12	2.27	3.52	0.58	1.19	N/A	0.15	0.14	1.43	1.12	1.23	1.40	0.38	0.24	0.74	0.02	1.69	0.26	1.15	0.07	0.25	0.03	0.05	0.18	1.31	1.44	1.25
x	-2.13	-2.49	-1.30	-2.46	-3.66	-1.92	х	1.91	-1.86	4.14	1.56	-1.70	1.48	3.65	-1.37	-1.18	5.33	1.50	1.46	-2.96	1.72	1.94	1.54	1.45	-3.84	-4.17	1.11	1.59	3.81	1.05	-1.23	-1.40	1.09	1.03	1.32
х	0.17	0.08	0.87	0.02	0.00	0.16	Х	0.02	0.04	0.07	0.16	0.04	0.08	0.00	0.26	0.56	0.00	0.32	0.16	0.01	0.03	0.04	0.73	0.00	0.00	0.01	0.69	0.02	0.29	0.02	0.75	0.02	0.96	0.38	0.19
x	-1.17	-1.13	-1.20	-1.64	-1.05	1.08	X	1.35	-1.24	1.23	1.35	1.17	-1.45	-1.01	-1.43	1.21	-1.15	-1.01	-1.28	1.65	1.06	1.24	-1.12	1.31	17.70	5.26	0.98	1.25	-1.50	-1.43	-1.11	1.10	1.03	1.16	-1.17
х	x	0.64	0.26	0.19	0.83	0.78	Х	0.48	0.54	0.60	0.17	0.41	0.17	0.80	0.17	0.69	0.10	0.95	0.12	0.01	0.96	0.17	0.61	0.33	0.00	0.00	0.53	0.20	0.20	0.01	0.81	0.66	0.96	0.25	0.28
N/A	-1.65	-1.81	-1.25	-2.05	-2.36	-0.42	N/A	1.63	-1.55	2.69	1.46	-0.27	0.02	1.32	-1.40	0.02	2.09	0.25	0.09	-0.66	1.39	1.59	0.21	1.38	6.93	0.55	1.05	1.42	1.16	-0.19	-1.17	-0.15	1.06	1.10	0.08
N/A	0.48	0.68	0.05	0.41	1.31	1.50	N/A	0.28	0.31	1.46	0.11	1.44	1.47	2.33	0.03	1.20	3.24	1.26	1.37	2.31	0.33	0.35	1.33	0.07	10.77	4.72	0.06	0.17	2.66	1.24	0.06	1.25	0.03	0.06	1.25
-1.17	×	-1.21	1.07	-1.28	-1.77	-0.97	-1.02	1.62	1.03	1.10	-1.03	-1.54	1.08	1.02	-1.02	1.24	2.48	1.26	-1.26	-1.57	-1.43	0.97	-1.13	1.07	1.55	1.12	2.08	-1.13	-1.33	1.12	1.60	-1.54	1.43	1.02	-1.24
Х	×	0.30	0.81	0.25	0.08	0.84	Х	0.04	0.91	0.82	0.73	0.09	0.70	0.75	0.70	0.72	0.04	0.67	0.13	0.01	0.04	0.37	0.70	0.79	0.29	0.27	0.03	0.58	0.19	0.72	0.20	0.11	0.60	0.51	0.09
x	x	×	1.36	-1.52	-1.88	-1.32	x	1.83	2.21	-1.26	1.05	1.04	-1.53	-1.79	1.36	1.81	2.04	-1.41	-1.12	-1.56	-1.16	1.09	-1.14	0.99	6.59	1.98	1.75	-1.12	-1.71	1.24	1.00	-1.48	1.11	1.35	-1.29
х	x	x	0.09	Х	0.16	0.22	Х	0.37	0.09	0.63	0.54	0.07	0.02	0.01	0.25	0.14	0.08	0.34	0.18	0.00	0.31	0.77	0.87	0.86	0.01	0.02	0.03	0.13	0.14	0.30	0.96	0.06	0.97	0.22	0.19
N/A	N/A	N/A	1.22	-1.40	-1.83	-1.14	N/A	1.73	1.62	-0.08	0.01	-0.25	-0.23	-0.39	0.17	1.53	2.26	-0.08	-1.19	-1.57	-1.30	1.03	-1.14	1.03	4.07	1.55	1.92	-1.13	-1.52	1.18	1.30	-1.51	1.27	1.19	-1.27
N/A	N/A	N/A	0.15	0.12	0.05	0.18	N/A	0.11	0.59	1.18	1.04	1.29	1.31	1.41	1.19	0.29	0.22	1.34	0.07	0.01	0.14	0.06	0.01	0.04	2.52	0.43	0.17	0.00	0.19	0.06	0.30	0.03	0.16	0.17	0.03

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# Chapter 10: Appendix

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SPO2571	SPO2570	SPO2569	SPO2568	SPO2567	SPO2566	SPO2565	SPO2564	SPO2563	SPO2562	SPO2561	SPO2560	SPO2559	SPO2556	SPO2555	SPO2554	SPO2553	SPO2552	SPO2551	SPO2550	SPO2549	SPO2548	SPO2547	SPO2546	SPO2545	SPO2544	SPO2543	SPO2542	SPO2541	SPO2540	SPO2539	SPO2538	SPO2537	SPO2536	SPO2535	SPO2534
		ate							fucA		garR	gnl																							
DetM	glyoxalase TR AP C4-dicarbox vlate transport system permease	arginyl-tRNA-protein transferase (EC:2.3.2.8)	RDD family protein	hypothetical protein	glyoxalase	ribosome-associated GTPase	hypothetical protein	hydroxypyruvate isomerase	aldolase (EC:4.1.2.17)	hypothetical protein	2-hydroxy-3-oxopropionate reductase (EC:1.1.1.60)	gluconolactonase (EC:3.1.1.17)	allantoate amidohydrolase	hypothetical protein	peptide/opine/nickel uptake ABC transporter substrate-binding protein	peptide/opine/nickel uptake ABC transporter permease	peptude/opine/mckel ABC transporter permease/ATP-binding protein	binding protein	GntR family transcriptional regulator	phosphotransferase	zinc-binding dehydrogenase oxidoreductase	TRAP dicarboxylate transporter subunit DctM	TRAP dicarboxylate transporter subunit DctQ	TRAP dicarboxylate transporter subunit DctP	amidohydrolase	GntR family transcriptional regulator	biotin/lipoate binding domain-containing protein	propionyl-CoA carboxylase	carbamoyl-phosphate synthase	AMP-binding protein	acyl-CoA dehydrogenase	hypothetical protein	LuxR family transcriptional regulator	histone deacetylase/AcuC/AphA family protein	branched-chain amino acid ABC transporter substrate-binding protein
2.39	0.99	-1.31	1.29	0.92	-1.72	-1.11	х	-1.23	-1.22	-1.39	-1.04	-1.47	-1.97	×	-2.31	1.07	×	-1.08	х	1.31	-2.68	-1.56	-1.73	-1.44	-1.01	1.17	-3.45	-2.13	-1.39	-1.45	-1.53	-1.62	-1.41	-1.36	-2.37
0.16	0.63	0.57	0.18	0.02	0.01	0.92	×	0.06	0.06	0.03	0.39	0.44	x	×	0.26	1.00	x	0.96	x	x	0.13	0.09	0.25	0.24	0.99	0.81	0.01	0.00	0.06	0.17	0.07	0.05	0.28	0.44	0.05
-1.17	-1.04	1.02	-1.07	-1.48	1.52	1.09	-1.22	-1.13	1.02	-1.00	-1.10	-1.09	-1.66	-1.02	-1.18	-1.01	1.22	-1.04	-1.15	-1.14	-1.29	-1.05	-1.28	-1.23	1.09	1.03	-1.25	-1.15	-1.25	-1.10	1.02	-1.25	-1.15	-1.11	-1.20
0.34	0.83	0.93	0.38	0.10	0.02	0.87	0.79	0.33	0.65	0.98	0.25	0.78	0.67	0.98	0.90	1.00	0.30	0.98	0.31	0.71	0.03	0.82	0.44	0.18	0.93	0.66	0.16	0.62	0.07	0.37	0.87	0.48	0.58	0.90	0.47
0.61	-0.03	-0.15	0.11	-0.28	-0.10	-0.01	N/A	-1.18	-0.10	-1.20	-1.07	-1.28	-1.82	N/A	-1.75	0.03	N/A	-1.06	N/A	0.09	-1.99	-1.31	-1.51	-1.34	0.04	1.10	-2.35	-1.64	-1.32	-1.28	-0.26	-1.44	-1.28	-1.24	-1.79
1.78	1.01	1.17	1.18	1.20	1.62	1.10	N/A	0.05	1.12	0.20	0.03	0.19	0.16	N/A	0.57	1.04	N/A	0.02	N/A	1.23	0.70	0.25	0.23	0.11	1.05	0.07	1.10	0.49	0.07	0.18	1.28	0.19	0.13	0.13	0.59
3.47	1.07	-1.30	2.05	1.24	1.41	-1.32	-1.37	-1.52	-1.42	-1.31	1.06	-1.34	-3.30	-1.43	-3.16	1.11	-1.11	1.16	2.68	-1.47	-3.15	-2.40	-3.52	-1.86	-1.42	1.28	-7.01	-3.30	-1.87	-1.94	-1.79	-1.96	-2.05	-11.10	-7.94
0.05	0.62	0.53	0.01	0.53	0.03	0.85	0.90	0.08	0.06	0.08	0.41	0.68	×	0.88	0.26	0.96	0.17	0.90	0.03	0.80	0.00	0.41	0.11	0.61	0.93	0.19	0.00	0.01	0.03	0.03	0.63	0.06	0.15	0.04	0.20
1.94	1.15	1.28	-1.20	1.37	-1.16	-1.98	-1.11	2.11	1.88	2.06	1.98	1.89	-1.07	1.17	-1.30	-1.00	-1.12	1.16	1.06	-1.41	-1.16	-1.20	-0.95	-1.47	-1.20	1.05	-1.02	-1.54	-1.26	-1.17	-1.01	1.14	х	-1.32	-1.01
0.03	0.65	0.11	0.33	0.13	0.05	0.30	0.72	0.02	0.01	0.05	0.01	0.12	×	0.31	0.83	0.99	х	0.93	0.68	0.17	0.43	x	0.87	0.07	0.89	0.72	0.98	0.16	0.38	0.31	0.99	0.63	x	0.76	0.98
2.71	1.11	-0.01	0.43	1.31	0.13	-1.65	-1.24	0.30	0.23	0.38	1.52	0.28	-2.19	-0.13	-2.23	0.06	-1.12	1.16	1.87	-1.44	-2.16	-1.80	-2.24	-1.67	-1.31	1.17	-4.02	-2.42	-1.57	-1.56	-1.40	-0.41	N/A	-6.21	-4.48
0.77	0.04	1.29	1.63	0.07	1.29	0.33	0.13	1.82	1.65	1.69	0.46	1.62	1.12	1.30	0.93	1.05	0.01	0.00	0.81	0.03	1.00	0.60	1.28	0.19	0.11	0.12	3.00	0.88	0.31	0.39	0.39	1.55	N/A	4.89	3.47
1.40	1.54	1.32	1.54	-1.14	-1.39	-1.59	-1.15	1.04	1.08	1.30	1.18	1.20	-1.58	-1.36	1.19	-1.02	-0.97	-1.19	-1.58	-1.37	-1.28	1.38	-1.32	1.04	1.18	1.45	-1.29	-1.15	-1.12	-1.11	-1.33	-1.54	-1.31	-1.05	-1.14
0.37	0.07	0.11	0.12	0.07	0.06	0.44	0.73	0.85	0.52	0.33	0.10	0.32	0.68	0.33	0.88	1.00	0.62	0.91	0.02	0.16	0.03	0.62	0.40	0.77	0.86	0.01	0.16	0.49	0.61	0.44	0.68	0.14	0.07	0.98	0.72
1.89	1.03	1.37	-1.12	0.99	-1.46	-1.76	1.09	-0.97	-1.00	-0.97	1.12	1.11	1.59	1.14	1.91	1.33	x	-1.24	-1.96	-1.80	1.39	1.41	-1.45	-1.06	1.53	1.05	-1.93	-1.11	-1.12	-1.05	-1.13	-1.05	Х	1.04	-1.29
0.04	0.34	0.28	0.21	0.36	0.15	0.38	×	0.38	0.66	0.85	0.15	0.76	x	x	0.65	0.65	Х	0.94	0.06	0.54	0.05	x	x	0.90	0.77	0.45	0.07	0.94	0.94	0.70	0.91	x	x	x	0.71
1.65	1.29	1.35	0.21	-0.07	-1.43	-1.68	-0.03	0.03	0.04	0.17	1.15	1.16	0.01	-0.11	1.55	0.16	N/A	-1.22	-1.77	-1.59	0.05	1.40	-1.39	-0.01	1.36	1.25	-1.61	-1.13	-1.12	-1.08	-1.23	-1.30	N/A	-0.01	-1.22
0.25	0.25	0.03	1.33	1.07	0.04	0.09	1.12	1.01	1.04	1.13	0.03	0.04	1.59	1.25	0.36	1.18	N/A	0.03	0.19	0.22	1.34	0.02	0.06	1.05	0.18	0.20	0.32	0.02	0.00	0.03	0.10	0.25	N/A	1.05	0.08

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ilvB ilvH prfB mrca hemA-1	DNA-binding protein HU		SPO2612
ilvB ilvH mrca hemA-1	hypothetical protein		SPO2610
ilvB ilvH mrca hemA-1	alcohol dehydrogenase		SPO2609
ilvB ilvH mrca hemA-1	aldehyde dehydrogenase		SPO2608
ilvB ilvH mrca hemA-1	gamma-glutamylisopropylamide synthetase		SPO2607
ilvB ilvH prfB mrca hemA-1	solute-binding family 7 protein		SPO2606
ilvB ilvH prfB mrca hemA-1	TRAP dicarboxylate transporter subunit DctM		SPO2605
ilvB ilvH prfB mrca hemA-1	hypothetical protein		SPO2604
ilvB ilvH prfB mrca hemA-1	N-formylglutamate amidohydrolase		SPO2603
ilvB ilvH prfB mrca hemA-I	RpiR family transcriptional regulator		SPO2602
ilvB ilvH prfB mrca ispG hemA-1	pirin		SPO2601
ilvB ilvH prfB mrca ispG hemA-1	hypothetical protein		SPO2598
ilvB ilvH prfB mrca ispG hemA-1	hypothetical protein		SPO2597
ilvB ilvH mrca	5-aminolevulinate synthase (EC:2.3.1.37)	hemA-1	SPO2596
ilvB ilvH mrca	hypothetical protein		SPO2595
ilvB ilvH mrca	4-hydroxy-3-methylbut-2-en-1-yl diphosphate synthase (EC:1.17.7.1)	ispG	SPO2594
ilvB ilvH mrca	hypothetical protein		SPO2593
ilvB ilvH prfB mrca	beta-lactamase		SPO2592
ilvB ilvH prfB mrca	hypothetical protein		SPO2591
ilvB ilvH prfB mrca	M48 family peptidase		SPO2590
ilvB ilvH prfB mrca	class I and II aminotransferase		SPO2589
ilvB ilvH prfB	N-acetylmuramoyl-L-alanine amidase		SPO2588
ilvB ilvH	penicillin-binding protein 1A	mrca	SPO2587
hypothetical protein solute-binding family 7 sensor histidine kinase LuxR family transcripti TetR family transcripti hypothetical protein acctolactate synthase 3 <i>ilvH</i> (EC:2.2.1.6) hypothetical protein DNA-binding protein bypothetical protein amidase (EC:3.5.1.4) glyoxalase	type I secretion target repeat-containing protein		SPO2586
hypothetical protein solute-binding family 7 sensor histidine kinase LuxR family transcripti hypothetical protein acetolactate synthase 3 acetolactate synthase 3 (EC:2.2.1.6) hypothetical protein DNA-binding protein hypothetical protein amidase (EC:3.5.1.4) glyoxalase	peptide chain release factor 2	prfB	SPO2585
hypothetical protein solute-binding family 7 sensor histidine kinase LuxR family transcripti hypothetical protein acetolactate synthase 3 acetolactate synthase 3 acetolactate synthase 3 (EC:2.2.1.6) hypothetical protein DNA-binding protein hypothetical protein	glyoxalase		SPO2584
hypothetical protein solute-binding family 7 sensor histidine kinase LuxR family transcripti TetR family transcripti hypothetical protein acetolactate synthase 3 <i>ilvH</i> (EC:2.2.1.6) hypothetical protein DNA-binding protein	amidase (EC:3.5.1.4)		SPO2583
hypothetical protein solute-binding family 7 sensor histidine kinase LuxR family transcripti TetR family transcripti hypothetical protein acetolactate synthase 3 acetolactate synthase 3 ih/H (EC:2.2.1.6) hypothetical protein DNA-binding protein	hypothetical protein		SPO2582
hypothetical protein solute-binding family 7 sensor histidine kinase LuxR family transcripti TetR family transcripti hypothetical protein acetolactate synthase 3 <i>ilvH</i> (EC:2.2.1.6) hypothetical protein	DNA-binding protein		SPO2581
hypothetical protein solute-binding family 7 sensor histidine kinase LuxR family transcripti TetR family transcripti hypothetical protein acetolactate synthase 3 acetolactate synthase 3 ilvH (EC:2.2.1.6)	hypothetical protein		SPO2580
hypothetical protein solute-binding family 7 sensor histidine kinase LuxR family transcripti TetR family transcripti hypothetical protein acetolactate synthase 3	ŝ	ίlvH	SPO2579
	ω	ilvB	SPO2578
	hypothetical protein		SPO2577
	TetR family transcriptional regulator		SPO2576
	LuxR family transcriptional regulator		SPO2575
	sensor histidine kinase (EC:2.7.3)		SPO2574
	solute-binding family 7 protein		SPO2573
	hypothetical protein		SPO2572

-1.42	-2.39	1.66	1.95	1.34	-1.29	-1.56	-1.31	-1.51	-1.70	-1.74	х	1.88	3.05	1.29	1.21	-1.45	-1.13	2.00	1.15	1.57	-1.72	-1.57	1.52	-1.16	1.16	0.99	1.54	-1.70	-2.20	1.28	-1.17	-1.08	1.09	1.07	1.22	1.10	2.05
0.51	0.00	0.02	0.10	0.27	0.15	0.77	0.33	0.07	0.12	0.36	×	0.03	0.00	х	0.50	0.01	х	0.03	0.97	0.09	0.49	0.17	0.02	0.43	0.82	0.02	0.41	0.01	0.16	Х	0.25	0.54	0.85	1.00	0.39	0.07	0.56
-1.78	-1.71	-1.22	-1.18	-1.11	-1.78	-1.50	1.05	-1.04	-1.25	-1.29	1.28	1.19	1.75	1.52	-1.07	1.93	1.13	-1.35	-1.07	1.29	-1.05	-1.07	1.70	0.99	-1.05	1.24	1.25	-1.16	1.73	1.54	1.23	1.50	3.28	-1.11	-1.26	-3.16	-1.54
0.06	0.03	0.21	0.58	0.17	0.02	0.19	0.72	0.87	0.22	0.02	0.56	0.11	0.07	0.74	0.55	0.02	0.84	0.07	0.28	0.36	0.86	0.65	0.01	0.82	0.90	0.48	0.32	0.39	0.04	0.24	0.20	0.11	0.01	0.86	0.07	0.01	0.17
-1.60	-2.05	0.22	0.39	0.12	-1.54	-1.53	-0.13	-1.28	-1.48	-1.52	N/A	1.54	2.40	1.41	0.07	0.24	0.00	0.33	0.04	1.43	-1.39	-1.32	1.61	-0.09	0.05	1.11	1.40	-1.43	-0.24	1.41	0.03	0.21	2.19	-0.02	-0.02	-1.03	0.26
0.18	0.34	1.44	1.57	1.23	0.25	0.03	1.18	0.24	0.22	0.22	N/A	0.35	0.65	0.12	1.14	1.69	1.13	1.68	1.11	0.14	0.34	0.25	0.09	1.07	1.11	0.13	0.14	0.27	1.97	0.13	1.20	1.29	1.10	1.09	1.24	2.13	1.80
-2.29	-1.94	-1.66	-1.61	-2.26	-4.55	-2.34	-1.99	-1.87	-3.00	-2.75	2.67	2.81	4.82	-2.04	1.60	-3.71	2.98	1.69	1.02	3.18	-1.11	2.30	1.19	4.00	-1.07	0.94	1.69	-1.34	-3.26	3.83	2.76	1.84	-1.61	1.92	1.98	1.28	2.34
0.02	0.00	0.01	0.02	0.00	0.01	0.11	0.06	0.01	0.03	0.03	0.01	0.03	0.01	Х	0.06	0.01	0.21	0.06	0.13	0.03	0.78	0.08	0.17	0.01	0.97	0.24	0.10	0.16	0.00	0.01	0.05	0.03	0.43	0.69	0.01	0.26	0.14
1.10	-1.19	-1.21	-1.30	-1.22	-1.93	-1.88	-1.73	-1.81	-1.71	-1.22	-1.04	1.18	-1.04	-1.36	-1.48	2.01	1.03	1.58	1.58	-1.22	-1.29	-1.07	-1.54	-1.16	1.22	1.26	1.24	-1.05	2.91	-1.21	1.61	-2.07	-1.10	-1.29	1.25	1.76	1.88
0.71	0.03	0.01	0.02	0.01	0.04	0.16	0.49	0.04	0.04	0.22	0.88	0.34	0.42	х	0.06	0.09	0.95	0.12	0.21	0.45	0.03	0.59	0.13	0.10	0.62	0.11	0.52	0.74	0.01	0.06	0.01	0.01	0.63	0.76	0.13	0.08	0.05
-0.60	-1.57	-1.44	-1.46	-1.74	-3.24	-2.11	-1.86	-1.84	-2.36	-1.99	0.82	2.00	1.89	-1.70	0.06	-0.85	2.01	1.64	1.30	0.98	-1.20	0.62	-0.18	1.42	0.08	1.10	1.47	-1.20	-0.18	1.31	2.19	-0.12	-1.36	0.32	1.62	1.52	2.11
1.70	0.38	0.22	0.16	0.52	1.31	0.23	0.13	0.03	0.64	0.77	1.86	0.82	2.93	0.34	1.54	2.86	0.98	0.05	0.28	2.20	0.09	1.69	1.37	2.58	1.15	0.16	0.23	0.15	3.09	2.52	0.57	1.96	0.26	1.61	0.37	0.24	0.23
1.02	-1.03	-1.45	-1.52	-1.17	-1.03	-1.56	1.05	-1.16	-1.23	1.17	-1.26	1.36	-1.04	1.17	1.22	-1.28	1.43	1.14	1.34	1.14	1.14	2.48	-1.38	-1.23	1.21	1.34	1.29	-1.39	1.86	-1.05	-1.32	1.59	1.75	-1.06	-1.35	1.24	1.03
0.80	0.10	0.10	0.00	0.07	0.56	0.14	0.75	0.16	0.13	0.20	0.05	0.05	0.69	0.71	0.15	0.52	0.56	0.48	0.28	0.67	0.18	0.00	0.02	0.06	0.57	0.17	0.41	0.12	0.04	0.17	0.03	0.12	0.05	0.88	0.19	0.31	0.90
1.20	-1.33	-1.31	-1.45	-1.41	0.96	-1.61	-1.14	-2.28	-1.58	-1.05	-2.06	1.11	0.93	2.01	1.42	х	1.12	1.20	1.58	-1.30	-1.16	1.69	Х	-1.72	-0.97	1.18	1.29	-1.78	3.91	-1.58	-1.28	5.56	3.34	-1.17	-1.51	1.41	1.45
0.57	0.02	0.19	0.03	0.16	0.33	0.09	0.93	0.02	0.08	0.75	0.19	0.06	0.46	Х	0.11	Х	0.37	0.34	0.29	0.41	0.73	0.03	х	0.04	0.78	0.26	0.50	0.06	0.00	0.04	0.16	0.00	0.01	0.77	0.13	0.30	0.08
1.11	-1.18	-1.38	-1.49	-1.29	-0.03	-1.59	-0.04	-1.72	-1.41	0.06	-1.66	1.24	-0.06	1.59	1.32	N/A	1.28	1.17	1.46	-0.08	-0.01	2.09	N/A	-1.48	0.12	1.26	1.29	-1.59	2.89	-1.32	-1.30	3.58	2.55	-1.12	-1.43	1.33	1.24
0.09	0.15	0.07	0.04	0.12	1.00	0.03	1.10	0.56	0.18	1.11	0.40	0.13	0.98	0.42	0.10	N/A	0.16	0.03	0.12	1.22	1.15	0.39	N/A	0.24	1.09	0.08	0.00	0.20	1.03	0.27	0.02	1.99	0.80	0.05	0.08	0.09	0.21

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SPO2649	SPO2648	SPO2647	SPO2646	SPO2645	SPO2644	SPO2643	SPO2642	SPO2641	SPO2640	SPO2639	SPO2638	SPO2637	SPO2636	SPO2635	SPO2634	SPO2633	SPO2632	SPO2631	SPO2630	SPO2629	SPO2628	SPO2627	SPO2626	SPO2625	SPO2624	SPO2623	SPO2622	SPO2621	SPO2620	SPO2619	SPO2618	SPO2617	SPO2616	SPO2615	SPO2614	SPO2613
											infC			$cy_{S}H$			cobA-I											tpiA					tgt			lon
hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	S2 family protease	von Willebrand factor A	hypothetical protein	hypothetical protein	hypothetical protein	XdhC/CoxI family protein	molybdopterin biosynthesis protein	translation initiation factor IF-3	ferredoxin-NADP reductase (EC:1.18.1.2)	hypothetical protein	pnospnoadenosme pnospnosuitate reductase (EC:1.8.4.8)	sulfite reductase	hypothetical protein	uroporpnyrin-ш с-meinyitransierase (ЕС:2.1.1.107)	AsnC family transcriptional regulator	С4-сисатоохугае папярот, sensor protein (ЕС:2.7.3)	C4-dicarboxylate transport transcriptional regulator	protein	TRAP dicarboxylate transporter subunit DctQ	TRAF C+-ticarooxytate transport system permease DefM	hypothetical protein	AsnC family transcriptional regulator	amino acid transporter LysE	acetyltransferase	triosephosphate isomerase (EC:5.3.1.1)	iron-sulfur cluster assembly accessory protein	hypothetical protein	hypothetical protein	hypothetical protein	queuine tRNA-ribosyltransferase (EC:2.4.2.29)	Oye family is ADD-uependent havin oxidoreductase	hypothetical protein	ATP-dependent protease La (EC:3.4.21.53)
-1.67	-1.71	-1.55	1.09	-2.06	-1.05	1.13	1.16	x	-1.58	0.95	1.23	-1.31	1.14	1.40	-1.03	1.13	-1.45	1.63	-0.98	-0.94	-1.16	-1.13	-1.26	-1.03	-1.38	-2.41	-1.21	-0.99	0.91	-1.13	1.39	1.49	х	-1.30	-2.15	1.18
0.05	0.05	0.30	0.58	0.00	0.53	0.40	0.93	×	0.05	0.93	0.90	0.31	0.41	0.45	0.28	0.66	0.09	0.19	0.97	0.55	0.23	0.41	0.10	0.93	0.22	0.21	0.58	0.98	0.14	0.15	0.02	0.74	x	0.80	0.01	0.97
-1.60	-1.62	-1.61	2.44	-1.09	2.20	1.16	-1.44	-1.15	1.12	1.23	1.24	-2.45	-2.52	-2.99	-2.09	-3.37	-2.30	-1.72	-1.14	-1.08	-3.48	-2.13	1.10	1.16	1.12	-1.13	-1.15	1.22	-1.60	-1.34	-1.38	-1.08	1.71	-1.96	-1.84	-1.60
0.00	0.08	0.37	0.01	0.63	0.01	0.63	0.55	0.91	0.50	0.49	0.38	0.03	0.01	0.02	0.02	0.03	0.01	0.03	0.37	0.43	0.02	0.00	0.55	0.74	0.05	0.77	0.28	0.51	0.03	0.02	0.02	0.75	0.01	0.02	0.01	0.19
-1.64	-1.67	-1.58	1.77	-1.58	0.58	1.15	-0.14	N/A	-0.23	1.09	1.24	-1.88	-0.69	-0.80	-1.56	-1.12	-1.88	-0.05	-1.06	-1.01	-2.32	-1.63	-0.08	0.06	-0.13	-1.77	-1.18	0.11	-0.34	-1.24	0.01	0.21	N/A	-1.63	-2.00	-0.21
0.03	0.04	0.03	0.68	0.48	1.63	0.02	1.30	N/A	1.35	0.14	0.01	0.57	1.83	2.20	0.53	2.25	0.43	1.68	0.08	0.07	1.16	0.50	1.18	1.10	1.25	0.64	0.03	1.11	1.26	0.11	1.39	1.29	N/A	0.33	0.16	1.39
1.02	1.02	1.11	-2.19	-5.52	-2.18	-1.16	1.48	-1.11	-3.92	-1.80	2.98	7.66	34.30	23.00	21.10	8.57	6.00	7.56	2.67	3.72	17.40	29.00	9.23	-1.28	-0.94	-2.67	1.13	2.23	1.28	1.18	-1.30	0.88	4.52	6.46	20.60	2.64
0.68	0.77	0.87	0.00	0.00	0.02	0.29	0.17	×	0.00	0.73	0.13	0.00	0.03	0.00	0.00	0.00	0.00	0.02	0.11	0.00	0.00	0.00	0.00	0.84	0.33	0.24	0.15	0.48	0.47	0.41	0.06	0.75	0.00	0.17	0.00	0.28
-1.24	-1.19	-1.12	1.28	1.24	1.30	-1.02	0.99	1.07	1.87	1.53	1.29	2.52	3.54	2.89	2.63	3.09	3.77	1.73	1.57	1.53	2.94	1.33	1.44	1.11	-1.26	-1.43	1.43	-1.27	1.30	1.31	1.30	1.42	-1.70	5.36	7.08	-1.21
4 0.20		2 0.56	3 0.06	0.09	0.12	2 X	0.91	x	7 0.01	3 0.36		0.01	0.03	0.01	3 0.01	0.00	0.01		0.06	3 0.02	0.00	3 0.34	I 0.03	0.86	6 0.19	3 0.56		7 0.44	0.25	0.14	0.13	0.36	0 0.01	0.01	3 0.00	0.42
0 -0.1	0 -0.09			9 -2.14		-1.09	1 1.24	-0.02	1 -1.03	6 -0.14		1 5.09	3 18.92				1 4.89	6 4.65		2 2.63	0 10.17		3 5.34	6 -0.09				•		4 1.25	3 0.00	6 1.15	1 1.41	1 5.91	0 13.84	
1.13	)9 1.11	1.12	1.74	14 3.38	1.74	0.07	4 0.24		2.90	14 1.67	4 0.85	9 2.57	15.38		37 9.24	3 2.74		5 2.92		3 1.10	17 7.23	13.84	4 3.90	1.20	0 0.16	0.62	8 0.15	8 1.75	9 0.01	5 0.07		5 0.27	-1 3.11	0.55		2 1.93
								1.09						10.06 -1			1.12 1.														1.30 -1					
-1.07 0.	-1.04 0.	1.07 0.			1.47 0.	1.22 0.	-1.35 0.	X	1.34 0.	1.82 0.	1.45 0.	1.05 0.	0.98 0.	1.39 0.			1.36 0.		1.35 0.	1.31 0.	1.45 0.	1.09 0.	2.44 <mark>0.</mark>	1.41 0.	1.18 0.	1.18 0.	1.14 0.	1.37 0.	1.52 0.	1.00 0.	1.07 0.	1.47 0.	-1.35 0.	1.71 0.		2.10 0.
0.65 -1		0.91 1.	0.05 1.	0.96 -0	0.07 -1	0.25	0.43 2	×	0.18 1.	0.09 1.	0.13 1.	0.98 1.	0.73 1.	0.09 0.	0.35 1.	0.07 -1	0.23 2.	0.48 -1	0.20 1.	0.20 -1	0.19 1.	0.64 1.	0.01 2.	0.57 1.	0.44 -1	0.79 -1		0.32 1.	0.11 2	0.83 1.	0.67 1.	0.34 1.	0.19 -2	0.14 1.		
1.19 0	-1.05 0	1.08 0	1.29 0		-1.08 0	Х	2.24 0	X	1.68 0			1.11 0		0.97 0			2.14 0	-1.41 0	1.09 0	-1.25 0	1.91 0		2.47 0	1.42 0			Ŭ			1.17 0	1.18 0	1.46 0	2.54 0	1.98 0	1.16 0	
0.07 -	0.70 -	0.93 1	0.07 1	0.33 -(	0.78 (	X	0.05 (	X	0.20	0.25	0.88 1	0.79 1	0.59 1	0.18 -(	0.57 1	0.11	0.07 1	0.21 -	0.62 1	0.37 (	0.01 1	0.18 1	0.04 2	0.44 1	0.18 -(	0.94 (		0.78 1	0.08 1	0.62 1	0.32 (	0.31 1	0.01 -	0.09 1		
1.13	-1.05	1.08	1.33	-0.99	0.20	N/A	0.45	N/A	1.51	1.64	1.30	1.08	1.03	-0.21	1.23	-1.40	1.75	-1.27	1.22	0.03	1.68	1.31	2.46	1.42	0.12	0.06	0.08	1.24	1.79	1.09	0.05	1.47	1.95	1.85	1.09	2.18
0.06	0.01	0.01	0.04	0.03	1.28	N/A	1.80	N/A	0.17	0.19	0.16	0.03	0.04	1.18	0.04	0.24	0.39	0.14	0.13	1.28	0.23	0.21	0.02	0.01	1.30	1.12	1.06	0.13	0.27	0.09	1.13	0.01	0.60	0.14	0.07	0.08

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SPO2686	SPO2685	SPO2684	SPO2683	SPO2682	SPO2681	SPO2680	SPO2679	SPO2678	SPO2677	SPO2676	SPO2675	SPO2674	SPO2673	SPO2672	SPO2671	SPO2670	SPO2669	SPO2668	SPO2667	SPO2666	SPO2665	SPO2664	SPO2663	SPO2662	SPO2661	SPO2660	SPO2659	SPO2658	SPO2657	SPO2656	SPO2655	SPO2654	SPO2653	SPO2652	SPO2651	SPO2650
				tatB	tatC				purF			radA				alr							aspA		gltL	gltK	gltJ	gltI			dnaB	pyrE				
LysM/M23/M37 peptidase	hypothetical protein	hypothetical protein	twin arginine-targeting protein translocase	twin-arginine translocation protein TatB	twin-arginine translocation protein TatC	hypothetical protein	short chain denydrogenase/reductase oxidoreductase	hypothetical protein	amidophosphoribosyltransferase (EC:2.4.2.14)	hypothetical protein	colicin V production protein CvpA	DNA repair protein RadA	paraquat-inducible protein A	ABC transporter ATP-binding protein	hypothetical protein	alanine racemase (EC:5.1.1.1)	20G-Fe(II) oxygenase	LysR family transcriptional regulator	potar annuo actu ADC transporter suostrate-onionig	polar amino acid ABC transporter permease	polar amino acid ABC transporter permease	pota annuo acia Are nanspoter Are-oniung	aspartate ammonia-lyase (EC:4.3.1.1)	aspartate racemase	protein	glutamate/aspartate ABC transporter permease	glutamate/aspartate ABC transporter permease	giutantate/aspartate ABC transporter substrate- binding protein	D-cysteine desulfhydrase (EC:4.4.1.15)	LysR family transcriptional regulator	replicative DNA helicase (EC:3.6.1)	orotate phosphoribosyltransferase (EC:2.4.2.10)	dihydroorotase (EC:3.5.2.3)	hypothetical protein	glycosyl transferase family protein	FkbM family methyltransferase
-1.63	1.06	1.04	1.28	0.99	1.51	1.84	1.53	1.46	1.80	1.65	-1.21	1.21	1.43	1.84	1.93	2.00	-1.52	1.83	×	-1.44	х	-1.00	-1.52	х	×	Х	-1.88	-1.60	-1.56	-1.44	-1.50	1.55	1.80	1.53	2.02	-1.31
0.01	1.00	0.65	0.39	0.13	0.07	0.04	0.01	0.23	0.01	0.52	0.06	0.17	0.04	0.44	0.12	0.03	0.50	0.37	х	Х	×	0.98	0.67	x	х	х	0.02	0.33	Х	0.72	0.57	0.19	0.02	0.60	0.21	0.49
-1.10	1.09	-1.07	-1.38	-1.47	-1.08	0.99	1.86	1.41	1.65	2.02	0.99	1.19	-1.11	-1.03	1.34	1.01	-1.09	-0.99	-1.24	-1.20	х	-1.00	1.17	-1.07	-1.20	-1.37	-2.00	-2.80	-1.45	-1.24	0.99	1.86	1.12	-1.02	1.15	-1.22
0.41	0.90	0.74	0.27	0.07	0.46	0.34	0.24	0.33	0.03	0.02	0.78	0.47	0.72	0.89	0.18	0.93	0.47	0.79	0.20	0.88	×	1.00	0.75	0.87	0.60	0.41	0.00	0.04	0.84	0.36	0.84	0.17	0.54	0.87	0.44	0.20
-1.37	1.08	-0.02	-0.05	-0.24	0.22	1.41	1.70	1.44	1.73	1.84	-0.11	1.20	0.16	0.41	1.64	1.51	-1.31	0.42	N/A	-1.32	N/A	-1.00	-0.18	N/A	N/A	N/A	-1.94	-2.20	-1.51	-1.34	-0.25	1.71	1.46	0.26	1.59	-1.27
0.27	0.02	1.06	1.33	1.23	1.30	0.43	0.17	0.03	0.08	0.18	1.10	0.01	1.27	1.44	0.30	0.50	0.21	1.41	N/A	0.12	N/A	0.00	1.35	N/A	N/A	N/A	0.06	0.60	0.06	0.10	1.25	0.16	0.34	1.28	0.44	0.05
1.43	2.44	1.37	1.76	1.59	2.31	3.48	4.42	1.55	3.25	-1.16	1.58	1.24	1.36	2.74	3.04	2.50	1.21	-1.90	-1.59	-2.27	Х	1.15	-1.48	-1.77	-0.94	1.42	-2.15	-1.64	-1.80	-1.80	1.13	2.73	2.44	-1.93	1.26	1.14
0.03	0.62	0.47	0.13	0.09	0.02	0.00	0.00	0.34	0.01	0.86	0.08	0.03	0.07	0.22	0.11	0.00	0.66	0.09	0.39	0.74	×	0.72	0.75	0.12	0.21	0.11	0.04	0.38	0.54	0.57	0.70	0.03	0.02	0.30	0.73	0.85
1.01	1.21	1.32	1.08	1.03	-1.17	-1.04	-1.44	-1.12	1.14	1.90	-1.11	-1.05	1.75	0.97	-1.33	-1.35	1.62	1.27	-1.25	-1.31	Х	1.21	-1.03	-1.25	-1.30	-1.93	-1.72	-2.06	-1.29	-1.10	-1.12	-1.29	-1.11	1.04	1.04	-1.44
0.76	0.80	0.05	0.97	0.75	0.15	0.30	0.17	0.14	0.53	0.07	0.07	0.25	0.05	0.89	0.30	0.25	0.08	0.24	×	0.72	х	0.72	0.99	0.53	0.60	0.13	0.11	0.02	0.87	0.63	0.21	0.04	0.28	1.00	0.94	0.03
1.22	1.83	1.35	1.42	1.31	0.57	1.22	1.49	0.22	2.20	0.37	0.24	0.10	1.56	1.85	0.86	0.58	1.42	-0.32	-1.42	-1.79	N/A	1.18	-1.26	-1.51	-1.12	-0.26	-1.94	-1.85	-1.55	-1.45	0.00	0.72	0.67	-0.45	1.15	-0.15
0.21	0.62	0.03	0.34	0.28	1.74	2.26	2.93	1.34	1.06	1.53	1.35	1.15	0.19	0.89	2.19	1.93	0.21	1.59	0.17	0.48	N/A	0.03	0.23	0.26	0.18	1.68	0.21	0.21	0.26	0.35	1.13	2.01	1.78	1.49	0.11	1.29
1.15	-1.02	1.15	1.01	-1.40	-1.13	-1.43	1.07	2.10	-1.08	-1.32	-1.39	1.02	-1.63	-1.03	1.04	-1.03	-1.44	1.18	-1.25	-1.08	-1.40	-1.52	-1.00	-1.18	-1.12	-1.37	-1.54	-1.29	-1.41	-1.09	1.38	-1.52	-1.79	-2.30	-1.11	1.30
0.08	0.98	0.12	0.33	0.07	0.05	0.62	0.20	0.01	0.67	0.30	0.02	0.93	0.08	0.88	0.89	0.74	0.09	0.37	0.10	0.96	х	0.27	0.95	0.48	0.43	0.13	0.03	0.16	0.71	0.55	0.30	0.01	0.04	0.04	0.75	0.06
-1.26	-1.11	-1.09	1.16	-1.17	-1.44	-1.80	-1.39	1.41	-1.55	-1.51	-1.42	-1.08	-1.53	-1.30	-1.41	-1.31	1.05	-1.20	1.05	2.19	Х	-1.18	1.23	1.08	-0.98	-1.46	-1.24	-0.98	-1.21	1.06	1.38	-1.94	-1.78	-1.65	-1.09	-1.20
0.01	0.92	0.45	0.97	0.01	0.01	0.07	0.48	0.09	0.14	0.39	0.03	0.20	0.11	0.59	0.09	0.03	0.94	x	x	0.07	х	0.82	0.34	Х	0.61	0.22	0.46	0.55	x	0.82	0.29	0.02	0.05	0.10	0.74	0.45
-0.06	-1.07	0.03	1.09	-1.29	-1.29	-1.62	-0.16	1.76	-1.32	-1.42	-1.41	-0.03	-1.58	-1.17	-0.19	-1.17	-0.20	-0.01	-0.10	0.56	N/A	-1.35	0.12	-0.05	-1.05	-1.42	-1.39	-1.14	-1.31	-0.02	1.38	-1.73	-1.79	-1.98	-1.10	0.05
1.21	0.05	1.12	0.08	0.12	0.16	0.19	1.23	0.35	0.24	0.10	0.02	1.05	0.05	0.14	1.23	0.14	1.25	1.19	1.15	1.64	N/A	0.17	1.12	1.13	0.07	0.04	0.15	0.16	0.10	1.08	0.00	0.21	0.01	0.33	0.01	1.25

# Chapter 10: Appendix

SPO2724	SPO2723	SPO2722	SPO2721	SPO2720	SPO2719	SPO2718	SPO2717	SPO2716	SPO2715	SPO2714	SPO2713	SPO2712	SPO2711	SPO2710	SPO2709	SPO2708	SPO2707	SPO2706	SPO2705	SPO2704	SPO2703	SPO2702	SPO2701	SPO2700	SPO2699	SPO2698	SPO2697	SPO2696	SPO2695	SPO2694	SPO2693	SPO2692	SPO2691	SPO2690	SPO2689	SPO2688	SPO2687
											secA							caiD-1																		surE	pcm-2
hypothetical protein	protein SapC protein	type I secretion target repeat-containing protein	GAF domain-containing protein	M50 family peptidase	preprotein translocase subunit SecA	hypothetical protein	CaiB/BaiF family protein	3-oxoadipate enol-lactonase	endoribonuclease L-PSP	aldehyde dehydrogenase	luciferase	carnitinyl-CoA dehydratase	3-hydroxyacy1-CoA dehydrogenase	AraC family transcriptional regulator	hypothetical protein	opine/polyamine ABC transporter ATF-binding protein	protein	opine/polyamine ABC transporter permease	opine/polyamine ABC transporter permease	acyl-CoA dehydrogenase	acyl-CoA synthetase	beta-lactamase	peptide chain release factor 3	cytochrome c'	hypothetical protein	short chain denydrogenase/reductase oxidoreductase	hypothetical protein	hypothetical protein	iron ABC transporter ATP-binding protein	stationary phase survival protein SurE (EC:3.1.3.2)	protein-L-isoaspartate O-methyltransferase (EC:2.1.1.77)						
-2.17	-2.58	-3.02	-4.13	-1.91	-1.81	-1.06	-1.24	-1.54	-1.88	-1.40	-1.40	2.66	3.72	1.79	1.76	2.80	2.80	1.74	2.93	-1.60	-1.64	-1.40	1.41	1.17	х	×	-1.19	х	-1.26	-1.13	-1.03	1.38	1.30	1.40	×	1.30	-2.13
0.03	0.02	0.01	0.00	0.82	0.29	1.00	0.52	0.28	0.29	0.12	×	0.04	0.00	0.06	0.88	0.27	0.07	0.04	0.00	0.04	0.14	0.46	0.30	0.34	×	×	0.91	×	0.32	0.03	0.95	0.65	0.60	0.06	×	0.24	0.01
-1.69	-1.76	-2.04	-1.84	-1.41	1.03	1.47	1.22	1.48	1.06	-1.05	1.03	÷	1.12	1.40	1.54	1.14	1.36	1.17	1.17	-1.14	х	-2.05	-1.66	-1.03	-1.37	-1.19	-1.24	х	-1.06	-1.21	1.02	-1.16	1.02	1.27	1.32	-1.18	-1.45
0.02	0.04	0.00	0.06	0.83	0.75	0.41	0.25	0.04	0.87	0.92	0.52	0.73	0.56	0.04	0.71	0.48	0.19	0.22	0.59	0.44	×	0.04	0.14	0.98	0.61	0.53	0.78	×	0.10	0.03	0.95	0.47	0.95	0.17	0.18	0.37	0.05
-1.93	-2.17	-2.53	-2.99	-1.66	-0.39	0.21	-0.01	-0.03	-0.41	-1.23	-0.19	0.78	2.42	1.60	1.65	1.97	2.08	1.46	2.05	-1.37	N/A	-1.73	-0.13	0.07	N/A	N/A	-1.22	N/A	-1.16	-1.17	-0.01	0.11	1.16	1.34	N/A	0.06	-1.79
0.24	0.41	0.49	1.15	0.25	1.42	1.27	1.23	1.51	1.47	0.17	1.22	1.89	1.30	0.19	0.11	0.83	0.72	0.28	0.88	0.23	N/A	0.33	1.54	1.10	N/A	N/A	0.03	N/A	0.10	0.04	1.03	1.27	0.14	0.06	N/A	1.24	0.34
-1.95	-1.92	-1.84	-2.85	-3.52	1.09	-1.49	-2.79	-1.95	-3.09	-2.10	-2.06	-1.93	1.12	-1.72	1.12	-2.64	-1.34	4.83	1.24	-1.14	5.38	5.39	-1.26	1.41	-1.37	-1.01	-1.18	Х	3.91	-1.61	1.04	1.83	1.74	2.19	4.45	1.76	-1.75
0.04	0.04	0.01	0.00	0.72	0.18	0.65	0.13	0.15	0.17	0.02	0.12	0.10	0.15	0.19	0.98	0.04	0.83	0.00	0.06	0.51	0.08	0.03	0.98	0.44	0.85	0.22	0.98	x	0.00	0.02	0.84	0.28	0.30	0.01	0.00	0.00	0.01
1.50	2.28	2.98	3.02	1.28	-1.18	-0.98	-1.28	-1.09	1.47	1.50	1.67	1.18	1.22	1.18	1.12	1.12	1.41	1.16	1.17	-1.01	х	-1.30	-1.01	-1.41	-1.22	-0.98	-1.15	х	-1.05	1.85	-1.12	1.26	-1.09	-1.16	-1.44	-1.20	1.88
0.00	0.05	0.01	0.00	0.46	0.63	0.94	0.20	0.79	0.44	0.11	0.04	0.44	0.16	0.27	0.91	0.64	0.02	0.23	0.29	0.97	×	0.40	0.97	0.72	0.67	0.79	0.51	x	0.31	0.06	0.50	0.32	0.58	0.02	0.21		0.01
-0.23	0.18	0.57	0.09	-1.12	-0.04	-1.23	-2.04	-1.52	-0.81	-0.30	-0.20	-0.38	1.17	-0.27	1.12	-0.76	0.03	3.00	1.21	-1.08	N/A	2.05	-1.14	0.00	-1.30	-0.99	-1.17	N/A	1.43	0.12	-0.04	1.55	0.33	0.52	1.51	0.28	0.06
1.73		2.41	2.94	2.40	1.14	0.26	0.76	0.43	2.28	1.80	1.87	1.56	0.05	1.45	0.00	1.88	1.38	1.84	0.04	0.06	N/A	3.35	0.13	1.41	0.08	0.02	0.02	N/A	2.48	1.73	. 1.08	0.29	1.42	1.68	2.95	1.48	1.82
1.05		1.93	1.43	-1.08	¥ 1.12	5 1.30	1.23	3 1.79	3 1.29	1.04	1.20	-1.24	-1.00	1.34	1.40	-1.39	3 1.04	-1.12	-1.01	-1.25		-1.17	1.43	1.79	3 1.74	-1.11	-1.11	X	-2.38	-1.16		-1.12	1.57	-1.12	-1.14		1.04
0.65		3 0.07	3 0.06	8 0.92	2 0.18	0.61	3 0.35	9 0.04	€ 0.61	4 0.82	0.04	4 0.49	0 0.91	4 0.10	0.73	9 0.10	4 0.62	2 0.22	1 0.99	5 0.16	7 X	7 0.17	3 0.15			1 0.53	1 0.73	x	8 0.00	6 0.46	5 0.79	2 0.43	7 0.10	2 0.11			4 0.93
5 1.45		7 6.57	6 4.03	2 -0.90		1 1.26	5 1.41	4 1.37	1 1.32	2 1.15	4 X	9 1.04				0 -1.17	2 -0.99	2 -0.96	-1.04	6 -1.32	X	7 1.11	5 1.23	9 1.59	3 3.26	3 X	3 1.34	Х				3 0.98	0 1.11	-1.75			
5 0.04	_	7 0.00	3 0.00	0 0.65	0.09	6 0.45	1 0.13	7 0.09	2 0.52	5 0.28	X	4 0.84	0 0.55	0.52	0 X	0.83	0.56	0.47	0.88	32 0.66	x		3 X	9 0.61	8 9		4 X	X	55 <b>0.01</b>	)8 0.23		8 0.72	1 0.36	<sup>75</sup> 0.02			8 0.75
		00 4.25	00 2.73		09 -0.06												56 0.02		-1.03	-1.29	۲ N/A		۲.33 x	61 1.69	<b>ξ</b> 2.50	۲ N/A	ζ 0.12	۲ N/A		_	•	72 -0.07					
1.25 0.				0.99 0.		1.28 0.	1.32 0.	1.58 0.	1.31 0.	1.10 0.	N/A N	-0.10 1.	-1.00 0.		2.10 0.	-1.28 0.		-1.04 0.				-0.03 1.															1.11 0.
0.20	0.38	2.32	1.30	0.09	1.19	0.02	0.09	0.21	0.02	0.05	N/A	1.14	0.00	1.13	0.70	0.11	1.02	0.08	0.02	0.04	N/A	1.14	0.10	0.10	0.76	N/A	1.23	N/A	0.14	0.04	1.08	1.05	0.23	0.32	0.40	0.03	0.07

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SPO2766	SPO2765	SPO2764	SPO2763	SPO2762	SPO2761	SPO2760	SPO2759	SPO2758	SPO2757	SPO2756	SPO2755	SPO2754	SPO2753	SPO2752	SPO2751	SPO2750	SPO2749	SPO2748	SPO2747	SPO2746	SPO2745	SPO2744	SPO2743	SPO2742	SPO2741	SPO2740	SPO2739	SPO2738	SPO2736	SPO2735	SPO2734	SPO2733	SPO2731	SPO2730	SPO2729	SPO2726	SPO2725
nuoK	nuoL	nuoM	nuoN																							tmk				hsdR	hsdS	hsdM					
NADH dehydrogenase subunit	NADH dehydrogenase subunit L (EC:1.6.99.5)	NADH dehydrogenase subunit M (EC:1.6.5.3)	NADH dehydrogenase subunit N (EC:1.6.5.3)	(EC:6.3.4.15)	pantothenate kinase (EC:2.7.1.33)	metallo-beta-lactamase	NUDIX family hydrolase	hypothetical protein	EF hand domain-containing protein	RNA polymerase sigma factor	hypothetical protein	hypothetical protein	diguanylate cyclase,/response regulator	hypothetical protein	HAD-superfamily hydrolase	trimethylamine methyltransferase	hypothetical protein	hypothetical protein	diguanylate cyclase	hypothetical protein	hypothetical protein	malonate transporter	hypothetical protein	TatD family hydrolase	DNA polymerase III subunit delta' (EC:2.7.7.7)	thymidylate kinase (EC:2.7.4.9)	D-alanyl-D-alanine carboxypeptidase (EC:3.4.16.4)	lipoprotein	hypothetical protein	ty је т технискол-шонпсаноп system, is snothit (EC:3.1.21.3)	type I restriction-modification system subunit S	type 1 restriction-modification system, M subunit (EC:2.1.1.72)	ISSpo7, transposase	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein
2.17	1.86	1.68	2.08	1.32	1.10	-1.70	2.86	8.02	-1.36	1.13	-1.23	1.34	1.60	1.60	1.93	4.96	-1.02	1.27	-1.01	1.06	-1.49	1.81	-1.05	1.14	-1.08	1.08	-1.16	1.20	1.35	1.63	1.84	1.73	х	-1.04	-1.18	-1.23	1.50
0.07	0.08	0.05	0.01	0.31	0.80	0.63	0.03	0.05	0.02	0.74	0.30	0.76	0.06	0.06	0.84	0.03	0.41	0.12	1.00	0.21	0.71	0.01	0.58	0.51	0.69	0.59	0.76	0.99	0.71	0.64	0.13	0.01	х	0.80	0.89	0.77	×
1.19	1.12	1.11	1.04	1.18	1.15	1.20	2.49	3.69	1.00	1.40	-1.15	1.48	1.03	1.13	1.58	2.83	-1.24	1.40	-1.05	-1.25	1.32	1.14	1.11	-1.09	-1.47	-1.19	0.98	-1.71	1.38	1.56	1.23	1.35	1.32	-1.15	-1.18	-1.25	-1.89
0.61	0.10	0.66	0.92	0.22	0.76	0.47	0.00	0.00	0.99	0.43	0.27	0.30	0.91	0.57	0.34	0.01	0.52	0.09	0.76	0.13	0.78	0.18	0.64	0.77	0.31	0.44	0.55	0.13	0.11	0.15	0.34	0.15	0.05	0.93	0.76	0.84	×
1.68	1.49	1.40	1.56	1.25	1.13	-0.25	2.68	5.86	-0.18	1.27	-1.19	1.41	1.32	1.37	1.76	3.90	-1.13	1.34	-1.03	-0.10	-0.09	1.48	0.03	0.02	-1.28	-0.05	-0.09	-0.26	1.37	1.60	1.54	1.54	N/A	-1.10	-1.18	-1.24	-0.20
0.49	0.37	0.29	0.52	0.07	0.02	1.45	0.19	2.17	1.18	0.14	0.04	0.07	0.29	0.24	0.18	1.07	0.11	0.06	0.02	1.16	1.41	0.33	1.08	1.12	0.20	1.14	1.07	1.46	0.01	0.03	0.31	0.19	N/A	0.05	0.00	0.01	1.70
3.36	2.85	3.08	4.02	2.81	2.09	1.21	2.79	7.74	-2.26	-0.97	1.97	1.38	1.70	2.69	2.25	-2.57	1.02	-1.37	-1.87	0.84	-1.17	3.23	1.95	2.10	1.05	1.28	1.16	1.09	1.81	1.06	1.80	1.69	1.46	1.55	-2.49	-4.59	1.74
0.02	0.01	0.02	0.02	0.03	0.11	0.66	0.02	0.00	0.00	0.78	0.01	0.72	0.04	0.01	0.76	0.11	0.50	0.26	0.35	0.03	0.92	0.00	0.03	0.04	0.93	0.06	0.61	0.07	0.40	0.96	0.34	0.01	х	0.16	0.68	0.08	0.02
-1.07	-1.35	-1.08	-1.56	1.07	-1.08	1.12	1.94	1.55	1.05	-1.02	1.06	1.05	1.59	-1.14	-1.07	-1.12	1.08	1.19	1.07	1.55	1.17	-1.02	-1.05	1.05	-1.03	1.02	1.71	1.18	-1.00	1.13	-1.05	1.02	-1.41	-1.45	-1.11	-1.00	×
0.47	0.03	0.46	0.04	0.93	0.79	0.21	0.03	0.03	0.69	0.95	0.89	0.91	0.13	0.04	0.64	0.63	0.47	0.47	0.76	0.05	0.88	0.78	0.74	0.96	0.85	0.47	0.05	0.31	0.99	0.44	0.80	0.82	0.09	Х	0.81	1.00	×
1.15	0.75	1.00	1.23	1.94	0.51	1.17	2.37	4.65	-0.61	-1.00	1.52	1.22	1.65	0.78	0.59	-1.85	1.05	-0.09	-0.40	1.19	0.00	1.11	0.45	1.58	0.01	1.15	1.44	1.14	0.41	1.10	0.38	1.36	0.03	0.05	-1.80	-2.80	N/A
2.22	2.10	2.08	2.79	0.87	1.59	0.04	0.42	3.10	1.66	0.03	0.46	0.17	0.05	1.92	1.66	0.73	0.03	1.28	1.47	0.36	1.17	2.13	1.50	0.52	1.04	0.13	0.27	0.04	1.41	0.03	1.43	0.34	1.44	1.50	0.69	1.80	N/A
1.18	-1.15	1.02	-1.09	-1.10	-1.47	1.04	1.09	1.84	-1.00	-1.10	-1.26	1.48	-1.20	-1.49	1.29	1.32	1.04	-1.01	-1.27	1.62	1.17	-1.27	-1.66	-1.54	-1.41	1.62	1.93	1.14	-1.02	1.23	-1.12	0.99	-1.53	-1.05	-1.11	-0.98	-1.14
0.51	0.10	0.90	0.54	0.62	0.47	0.97	0.63	0.06	1.00	0.65	0.19	0.21	0.29	0.05	0.27	0.30	0.76	0.99	0.42	0.10	0.87	0.02	0.13	0.15	0.24	0.12	0.01	0.17	0.95	0.16	0.57	0.67	0.08	0.99	0.87	0.90	0.76
0.95	-1.59	0.99	-1.55	-1.56	-1.83	1.05	-1.00	1.67	1.34	-1.43	1.87	-0.98	2.18	-2.09	1.07	-1.17	-1.04	-1.49	-1.07	2.82	-0.97	-1.72	-3.00	-1.83	-1.41	1.08	1.93	1.26	-1.61	-1.19	-1.26	-1.16	-3.11	Х	2.08	2.07	1.27
0.29	0.01	0.60	0.00	0.10	0.23	0.90	0.95	0.08	0.05	0.34	0.04	0.77	0.08	0.08	0.89	0.70	0.96	0.11	0.94	0.01	0.92	0.04	0.01	0.09	0.31	0.21	0.05	0.48	0.09	0.29	0.42	0.10	0.11	Х	0.20	0.58	x
1.07	-1.37	1.00	-1.32	-1.33	-1.65	1.05	0.05	1.76	0.17	-1.27	0.31	0.25	0.49	-1.79	1.18	0.08	0.00	-1.25	-1.17	2.22	0.10	-1.50	-2.33	-1.69	-1.41	1.35	1.93	1.20	-1.32	0.02	-1.19	-0.08	-2.32	N/A	0.49	0.55	0.07
0.11	0.22	0.02	0.23	0.23	0.18	0.01	1.04	0.09	1.17	0.16	1.57	1.23	1.69	0.30	0.11	1.25	1.04	0.24	0.10	0.60	1.07	0.22	0.67	0.15	0.00	0.27	0.00	0.06	0.30	1.21	0.07	1.08	0.79	N/A	1.60	1.52	1.21

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SPO2802	SPO2801	SPO2800	SPO2799	SPO2798	SPO2797	SPO2796	SPO2795	SPO2794	SPO2793	SPO2792	SPO2791	SPO2790	SPO2789	SPO2788	SPO2787	SPO2786	SPO2785	SPO2784	SPO2783	SPO2782	SPO2781	SPO2780	SPO2779	SPO2778	SPO2777	SPO2776	SPO2775	SPO2774	SPO2773	SPO2772	SPO2771	SPO2770	SPO2769	SPO2768	SPO2767
	lpcC	kdtA	moaA		glmU	gph-2			ivD	omp W	acsA	mccB	mccA	mvaB		nuoA	nuoB	nuoC		nuoD		nuoE			nuoF			nuoG		nuoH		nuol			nuoJ
bmp family protein	Ipopolysaccharide core biosynthesis mannosyltransferase (EC:2.4.1)	3-deoxy-D-manno-octulosonic acid transferase (EC:2)	molybdenum cofactor biosynthesis protein A	hypothetical protein	bifunctional N-acetylglucosamine-1-phosphate uridyltransferase/glucosamine-1-phosphate acetyltransferase (EC:2.7.7.23)	phosphoglycolate phosphatase (EC:3.1.3.18)	aminotransferase	hypothetical protein	isovaleryI-CoA dehydrogenase (EC:1.3.99.10)	outer membrane protein OmpW	acetyl-coenzyme A synthetase (EC:6.2.1.1)	methylcrotonyl-CoA carboxylase subunit beta (EC:6.4.1.4)	EC:6.4.1.4)	hydroxymethylglutaryl-CoA lyase (EC:4.1.3.4)	enoyl-CoA hydratase (EC:4.2.1.17)	NADH dehydrogenase subunit A (EC:1.6.5.3)	NADH dehydrogenase subunit B (EC:1.6.5.3)	NADH dehydrogenase subunit C (EC:1.6.5.3)	hypothetical protein	NADH dehydrogenase subunit D (EC:1.6.5.3)	lipoprotein	NADH dehydrogenase subunit E (EC:1.6.5.3)	hypothetical protein	hypothetical protein	NADH dehydrogenase I subunit F (EC:1.6.99.5)	hypothetical protein	hypothetical protein	NADH dehydrogenase subunit G (EC:1.6.5.3)	lipoprotein	NADH dehydrogenase subunit H (EC:1.6.5.3)	hypothetical protein	NADH dehydrogenase subunit I (EC:1.6.5.3)	hypothetical protein	4-carboxymuconolactone decarboxylase	NADH dehydrogenase subunit J (EC:1.6.5.3)
-3.77	1.25	-0.95	1.24	1.54	1.03	1.37	-2.20	-2.36	1.26	1.52	-1.89	0.97	1.23	1.10	-1.04	1.43	1.37	1.30	-1.15	1.30	1.25	1.56	-1.47	1.10	1.56	1.61	1.14	0.92	1.22	2.42	2.16	2.46	1.09	1.47	1.24
0.00	0.62	0.40	0.56	0.07	0.82	0.79	0.17	0.25	0.58	x	0.03	0.09	0.46	0.50	0.14	0.10	0.09	0.22	0.82	0.72	0.39	0.10	0.68	0.81	0.01	0.01	0.70	0.44	0.98	0.05	0.13	0.17	0.89	0.00	0.44
-2.60	1.33	-1.23	-1.59	-1.03	-1.21	1.60	1.23	-1.15	-1.25	3.38	-1.23	-1.12	1.17	1.37	1.22	1.05	1.09	1.23	1.61	0.99	1.28	-1.11	1.34	1.07	1.18	1.26	1.03	1.59	1.93	1.02	-1.22	1.27	1.29	1.22	-1.06
0.01	0.24	0.03	0.23	0.75	0.35	0.17	0.73	0.56	0.46	0.21	0.35	0.44	0.11	0.07	0.50	0.97	0.79	0.27	0.00	0.73	0.55	0.14	0.28	0.82	0.45	0.19	0.92	0.02	0.01	0.97	0.40	0.34	0.15	0.34	0.37
-3.19	1.29	-1.09	-0.18	0.26	-0.09	1.49	-0.49	-1.76	0.01	2.45	-1.56	-0.08	1.20	1.24	0.09	1.24	1.23	1.27	0.23	1.15	1.27	0.23	-0.06	1.09	1.37	1.44	1.09	1.26	1.58	1.72	0.47	1.87	1.19	1.35	0.09
0.58	0.04	0.14	1.42	1.29	1.12	0.12	1.72	0.61	1.26	0.93	0.33	1.04	0.03	0.14	1.13	0.19	0.14	0.04	1.38	0.15	0.02	1.34	1.41	0.02	0.19	0.18	0.05	0.33	0.36	0.70	1.69	0.60	0.10	0.13	1.15
-14.60	1.61	3.50	2.04	3.90	3.75	2.22	-1.34	-3.57	-1.96	8.08	-2.20	-2.51	-1.45	-1.89	-1.41	3.42	2.77	3.04	2.63	3.03	1.84	2.63	0.92	1.93	2.69	3.67	1.97	1.28	1.29	3.30	2.70	3.43	1.81	2.92	2.32
0.00	0.32	0.01	0.03	0.01	0.00	0.54	0.42	0.01	0.12	0.03	0.07	0.00	0.23	0.00	0.17	0.02	0.01	0.01	0.01	0.08	0.03	0.01	0.22	0.02	0.02	0.01	0.09	0.28	0.95	0.05	0.07	0.07	0.12	0.01	0.02
-2.11	1.30	1.02	-1.12	-1.09	-1.53	-1.49	1.27	-0.99	1.16	1.19	-1.16	1.10	1.22	1.36	1.34	-1.09	1.21	-1.23	-1.49	-1.13	1.20	-1.12	1.50	-1.07	1.10	0.99	0.99	-1.07	1.14	1.04	1.05	1.34	-1.17	1.25	1.14
0.01	0.29	1.00	0.51	0.27	0.01	0.21	0.60	0.90	0.52	0.83	0.32	0.33	0.52	0.06	0.16	0.10	0.25	0.08	0.02	0.46	0.14	0.02	0.11	0.19	0.70	0.51	0.65	0.54	0.71	0.85	0.69	0.13	0.52	0.40	0.57
-8.36	1.46	2.26	0.46	1.41	1.11	0.37	-0.04	-2.28	-0.40	4.64	-1.68	-0.71	-0.12	-0.27	-0.03	1.17	1.99	0.91	0.57	0.95	1.52	0.76	1.21	0.43	1.90	2.33	1.48	0.11	1.22	2.17	1.88	2.39	0.32	2.09	1.73
6.25	0.16	1.24	1.58	2.50	2.64	1.86	1.31	1.29	1.56	3.45	0.52	1.81	1.34	1.63	1.38	2.26	0.78	2.14	2.06	2.08	0.32	1.88	0.29	1.50	0.80	1.34	0.49	1.18	0.08	1.13	0.83	1.05	1.49	0.84	0.59
1.31	1.37	-1.18	-1.12	0.99	-1.29	1.11	1.25	-1.06	1.12	-0.98	-1.04	-1.16	1.23	1.07	-1.02	1.21	-1.07	1.21	1.65	-1.05	1.10	-1.10	-1.33	-1.56	-1.23	-1.16	-1.33	1.02	-1.11	-1.26	-1.52	-1.12	-1.04	0.98	1.03
0.14	0.15	0.21	0.46	0.83	0.05	0.71	0.50	0.89	0.59	0.94	0.94	0.13	0.14	0.56	0.90	0.26	0.96	0.19	0.00	0.73	0.17	0.41	0.26	0.03	0.10	0.19	0.32	0.90	0.16	0.17	0.04	0.42	0.40	0.29	
1.22	-1.05	-1.66	-1.12	-1.12	-1.42	-1.30	1.16	1.76	1.48	1.82	-1.31	-1.08	1.11	1.01	-1.00	0.92	1.05	-1.25	0.97	0.96	-1.10	0.98	-1.28	-1.71	-1.19	-1.65	-1.73	-1.34	-1.44	-1.47	-1.56	-1.22	-1.29	0.98	0.95
0.24	0.95	5 <u>0.04</u>	2 0.69	0.47	0.14	0.28			0.21	x	0.60	3 0.92	0.27	0.93	1.00	0.20	0.40	0.09	0.40	0.48	0.39	0.44	3 0.06	0.02			_	4 0.02		0.13	5 0.03	2 0.07	0.03	0.39	0.09
1.27	0.16	4 -1.42	-1.12	-0.07		-0.10	1.21	1 0.35	1.30	0.42	-1.18	-1.12	7 1.17	3 1.04	-1.01		-0.01	-0.02	1.31	-0.05	0.00		-1.31					-	-1.28	-1.37	-1.54	-1.17	3 -1.17		0.99
0.05	1.21	2 0.24	2 0.00	7 1.05		0 1.21	0.05	5 1.41	0.18	1.40	8 0.14	2 0.04	0.06	0.03	0.01	0.14	1 1.06	2 1.23	0.34	5 1.01	) 1.10	6 1.04	0.03	4 0.08	0.02	0.25	3 0.20	6 1.18		7 0.11	4 0.02	7 0.05	7 0.13		0.04
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### Chapter 10: Appendix

											iorA		kynU	bdh-2			pntA	pntB							argE-1									
hypothetical protein	DNA-binding protein	dipeptide ABC transporter substrate-binding protein	dipeptide ABC transporter permease	dipeptide ABC transporter permease	dipeptide ABC transporter ATP-binding protein	oligopeptide ABC transporter ATP-binding protein	hypothetical protein	dipeptidase domain-containing protein	type I secretion target repeat-containing protein	isoquinoline 1-oxidoreductase subunit beta	Isoquinoline 1-oxidoreductase subunit alpha (EC:1.3.99.16)	TetR family transcriptional regulator	kynureninase (EC:3.7.1.3)	D-beta-hydroxybutyrate denydrogenase (EC:1.1.1.30)	soxH protein-like protein	isoprenylcysteine carboxyl methyltransferase	(EC:1.6.1.2) (EC:1.6.1.2)	NAD(P) transnydrogenase subunit beta (EC:1.6.1.1) NAD(D) waarda da baaraan a da baaraan a	hypothetical protein	hypothetical protein	permease	peptue/nickel/opine uptake ABC transporter peptide/nickel/opine uptake ABC transporter	substrate-binding protein	binding protein	acetylornithine deacetylase (EC:3.5.1.16)	amidohydrolase	amidohydrolase	amidohydrolase	amidohydrolase	hypothetical protein	cytosine deaminase (EC:3.5.4.1)	sugar ABC transporter permease	sugar ABC transporter permease	sugar ABC transporter ATP-binding protein
-2.67	-2.79	-1.74	х	x	1.41	-1.03	-2.35	-1.17	1.05	-2.21	-1.70	-3.00	1.35	2.15	1.81	2.06	1.91	1.80	-1.88	-0.96	-1.63	-1.31	-2.25	-1.17	1.19	-1.07	1.07	1.09	1.22	-1.22	-1.68	-2.61	×	-4.18
0.00	X	x	×	×	×	x	0.00	0.81	0.74	0.03	0.70	0.00	0.81	0.02	0.08	0.01	0.17	0.23	0.19	0.30	0.34	0.73	0.00	0.16	0.29	0.76	0.78	0.56	0.22	0.42	0.56	0.02	×	0.00
-1.38	-1.32	-1.69	-1.40	х	1.18	-1.10	-1.31	1.08	1.02	-1.40	-1.46	-1.48	1.32	-1.39	-1.25	1.20	1.78	1.67	1.20	-3.60	1.60	-1.39	-2.67	-1.03	1.08	-1.61	-1.16	-1.57	-1.49	1.23	-1.12	-1.31	-1.69	-1.92
0.21	0.08	0.01	0.27	Х	x	0.57	0.35	0.83	0.79	0.15	0.03	0.01	0.66	0.07	0.77	0.16	0.03	0.02	0.33	0.01	0.49	0.19	0.03	0.50	0.50	0.04	0.59	0.09	0.05	0.33	0.61	0.34	0.23	0.09
-2.03	-2.06	-1.72	N/A	N/A	1.30	-1.07	-1.83	-0.04	1.04	-1.81	-1.58	-2.24	1.34	0.38	0.28	1.63	1.85	1.74	-0.34	-2.28	-0.01	-1.35	-2.46	-1.10	1.14	-1.34	-0.04	-0.24	-0.14	0.01	-1.40	-1.96	N/A	-3.05
0.64	0.74	0.03	N/A	N/A	0.12	0.04	0.52	1.13	0.02	0.41	0.12	0.76	0.02	1.77	1.53	0.43	0.06	0.07	1.54	1.32	1.62	0.04	0.21	0.07	0.05	0.27	1.12	1.33	1.36	1.23	0.28	0.65	N/A	1.13
-1.87	-0.94	3.32	2.10	х	-0.89	-1.09	-4.65	1.18	-1.58	-1.08	-1.76	-1.57	1.12	-2.82	1.20	5.36	1.74	1.82	-1.20	19.30	-1.13	-1.38	-1.81	-1.37	1.21	0.84	1.32	0.96	1.22	-1.15	-1.87	-3.37	-4.37	-8.66
0.01	0.64	0.01	0.11	×	0.86	0.46	0.00	0.57	0.07	0.93	0.66	0.04	0.81	0.01	0.18	0.01	0.05	0.22	0.78	0.00	0.68	0.13	0.01	0.02	0.34	0.16	0.05	0.46	0.85	0.31	0.35	0.01	x	0.00
1.40	-1.26	1.12	-1.04	х	-1.10	-0.98	-1.04	1.06	-1.13	-1.23	-1.10	-1.39	-1.61	1.95	1.41	1.24	1.76	1.35	-1.69	1.92	1.41	2.21	1.76	1.25	1.45	1.62	2.17	1.65	1.46	1.34	1.08	-1.27	×	-1.47
0.13	0.16	0.13	0.98	x	х	0.72	0.93	0.82	0.32	0.30	0.50	0.03	0.45	0.02	0.56	0.02	0.04	0.16	0.12	0.01	0.57	0.03	0.06	0.12	0.06	0.01	0.00	0.12	0.04	0.06	0.70	0.38	×	0.17
-0.24	-1.10	2.22	0.53	N/A	-0.99	-1.04	-2.85	1.12	-1.36	-1.16	-1.43	-1.48	-0.25	-0.44	1.31	3.30	1.75	1.59	-1.45	10.61	0.14	0.42	-0.03	-0.06	1.33	1.23	1.75	1.30	1.34	0.10	-0.40	-2.32	N/A	-5.07
1.64	0.16	1.10	1.57	N/A	0.11	0.05	1.81	0.06	0.23	0.08	0.33	0.09	1.37	2.39	0.11	2.06	0.01	0.24	0.25	8.69	1.27	1.80	1.79	1.31	0.12	0.39	0.43	0.35	0.12	1.25	1.48	1.05	N/A	3.60
1.32	1.35	-1.24	-1.38	Х	-1.02	-1.04	-1.43	-1.08	-1.99	1.29	1.20	1.37	-1.11	1.29	-1.07	-1.91	1.34	-1.25	1.35	-1.38	2.62	-1.18	1.65	1.23	1.28	1.35	1.02	-1.04	1.15	1.15	1.11	1.12	1.28	1.13
0.09	0.10	0.24	0.47	×	0.94	0.90	0.35	0.94	0.01	0.08	0.33	0.03	0.84	0.23	0.94	0.01	0.06	0.12	0.11	0.13	0.15	0.15	0.07	0.17	0.04	0.20	0.94	0.79	0.43	0.20	0.53	0.87	0.13	0.25
1.07	-1.06	-1.28	-1.05	х	х	х	1.91	-1.22	-1.45	1.50	1.94	1.61	-1.19	3.27	1.13	-2.05	1.65	-1.45	-1.10	1.10	1.70	1.37	3.36	1.33	1.32	1.45	1.10	1.46	1.54	1.19	1.29	2.63	×	1.29
0.81	0.75	0.50	0.49	x	х	х	0.25	0.79	0.12	0.07	0.03	0.02	0.83	0.01	0.69	0.03	0.01	0.03	0.97	0.82	0.53	0.35	0.02	0.04	0.15	0.08	0.33	0.32	0.10	×	0.37	0.19	×	0.17
1.20	0.15	-1.26	-1.22	N/A	N/A	N/A	0.24	-1.15	-1.72	1.40	1.57	1.49	-1.15	2.28	0.03	-1.98	1.50	-1.35	0.13	-0.14	2.16	0.10	2.51	1.28	1.30	1.40	1.06	0.21	1.35	1.17	1.20	1.88	N/A	1.21
0.13	1.21	0.02	0.17	N/A	N/A	N/A	1.67	0.07	0.27	0.11	0.37	0.12	0.04	0.99	1.10	0.07	0.16	0.10	1.23	1.24	0.46	1.28	0.86	0.05	0.02	0.05	0.04	1.25	0.20	0.02	0.09	0.75	N/A	0.08

SPO2817 SPO2818

SPO2813 SPO2814 SPO2815 SPO2816

SPO2819

SPO2810 SPO2811 SPO2812 SPO2809

SPO2805 SPO2806 SPO2807 SPO2808

SPO2803 SPO2804

SPO2836 SPO2837

SPO2835

SPO2830 SPO2831 SPO2832 SPO2833 SPO2834 SPO2826 SPO2827

SPO2828 SPO2829 SPO2823 SPO2824 SPO2825 SPO2820 SPO2821 SPO2822

SPO2874	SPO2873	SPO2872	SPO2871	SPO2870	SPO2869	SPO2868	SPO2867	SPO2866	SPO2865	SPO2864	SPO2863	SPO2862	SPO2861	SPO2860	SPO2859	SPO2858	SPO2857	SPO2856	SPO2855	SPO2854	SPO2853	SPO2852	SPO2851	SPO2850	SPO2849	SPO2848	SPO2847	SPO2846	SPO2845	SPO2844	SPO2843	SPO2842	SPO2841	SPO2840	SPO2839	SPO2838
cobF	cobA-2	cobB	cobM		cobL	cobK	cobJ	cobI	cobH	cobG	cobN	cobW								nucH							gatB						pepN		glcB	
precorrin 6A synthase (EC:2.1.1.152)	(EC:2.1.1.107)	cobyrinic acid a,c-diamide synthase	precorrin-4 C(11)-methyltransferase (EC:2.1.1.133)	cobalamin biosynthesis domain-containing protein	precorrin-6Y C5,15-methyltransferase (EC:2.1.1.132)	cobalt-precorrin-6x reductase (EC:1.3.1.54)	precorrin-5B C(17)-methyltransterase (EC:2.1.1.131)	precorrin-2 C(20)-methyltransferase (EC:2.1.1.130)	precorrin-8X methylmutase (EC:5.4.1.2)	CobG	cobaltochelatase subunit CobN (EC:6.6.1.2)	CobW	hypothetical protein	LysR family transcriptional regulator	3-hydroxyisobutyrate dehydrogenase	hypothetical protein	M24 family metallopeptidase	hypothetical protein	cobalt chelatase large subunit	thermonuclease (EC:3.1.31.1)	cobalt chelatase, CobS subunit (EC:6.6.1.2)	CzcN domain-containing protein	hypothetical protein	DnaJ domain-containing protein	BolA protein, truncation	hypothetical protein	aspar.yr.glutannyi-txxxA annidotransierase subunit B (EC:6.3.5)	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	sitori citatii denyu ogenase/reductase oxidoreductase	aminopeptidase (EC:3.4.11.2)	diguanylate cyclase	malate synthase G (EC:2.3.3.9)	hypothetical protein
1.27	2.19	1.65	1.73	1.36	1.65	-1.40	1.06	1.89	1.15	1.74	1.27	2.18	1.33	-2.13	3.20	2.97	2.19	-1.48	-2.00	-1.55	1.58	×	-1.95	-1.62	1.25	-1.07	1.36	-1.51	1.30	-3.26	-1.35	1.44	1.36	-1.31	2.69	1.96
0.18	0.78	0.32	0.06	0.89	0.88	0.65	0.87	0.17	0.95	0.01	0.69	0.00	0.63	×	0.04	0.05	0.15	0.01	0.01	0.07	0.06	×	x	0.01	0.71	0.28	0.88	0.18	0.54	0.17	0.55	0.11	0.08	0.27	0.00	0.77
1.57	1.47	1.73	1.61	1.37	1.48	1.06	1.50	1.43	1.58	1.13	1.36	1.56	1.25	-1.42	-1.16	1.70	-1.31	-1.27	1.19	1.12	1.07	-1.18	-1.10	-1.47	1.39	-1.78	1.68	1.13	1.40	-1.35	-1.20	1.07	1.17	-1.30	-1.43	-1.06
0.04	0.07	0.04	0.03	0.13	0.45	0.69	0.25	0.14	0.03	0.64	0.08	0.18	0.48	0.10	0.36	0.06	0.31	0.34	0.34	0.59	0.33	0.22	0.94	0.03	0.71	0.02	0.15	0.38	0.42	0.36	0.55	0.42	0.07	0.64	0.02	0.90
1.42	1.83	1.69	1.67	1.37	1.57	-0.17	1.28	1.66	1.37	1.44	1.32	1.87	1.29	-1.78	1.02	2.34	0.44	-1.38	-0.41	-0.22	1.33	N/A	-1.53	-1.55	1.32	-1.43	1.52	-0.19	1.35	-2.31	-1.28	1.26	1.27	-1.31	0.63	0.45
0.15	0.36	0.04	0.06	0.01	0.09	1.23	0.22	0.23	0.22	0.31	0.05	0.31	0.04	0.36	2.18	0.64	1.75	0.11	1.60	1.34	0.25	N/A	0.43	0.08	0.07	0.36	0.16	1.32	0.05	0.96	0.08	0.19	0.10	0.01	2.06	1.51
2.18	2.83	2.48	2.78	2.33	2.49	-1.39	4.92	2.68	1.21	3.74	-1.15	4.26	1.29	-1.31	-2.07	2.41	2.64	-1.43	1.12	2.29	5.48	-0.92	-1.22	1.10	-1.35	2.65	2.23	-1.83	-1.37	-5.43	1.26	2.13	1.20	-2.25	1.58	1.38
0.02	0.63	0.03	0.00	0.63	0.72	0.69	0.17	0.06	0.84	0.00	0.82	0.00	0.73	0.88	0.14	0.02	0.12	0.08	0.56	0.01	0.00	0.03	0.97	0.40	0.10	0.00	0.35	0.25	0.44	0.07	0.26	0.01	0.16	0.04	0.06	0.90
-1.34	-1.24	-1.49	-1.25	-1.10	-1.13	-1.22	-1.29	-1.67	-1.23	1.12	-1.06	-1.11	1.15	-1.62	-1.04	1.45	1.26	1.40	1.19	1.06	-1.16	-1.33	-1.06	0.96	1.11	-1.22	-1.19	-1.14	-1.15	1.14	1.48	1.31	1.03	1.23	2.44	1.79
0.06	0.43	0.02	0.20	0.53	0.49	0.24	0.07	0.03	0.10	0.17	0.23	0.16	0.41	0.37	0.92	0.03	0.22	0.11	0.15	0.91	0.11	0.23	0.95	0.27	0.93	0.08	0.22	0.63	0.68	0.69	0.06	0.06	0.93	0.71	0.05	0.31
0.42	0.80	0.50	0.77	0.62	0.68	-1.31	1.82	0.51	-0.01	2.43	-1.11	1.58	1.22	-1.47	-1.56	1.93	1.95	-0.02	1.16	1.68	2.16	-1.13	-1.14	1.03	-0.12	0.72	0.52	-1.49	-1.26	-2.15	1.37	1.72	1.12	-0.51	2.01	1.59
1.76	2.04	1.99	2.02	1.72	1.81	0.09	3.11	2.18	1.22	1.31	0.04	2.69	0.07	0.16	0.52	0.48	0.69	1.42	0.03	0.62	3.32	0.20	0.08	0.07	1.23	1.94	1.71	0.35	0.11	3.29	0.11	0.41	0.09	1.74	0.43	0.20
-1.18	-1.14	-1.15	-1.21	-1.69	-1.04	-1.37	-1.28	-1.46	-1.71	-1.96	-1.12	-1.18	-1.70	-1.11	-0.98	1.87	-1.12	1.05	-1.05	-1.20	1.20	-1.22	1.24	-1.24	1.77	-1.31	1.14	-1.04	1.04	-1.09	-1.13	-1.13	1.13	-1.06	-1.11	1.06
0.11	0.39	0.35	0.17	0.07	0.79	0.11	0.07	0.03	0.01	0.01	0.29	0.18	0.02	0.49	0.71	0.02	0.61	0.75	0.64	0.21	0.21	0.16	0.83	0.00	0.54	0.11	0.67	0.84	0.97	0.85	0.18	0.44	0.18	0.38	0.33	0.86
-2.18	-1.71	-2.20	-2.00	-2.25	-1.12	-1.25	-2.15	-2.34	-2.21	-2.50	-1.30	-1.68	-1.91	-1.50	-1.02	1.19	1.05	1.20	-1.30	-1.36	0.97	×	1.27	-1.41	1.25	-1.22	-1.12	-1.02	-1.17	1.89	1.35	1.02	1.59	1.38	-1.81	-1.13
0.04	0.12	0.01	0.02	0.05	0.75	0.32	0.00	0.03	0.02	0.00	0.38	0.11	0.00	0.12	0.54	0.19	0.89	0.37	0.01	0.06	0.09	x	0.72	0.04	0.63	0.04	0.40	x	0.30	0.26	0.20	0.97	0.01	0.11	0.02	0.82
-1.68	-1.43	-1.68	-1.61	-1.97	-1.08	-1.31	-1.72	-1.90	-1.96	-2.23	-1.21	-1.43	-1.81	-1.31	-1.00	1.53	-0.04	1.13	-1.18	-1.28	1.09	N/A	1.26	-1.33	1.51	-1.27	0.01	-1.03	-0.06	0.40	0.11	-0.05	1.36	0.16	-1.46	-0.03
	0.29	0.53	0.40	0.28	0.04	0.06	0.44	0.44	0.25	0.27	0.09	0.25	0.11	0.19	0.02	0.34	1.09	0.08	0.13	0.08	0.11	N/A	0.02	0.09	0.26	0.05	1.13	0.01	1.11	1.49	1.24	1.08	0.23	1.22	0.35	1.10

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#### M. Kirkwood

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SPO2913	SPO2912	SPO2911	SPO2910	SPO2909 dinF	SPO2908 arsC-	SPO2907 pyrD	SPO2906	SPO2905	SPO2904	SPO2903	SPO2902	SPO2901	SPO2900	SPO2899	SPO2898 phaG	SPO2897 phaF	SPO2896 phaE	SPO2895 phaD	SPO2894 phaC	SPO2893 phaA	SPO2892	SPO2891	SPO2890	SPO2889 mdoG	SPO2888	SPO2887	SPO2886	SPO2885	SPO2884	SPO2883	SPO2882	SPO2881	SPO2880	SPO2879	SPO2878	SPO2877	SPO2876	SPO2875
hypothetical protein	MerR family transcriptional regulator	thioesterase	thioesterase	DNA-damage-inducible protein F	I arsenate reductase (EC:1.20.4.1)	dihydroorotate dehydrogenase 2	hypothetical protein	hypothetical protein	Ser/Thr protein phosphatase/nucleotidase	TPR/sulfotransferase domain-containing protein	glutathione S-transferase	hypothetical protein	tRNA 2-selenouridine synthase	selenide, water dikinase	monovalent cation/H+ antiporter subunit G	monovalent cation/H+ antiporter subunit F	monovalent cation/H+ antiporter subunit E	monovalent cation/H+ antiporter subunit D	monovalent cation/H+ antiporter subunit C	monovalent cation/H+ antiporter subunit A	hypothetical protein	hypothetical protein	glucosyltransferase MdoH	glucan biosynthesis protein G	hypothetical protein	hypothetical protein	CAIB/BAIF family protein	acetolactate synthase, catabolic	hypothetical protein	hypothetical protein	dihydrodipicolinate synthase	xanthine dehydrogenase, large subunit	L-aspartate dehydrogenase (EC:1.4.1.21)	LysR family transcriptional regulator	hypothetical protein	hypothetical protein	hypothetical protein	ny pomencar protein

1.19	1.43	1.39	-1.19	1.33	1.58	1.20	1.13	1.24	-1.34	1.69	-2.18	-2.14	1.10	-1.43	1.01	1.26	-1.13	1.11	1.06	1.08	1.20	1.00	1.76	1.11	1.44	1.42	2.30	1.34	-1.09	1.14	-1.26	1.80	1.09	-1.54	1.67	2.60	2.04	1.59
0.94	0.23	0.02	0.55	×	0.85	0.06	0.89	0.70	0.05	0.27	0.03	0.03	0.29	0.59	0.67	0.20	0.46	0.67	0.93	0.78	0.89	0.97	0.14	0.24	0.23	0.06	0.21	0.04	0.78	0.32	0.73	0.00	×	0.01	0.02	0.01	0.22	0.04
1.28	-1.16	1.39	1.09	1.44	1.01	-1.11	-1.15	-1.20	-1.37	1.71	-1.52	-1.14	-1.25	1.08	-1.55	-1.22	-1.12	-1.16	1.15	-1.25	1.64	1.56	1.32	-1.24	1.04	-1.16	1.47	1.56	-1.07	-1.02	-1.02	-1.10	1.09	-1.22	1.42	1.33	1.54	1.08
0.62	0.45	0.01	0.44	0.60	1.00	0.16	0.39	0.28	0.04	0.46	0.01	0.65	0.04	0.85	0.13	0.29	0.17	0.28	0.71	0.04	0.15	0.33	0.19	0.38	0.92	0.10	0.03	0.12	0.90	0.87	1.00	0.86	0.57	0.01	0.03	0.34	0.29	0.92
1.24	0.14	1.39	-0.05	1.39	1.30	0.04	-0.01	0.02	-1.36	1.70	-1.85	-1.64	-0.08	-0.18	-0.27	0.02	-1.13	-0.02	1.11	-0.09	1.42	1.28	1.54	-0.06	1.24	0.13	1.89	1.45	-1.08	0.06	-1.14	0.35	1.09	-1.38	1.55	1.97	1.79	1.34
0.05	1.30	0.00	1.14	0.05	0.29	1.16	1.14	1.22	0.02	0.01	0.33	0.50	1.18	1.26	1.28	1.24	0.00	1.14	0.04	1.17	0.22	0.28	0.22	1.18	0.20	1.29	0.42	0.11	0.01	1.08	0.12	1.45	0.00	0.16	0.13	0.64	0.25	0.26
0.89	1.79	1.95	-1.15	2.24	2.13	2.60	2.66	2.72	-2.30	1.63	-2.34	-1.75	3.63	1.36	1.03	1.65	1.26	1.91	1.87	1.66	1.34	2.00	2.24	2.09	2.39	1.09	2.46	1.83	-3.62	1.69	-2.75	-1.51	-1.32	-1.03	3.53	2.32	1.13	1.09
0.63	0.07	0.02	0.59	0.21	0.72	0.02	0.03	0.18	0.01	0.45	0.04	0.05	0.00	0.33	0.71	0.08	0.76	0.01	0.11	0.18	0.78	0.77	0.01	0.00	0.03	0.99	0.02	0.02	0.04	0.02	0.26	0.19	0.96	0.62	0.00	0.04	0.58	0.01
1.30	1.42	1.08	-1.08	-1.13	-1.13	-1.41	-1.14	-1.14	1.52	1.10	1.13	1.40	-1.47	1.19	1.30	1.40	1.16	-1.10	-1.10	1.09	1.04	-1.15	1.04	-1.54	-1.75	-1.44	1.12	1.13	1.05	1.16	1.11	1.21	1.14	1.02	-1.08	1.60	2.68	1.53
0.47	0.07	0.47	0.30	0.80	0.76	0.01	0.47	0.48	0.20	0.90	0.30	0.05	0.04	0.45	0.50	0.02	0.57	0.18	0.29	0.81	0.95	0.69	0.92	0.01	0.02	0.08	0.55	0.20	0.93	0.21	0.88	х	0.70	0.82	0.38	0.02	0.01	0.08
1.09	1.61	1.52	-1.12	0.56	0.50	0.60	0.76	0.79	-0.39	1.37	-0.61	-0.18	1.08	1.28	1.17	1.53	1.21	0.41	0.39	1.38	1.19	0.43	1.64	0.28	0.32	-0.18	1.79	1.48	-1.29	1.43	-0.82	-0.15	-0.09	-0.01	1.23	1.96	1.91	1.31
0.21	0.18	0.43	0.03	1.69	1.63	2.01	1.90	1.93	1.91	0.27	1.74	1.58	2.55	0.09	0.14	0.13	0.05	1.51	1.49	0.29	0.15	1.58	0.60	1.82	2.07	1.27	0.67	0.35	2.34	0.27	1.93	1.36	1.23	1.03	2.31	0.36	0.78	0.22
1.03	1.06	-1.19	-1.86	-1.05	1.21	-1.19	-1.06	1.03	2.11	1.56	-1.12	-1.26	-2.02	1.37	-1.23	-1.12	1.38	1.33	1.51	1.34	1.76	1.44	1.92	-1.42	1.23	-1.81	1.23	-1.58	-1.22	1.16	-1.09	-1.13	1.26	1.09	-1.11	1.14	1.29	1.35
0.95	0.95	0.07	0.00	0.94	0.79	0.02	0.73	0.98	0.02	0.38	0.39	0.08	0.00	0.32	0.38	0.13	0.01	0.02	0.08	0.14	0.12	0.04	0.01	0.14	0.32	0.02	0.14	0.00	0.75	0.40	0.93	0.59	0.06	0.59	0.28	0.03	0.37	0.20
-1.04	1.34	-1.27	-1.29	-1.20	-1.08	-1.87	-1.58	-1.25	3.20	1.30	-1.00	-1.00	-3.18	1.30	1.43	0.99	1.60	1.22	1.22	1.78	1.26	1.19	2.15	-1.53	-1.50	-1.91	-1.13	-2.04	-0.98	-1.05	1.40	Х	-1.16	-1.02	-1.47	1.25	1.66	2.21
0.91	0.14	0.10	0.24	0.89	0.86	0.02	0.03	0.53	0.01	0.57	0.98	0.88	0.01	0.49	0.31	0.60	0.06	0.23	0.55	0.10	0.29	0.34	0.02	0.02	0.04	0.04	0.20	0.02	0.78	0.68	0.48	×	0.91	0.69	0.01	0.20	0.10	0.08
-0.01	1.20	-1.23	-1.58	-1.13	0.06	-1.53	-1.32	-0.11	2.66	1.43	-1.06	-1.13	-2.60	1.34	0.10	-0.07	1.49	1.28	1.37	1.56	1.51	1.32	2.04	-1.48	-0.14	-1.86	0.05	-1.81	-1.10	0.05	0.16	N/A	0.05	0.04	-1.29	1.20	1.48	1.78
1.04	0.14	0.04	0.28	0.08	1.15	0.34	0.26	1.14	0.55	0.13	0.06	0.13	0.58	0.04	1.33	1.05	0.11	0.06	0.15	0.22	0.25	0.13	0.12	0.06	1.37	0.05	1.18	0.23	0.12	1.11	1.25	N/A	1.21	1.06	0.18	0.06	0.18	0.43

### Chapter 10: Appendix

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												glyA	ppnK					prpE				cdd	deoA	deoB	add	upp						fabJ-1						
TrkA domain-containing protein	hypothetical protein	hypothetical protein	endoribonuclease L-PSP	AraC family transcriptional regulator	cupin	monooxygenase domain-containing protein	rhodanese-like domain-containing protein	hypothetical protein	alpha/beta hydrolase	hypothetical protein	hypothetical protein	serine hydroxymethyltransferase (EC:2.1.2.1)	Inorganic polyphosphate/ATP-NAD kinase (EC:2.7.1.23)	amidase (EC:3.5.1.4)	SCO1/SenC family protein	hypothetical protein	hypothetical protein	propionateCoA ligase (EC:6.2.1.17)	hypothetical protein	malic enzyme (EC:1.1.1.39)	hypothetical protein	cytidine deaminase (EC:3.5.4.5)	thymidine phosphorylase (EC:2.4.2.4)	phosphopentomutase (EC:5.4.2.7)	adenosine deaminase (EC:3.5.4.4)	uracil phosphoribosyltransferase (EC:2.4.2.9)	hypothetical protein	fatty oxidation complex subunit alpha	hypothetical protein	acetyl-CoA acetyltransferase (EC:2.3.1.9)	glutathione S-transferase	hypothetical protein	acyl-CoA dehydrogenase	MerR family transcriptional regulator				
-1.11	-1.51	-1.04	1.43	-1.91	2.30	3.33	1.19	-1.04	-1.74	1.16	-3.04	х	-1.00	1.75	1.72	1.81	1.74	5.89	-1.58	2.94	1.27	-1.31	-1.39	1.03	1.03	1.29	х	1.08	2.30	1.05	2.02	1.95	2.26	1.90	1.59	1.63	1.53	1.26
0.80	0.06	0.26	0.21	0.09	0.00	0.04	0.31	0.61	0.08	0.97	×	×	0.91	0.50	0.09	0.00	0.00	0.00	0.03	0.00	0.33	0.11	0.31	0.80	0.95	0.29	×	0.93	0.04	0.36	0.07	0.09	0.00	0.02	0.00	0.54	0.29	0.86
1.22	1.30	-1.31	1.60	-1.18	1.06	-1.07	-1.11	-1.04	1.15	0.99	1.66	х	1.23	1.00	1.23	1.06	-1.04	2.29	-1.24	1.78	-1.61	-1.08	-1.04	-1.05	-1.27	1.14	-1.02	-1.14	1.38	1.17	3.84	-1.40	-1.71	-1.66	-2.01	-1.40	-1.47	-1.73
0.10	0.15	0.07	0.03	0.36	0.86	0.73	0.78	0.90	0.16	0.97	0.09	x	0.13	0.96	0.27	0.75	0.71	0.01	0.40	0.04	0.00	0.60	0.87	0.69	0.07	0.24	x	0.31	0.11	0.20	0.00	0.05	0.06	0.04	0.02	0.35	0.17	0.14
0.05	-0.11	-1.18	1.52	-1.55	1.68	1.13	0.04	-1.04	-0.30	1.08	-0.69	N/A	0.12	1.38	1.48	1.44	0.35	4.09	-1.41	2.36	-0.17	-1.20	-1.22	-0.01	-0.12	1.22	N/A	-0.03	1.84	1.11	2.93	0.28	0.28	0.12	-0.21	0.12	0.03	-0.24
1.17	1.41	0.14	0.09	0.37	0.62	2.20	1.15	0.00	1.45	0.08	2.35	N/A	1.12	0.38	0.24	0.38	1.39	1.80	0.17	0.58	1.44	0.12	0.18	1.04	1.15	0.08	N/A	1.11	0.46	0.06	0.91	1.68	1.99	1.78	1.80	1.52	1.50	1.50
1.23	-2.92	-1.71	1.59	1.69	-6.48	-2.74	-1.76	-1.07	-1.82	1.36	-2.97	х	1.85	-2.32	-1.21	-1.59	-2.76	1.61	0.92	8.67	1.16	-1.34	-1.37	-1.12	-1.23	-1.39	x	1.21	-1.41	1.59	-3.47	-1.46	-2.11	-1.95	-3.17	-1.49	-1.47	-2.14
0.40	0.01	0.03	0.12	0.08	0.00	0.04	0.05	1.00	0.04	0.93	0.01	х	0.00	0.26	0.74	0.04	0.00	0.02	0.29	0.00	0.39	0.08	0.63	0.68	0.38	0.02	х	0.07	0.37	0.01	0.01	0.12	0.00	0.01	0.01	0.08	0.17	0.20
1.10	1.89	1.59	-1.21	1.21	1.34	1.20	-1.05	-1.08	-1.29	1.03	-1.43	х	1.72	1.73	1.45	1.15	1.47	13.70	1.33	1.52	1.31	-1.90	-1.48	-1.70	-1.62	1.00	х	1.20	-1.07	1.48	1.64	-1.09	1.63	1.47	1.27	-1.08	-1.11	1.32
0.21	0.02	0.01	0.20	0.33	0.47	0.31	0.80	0.74	0.05	1.00	0.56	×	0.05	0.16	0.05	0.44	0.05	0.00	0.42	0.05	0.16	0.01	0.24	0.10	0.08	0.66	×	0.19	0.76	0.06	0.04	0.46	0.02	0.02	0.20	0.55	0.25	0.27
1.17	-0.52	-0.06	0.19	1.45	-2.57	-0.77	-1.41	-1.08	-1.56	1.20	-2.20	N/A	1.79	-0.30	0.12	-0.22	-0.65	7.66	1.13	5.10	1.24	-1.62	-1.43	-1.41	-1.43	-0.20	N/A	1.21	-1.24	1.54	-0.92	-1.28	-0.24	-0.24	-0.95	-1.29	-1.29	-0.41
0.06	2.41	1.65	1.40	0.24	3.91	1.97	0.36	0.01	0.27	0.17	0.77	N/A	0.07	2.03	1.33	1.37	2.12	6.05	0.20	3.58	0.08	0.28	0.05	0.29	0.20	1.19	N/A	0.01	0.17	0.06	2.56	0.19	1.87	1.71	2.22	0.20	0.18	1.73
1.14	1.59	-1.35	1.13	1.06	4.28	2.24	1.14	1.28	1.04	1.42	-1.27	х	-1.12	1.51	1.10	1.27	-1.38	-1.22	-1.22	1.18	-1.03	1.02	1.29	1.17	-1.11	1.43	1.33	1.01	1.16	1.42	1.14	1.93	0.99	1.10	1.01	1.14	1.20	-1.06
0.08	0.01	0.03	0.68	0.76	0.01	0.02	0.44	0.41	0.80	0.64	0.14	×	0.48	0.32	0.08	0.03	0.12	0.06	0.19	0.19	0.62	0.89	0.70	0.37	0.32	0.02	0.10	0.95	0.09	0.04	0.22	0.02	0.74	0.60	0.91	0.65	0.37	0.52
-1.16	1.65	-1.09	-1.34	1.46	3.02	2.19	1.13	1.31	-2.06	1.82	-1.04	х	-1.33	2.91	-1.29	1.23	-1.16	-1.11	1.30	1.73	1.56	-1.49	1.08	1.04	-1.38	1.36	x	-1.05	-0.98	1.16	-1.36	3.44	2.99	2.65	2.38	2.19	2.61	2.50
0.40	0.01	0.81	0.12	0.23	0.01	0.08	0.51	0.03	0.04	0.56	0.98	х	0.28	0.04	0.16	0.55	0.29	0.09	0.31	0.03	0.04	0.12	0.67	0.94	0.07	0.07	x	0.58	0.46	0.08	0.08	0.01	0.00	0.01	0.04	0.06	0.07	0.07
-0.01	1.62	-1.22	-0.11	1.26	3.65	2.22	1.14	1.30	-0.51	1.62	-1.16	N/A	-1.23	2.21	-0.10	1.25	-1.27	-1.17	0.04	1.46	0.27	-0.24	1.19	1.11	-1.25	1.40	N/A	-0.02	0.09	1.29	-0.11	2.69	1.99	1.88	1.70	1.67	1.91	0.72
1.15	0.03	0.13	1.24	0.20	0.63	0.03	0.01	0.02	1.55	0.20	0.12	N/A	0.11	0.70	1.20	0.02	0.11	0.05	1.26	0.27	1.30	1.26	0.11	0.06	0.13	0.03	N/A	1.03	1.07	0.13	1.25	0.75	1.00	0.78	0.69	0.53	0.71	1.78

SPO2930 SPO2931 SPO2932 SPO2933 SPO2934

SPO2928 SPO2929

SP02921 SP02922 SP02923 SP02924 SP02924 SP02925 SP02926 SP02927

SPO2916 SPO2917 SPO2918 SPO2919 SPO2920

SPO2914 SPO2915

SPO2946 SPO2947 SPO2948 SPO2949 SPO2950 SPO2951 SPO2952

SPO2943 SPO2944 SPO2945

SPO2939 SPO2940 SPO2941 SPO2942 SPO2935 SPO2936 SPO2937 SPO2938

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dtd																					gatC	gatA			rpmG										mgtE		
D-tyrosyl-tRNA(Tyr) deacylase (EC:3.1)	nitroreductase	hypothetical protein	cytochrome b562	hypothetical protein	lipoprotein	tellurite resistance protein	hypothetical protein	RNA pseudouridylate synthase	cytidine and deoxycytidylate deaminase	hypothetical protein	hypothetical protein	peptidyl-arginine deiminase	hypothetical protein	guanylate cyclase	guanylate cyclase	metallo-beta-lactamase	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	C (EC:6.3.5)	asparty/glutamy-tKNA amidotransferase subunit A (EC:6.3.5) asparty/ollutamy/tRNA amidotransferase subunit	hypothetical protein	N-acetylmuramoyl-L-alanine amidase	50S ribosomal protein L33	hypothetical protein	hypothetical protein	hypothetical protein	LysR family transcriptional regulator	zinc-binding dehydrogenase oxidoreductase	DNA-binding protein	inositol monophosphatase	пучно хучестного апаглис стиуталинон учтотахс (ЕС:3.5.99.3)	guanine deaminase	magnesium transporter	5-formyltetrahydrofolate cyclo-ligase	Ser/Thr protein phosphatase
1.39	1.54	-1.06	1.33	1.58	1.17	1.33	1.11	-1.15	0.96	2.59	1.44	-1.07	-1.37	-1.99	-1.58	-1.21	1.03	1.65	Х	-1.38	1.58	1.95	0.90	1.21	-1.69	-1.43	1.27	-3.65	-1.22	-1.21	-1.45	-1.31	1.37	-1.29	1.34	1.23	1.13
0.01	0.02	0.54	0.12	0.17	0.63	0.68	0.79	0.27	0.85	0.00	0.22	0.94	0.27	0.35	0.08	0.76	0.98	0.02	x	0.19	0.03	0.01	0.02	0.64	0.25	0.03	0.99	0.03	0.04	0.83	0.01	0.59	0.35	0.08	0.08	0.78	0.61
-1.24	-1.47	1.03	1.10	-1.20	-1.49	-1.20	-1.14	-1.12	1.89	2.01	1.33	1.17	1.20	-1.13	-1.13	-1.37	-1.52	1.13	1.21	1.02	1.39	-1.11	-1.11	-1.18	1.58	1.34	-1.61	-1.66	-1.24	-1.38	1.22	1.12	1.33	1.00	1.28	1.55	1.09
0.09	0.08	0.85	0.53	0.12	0.04	0.12	0.49	0.33	0.60	0.05	0.13	0.70	0.14	0.75	0.64	0.16	0.02	0.07	0.25	0.94	0.30	0.33	0.58	0.28	0.16	0.20	0.13	0.02	0.07	0.10	0.09	0.47	0.27	0.97	0.06	0.26	0.77
0.08	0.04	-0.02	1.22	0.19	-0.16	0.07	-0.01	-1.14	1.42	2.30	1.39	0.05	-0.09	-1.56	-1.36	-1.29	-0.25	1.39	N/A	-0.18	1.49	0.42	-0.10	0.02	-0.05	-0.04	-0.17	-2.66	-1.23	-1.30	-0.12	-0.10	1.35	-0.15	1.31	1.39	1.11
1.32	1.51	1.05	0.12	1.39	1.33	1.27	1.13	0.01	0.47	0.29	0.05	1.12	1.29	0.43	0.23	0.08	1.28	0.26	N/A	1.20	0.10	1.53	1.01	1.20	1.64	1.39	1.44	1.00	0.01	0.09	1.34	1.22	0.02	1.14	0.03	0.16	0.02
2.05	1.58	1.56	1.81	1.77	1.17	1.35	1.21	2.82	0.93	2.85	1.66	-1.39	-1.94	-1.96	-2.55	-2.20	1.71	2.03	х	-1.40	4.69	3.74	1.27	1.75	2.66	-1.65	1.29	4.22	1.16	1.33	0.76	1.16	1.79	-1.40	-1.36	1.30	2.59
0.02	0.02	0.07	0.04	0.02	0.79	0.65	0.15	0.00	0.82	0.00	0.07	0.17	0.06	0.52	0.03	0.05	0.06	0.01	x	0.67	0.00	0.00	0.84	0.07	0.01	0.03	0.66	0.01	0.04	0.25	0.00	0.18	0.06	0.09	0.07	0.65	0.02
-1.03	1.19	1.19	1.37	1.16	1.05	1.13	1.05	1.22	1.32	1.13	1.37	1.17	1.16	1.27	1.32	1.30	-1.70	-1.55	Х	-1.22	-1.06	-1.08	1.11	-1.15	1.11	1.64	-1.13	1.65	-1.04	-1.22	2.49	2.23	2.48	-1.20	1.03	-1.11	-1.29
0.61	0.28	0.25	0.03	0.40	1.00	0.73	0.96	0.40	0.86	0.21	0.03	0.38	0.16	0.62	0.37	0.08	0.03	0.08	x	0.68	0.43	0.10	0.93	0.11	0.74	0.21	0.26	0.02	0.31	0.23	0.00	0.02	0.02	0.28	0.98	0.51	0.15
0.51	1.39	1.38	1.59	1.47	1.11	1.24	1.13	2.02	1.13	1.99	1.52	-0.11	-0.39	-0.35	-0.62	-0.45	0.01	0.24	N/A	-1.31	1.82	1.33	1.19	0.30	1.89	-0.01	0.08	2.94	0.06	0.06	1.63	1.70	2.14	-1.30	-0.17	0.10	0.65
1.54	0.20	0.19	0.22	0.31	0.06	0.11	0.08	0.80	0.19	0.86	0.15	1.28	1.55	1.62	1.94	1.75	1.71	1.79	N/A	0.09	2.88	2.41	0.08	1.45	0.78	1.65	1.21	1.29	1.10	1.28	0.86	0.54	0.35	0.10	1.20	1.21	1.94
1.06	-1.24	-1.55	1.17	1.04	-1.38	1.15	-1.08	-1.80	1.18	1.05	1.32	1.14	-1.22	1.19	1.11	1.09	1.38	1.10	1.99	1.24	-1.19	-1.53	1.31	1.03	-1.59	1.12	1.51	1.16	1.08	-1.66	2.22	2.05	1.48	-0.99	1.08	1.11	1.10
0.65	0.13	0.03	0.28	0.97	0.07	0.42	0.68	0.09	0.89	0.57	0.09	0.61	0.38	0.61	0.60	0.31	0.10	0.23	0.17	0.56	0.12	0.00	0.01	0.93	0.01	0.08	0.26	0.10	0.28	0.02	0.00	0.07	0.07	0.99	0.73	0.46	0.26
-1.14	-1.40	-1.74	1.04	1.08	-1.48	1.26	1.30	-1.93	1.23	-1.11	-1.10	1.10	-1.38	1.30	1.14	1.27	-1.04	-0.95	Х	1.45	-1.72	-1.51	1.23	-1.29	-2.21	1.21	2.20	2.09	1.19	-1.78	3.27	2.17	-0.97	-1.16	-1.03	-1.29	-1.31
0.49	0.12	0.01	0.97	0.94	0.03	0.44	0.14	0.11	0.83	0.39	0.94	0.42	0.37	0.52	0.57	0.06	0.69	0.26	x	0.33	0.01	0.01	0.83	0.34	0.01	0.10	0.09	0.01	0.02	0.05	0.00	0.04	0.15	0.54	0.86	0.59	0.18
-0.04	-1.32	-1.65	1.11	1.06	-1.43	1.21	0.11	-1.87	1.21	-0.03	0.11	1.12	-1.30	1.25	1.13	1.18	0.17	0.08	N/A	1.35	-1.46	-1.52	1.27	-0.13	-1.90	1.17	1.86	1.63	1.14	-1.72	2.75	2.11	0.26	-1.08	0.03	-0.09	-0.11
1.10	0.08	0.10	0.06	0.02	0.05	0.06	1.19	0.06	0.03	1.08	1.21	0.02	0.08	0.06	0.01	0.09	1.21	1.03	N/A	0.11	0.26	0.01	0.04	1.16	0.31	0.04	0.35	0.47	0.05	0.06	0.53	0.06	1.22	0.08	1.06	1.20	1.21

SPO2990 SPO2991 SPO2992

SPO2988 SPO2989

SPO2982 SPO2983 SPO2984 SPO2985 SPO2986 SPO2987

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SPO2975 SPO2976 SPO2977 SPO2978 SPO2979

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SPO2961 SPO2962 SPO2963

SPO2957 SPO2958 SPO2959 SPO2960 SPO2955 SPO2956

SPO2953 SPO2954

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SPO3031	SPO3030	SPO3029	SPO3028	SPO3027	SPO3026	SPO3025	SPO3024	SPO3023	SPO3022	SPO3021	SPO3020	SPO3019	SPO3018	SPO3017	SPO3016	SPO3015	SPO3014	SPO3013	SPO3012	SPO3011	SPO3010	SPO3009	SPO3008	SPO3005	SPO3004	SPO3003	SPO3002	SPO3001	SPO3000	SPO2999	SPO2998	SPO2997	3F02990	CDCCCC	SPO2995	SPO2994	SPO2993
eda-1									valS						metF	metR					putA	putR															
2-dehydro-3-deoxyphosphooctonate aldolase	etnux ABC transporter transmemorane ATF- binding protein	serineglyoxylate transaminase	hypothetical protein	histidinol-phosphate aminotransferase (EC:2.6.1.9)	strictosidine synthase	enoyl-CoA hydratase (EC;4.2.1.17)	adenylate cyclase	hypothetical protein	valyl-tRNA synthetase (EC:6.1.1.9)	phytanoyl-CoA dioxygenase	LysR family transcriptional regulator	xanthine dehydrogenase, large subunit	hypothetical protein	thioesterase	5,10-methylenetetrahydrofolate reductase	transcriptional regulator MetR	hypothetical protein	L4BD family NADP-dependent oxidoreductase	inositol monophosphatase	rhomboid family protein	carboxylate dehydrogenase (EC:1.5.1.12 1.5.99.8)	proline dehydrogenase hifunctional proline dehydrogenase/nyrroline-5-	hypothetical protein	ABC transporter ATP-binding protein/permease	LysM domain/BON superfamily protein	AMP-binding protein (EC:2.3.1.86)	lipase	hypothetical protein	hypothetical protein	hypothetical protein	binding protein	permease	pertitease peptide/nickel/opine uptake ABC transporter	peptide/nickel/opine uptake ABC transporter	peptide/nickel/opine uptake ABC transporter substrate-binding protein	X-Pro dipeptidyl-peptidase	MOFRL family protein
1.68	-1.19	1.50	-1.20	1.16	1.62	1.94	1.97	-1.72	1.15	3.15	×	1.02	1.06	-1.34	-74.30	-1.94	-5.00	-1.45	-1.15	1.35	-1.08	1.14	1.32	1.42	-1.26	-1.42	-1.09	1.00	2.00	-1.35	-1.99	-1.51	-1.39	- 50	-1.32	3.46	1.55
0.83	0.27	0.32	0.30	0.57	0.06	0.03	0.01	0.04	0.97	0.03	x	0.49	0.96	0.70	0.00	0.09	0.00	0.02	0.87	0.02	0.36	0.06	0.34	0.92	0.03	0.10	0.88	0.88	0.03	0.39	0.23	0.49	0.15	0 12	0.04	0.04	0.63
1.57	-1.05	1.18	1.67	1.41	1.33	1.17	-1.26	-1.05	1.22	-1.24	x	1.23	1.11	-3.15	-22.40	-1.36	-1.27	-1.13	1.54	1.20	1.29	-1.02	1.06	1.95	-2.53	-1.86	-1.36	-1.05	1.03	1.05	-1.36	-1.51	-2.12	2 12	-2.66	1.57	-1.24
0.51	0.51	0.39	0.21	0.04	0.04	0.65	0.07	0.67	0.61	0.89	×	0.12	0.58	0.00	0.02	0.11	0.01	0.77	0.26	0.04	0.24	0.93	0.36	0.04	0.08	0.17	0.22	0.53	0.87	0.86	0.39	0.15	0.01	001	0.00	0.05	0.54
1.63	-1.12	1.34	0.24	1.29	1.48	1.56	0.36	-1.39	1.19	0.96	N/A	1.13	1.09	-2.25	-48.35	-1.65	-3.14	-1.29	0.20	1.28	0.11	0.06	1.19	1.69	-1.90	-1.64	-1.23	-0.03	1.52	-0.15	-1.68	-1.51	-1.00	1 0 6	-1.99	2.52	0.16
0.05	0.07	0.16	1.44	0.13	0.15	0.39	1.62	0.34	0.04	2.20	N/A	0.11	0.03	0.90	25.95	0.29	1.87	0.16	1.35	0.08	1.19	1.08	0.13	0.26	0.64	0.22	0.14	1.02	0.49	1.20	0.32	0.00	0.27	76.0	0.67	0.95	1.40
1.42	1.47	1.30	-1.23	6.37	4.55	-1.58	-1.73	-2.45	3.16	1.69	х	0.95	1.03	9.78	4.39	1.35	-12.00	-1.56	1.64	2.16	-1.81	-2.28	-1.98	2.62	2.80	-4.22	-1.93	-1.19	2.05	-1.10	-1.48	-1.60	-1.30	1 20	-1.40	-1.76	1.14
0.88	0.06	0.27	0.85	0.00	0.00	0.00	0.36	0.02	0.46	0.58	x	0.59	0.82	0.00	0.00	0.10	0.00	0.26	0.68	0.00	0.03	0.01	0.00	0.53	0.02	0.06	0.52	0.78	0.05	0.68	0.27	0.44	0.20	0,00	0.04	0.19	0.76
2.75	-1.04	1.07	-1.21	-1.15	-1.41	1.16	1.61	-1.02	-1.35	-1.20	X	1.31	1.21	1.80	2.56	1.21	1.21	1.93	-1.18	1.25	-1.39	-1.19	1.92	-1.31	1.58	1.47	1.50	1.17	1.33	-1.07	1.42	1.31	1.14	1 1 /	1.26	-1.00	1.27
0.14	0.39	0.59	0.55	0.29	0.03	0.63	0.03	0.82	0.07	0.85	×	0.15	0.42	0.02	0.01	0.06	0.14	0.21	0.56	0.05	0.04	0.31	0.09	0.09	0.15	0.34	0.15	0.23	0.11	0.83	0.15	0.33	0.07	ГЭ ()	0.09	0.84	0.43
2.09	0.22	1.19	-1.22	2.61	1.57	-0.21	-0.06	-1.74	0.91	0.25	N/A	1.13	1.12	5.79	3.48	1.28	-5.40	0.19	0.23	1.71	-1.60	-1.74	-0.03	0.66	2.19	-1.38	-0.22	-0.01	1.69	-1.09	-0.03	-0.15	-0.12	0 10	-0.07	-1.38	1.21
0.67	1.26	0.12	0.01	3.76	2.98	1.37	1.67	0.72	2.26	1.45	N/A	0.18	0.09	3.99	0.92	0.07	6.61	1.75	1.41	0.46	0.21	0.55	1.95	1.97	0.61	2.85	1.72	1.18	0.36	0.02	1.45	1.46	1.20	1 76	1.33	0.38	0.07
1.38	1.28	1.42	1.32	1.19	-1.11	1.37	-1.44	-1.19	-1.15	-1.21	х	1.65	1.39	-1.18	-1.50	1.14	2.59	1.05	-1.65	1.38	-1.11	1.04	1.17	1.19	1.01	1.13	1.61	1.31	-1.13	-1.05	1.39	1.47	1.37	1 27	1.27	-1.97	1.09
0.65	0.16	0.05	0.09	0.09	0.32	0.21	0.02	0.35	0.45	0.90	×	0.01	0.04	0.10	0.03	0.24	0.00	0.93	0.04	0.06	0.84	0.63	0.45	0.40	0.90	0.83	0.10	0.19	0.47	0.88	0.12	0.08	0.14	014	0.41	0.04	0.82
1.20	1.12	2.30	-1.38	-1.87	-1.62	2.60	х	-1.40	-1.20	1.72	х	1.53	1.08	1.66	1.14	1.13	5.36	1.42	-2.19	1.24	-1.02	-1.91	-1.47	1.33	-1.99	2.02	1.68	1.68	1.43	-1.13	1.49	1.62	1.39	1 50	2.13	-2.29	1.41
0.83	0.67	0.03	х	0.01	0.04	0.14	×	0.30	0.37	0.44	х	0.09	0.60	0.09	0.99	0.18	0.00	0.49	0.07	0.07	×	0.04	0.18	0.44	0.10	0.10	0.12	0.11	0.27	0.93	0.18	0.03	C2.0	20.0	0.05	0.05	0.41
1.29	1.20	1.86	-0.03	-0.34	-1.37	1.99	N/A	-1.30	-1.18	0.26	N/A	1.59	1.24	0.24	-0.18	1.14	3.98	1.24	-1.92	1.31	-1.07	-0.44	-0.15	1.26	-0.49	1.58	1.65	1.50	0.15	-1.09	1.44	1.55	1.40	1 /0	1.70	-2.13	1.25
0.09	0.08	0.44	1.35	1.53	0.25	0.62	N/A	0.11	0.03	1.47	N/A	0.06	0.16	1.42	1.32	0.01	1.39	0.19	0.27	0.07	0.05	1.48	1.32	0.07	1.50	0.45	0.03	0.18	1.28	0.04	0.05	0.08	0.11	011	0.43	0.16	0.16

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SPO3067	SPO3066	SPO3065	SPO3064	SPO3063	SPO3062	SPO3060	SPO3059	SPO3058	SPO3057	SPO3056	SPO3055	SPO3054	SPO3053	SPO3052	SPO3051	SPO3050	SPO3049	SPO3048	SPO3047	SPO3046	SPO3045	SPO3044	SPO3043	SPO3042	SPO3041	SPO3040	SPO3039	SPO3038	SPO3037	SPO3036	SPO3035	SPO3034	SPO3033	SPO3032	
	apt					mtaP																	glnQ										zwf-2	edd	
oxidoreductase, FAD-binding	adenine phosphoribosyltransferase (EC:2.4.2.7)	hypothetical protein	hypothetical protein	transmembrane efflux protein	hypothetical protein	(EC:2.4.2.28)	s' matheriticadappoint the shorthony land	sulfate transporter family protein	streptogramin acetyltransferase	hypothetical protein	acetyltransferase	hypothetical protein	NUDIX domain-containing protein	hypothetical protein	short chain dehydrogenase/reductase oxidoreductase	hypothetical protein	protein	dipeptide ABC transporter permease di peptide ABC transporter substrate-binding	dipeptide ABC transporter permease	oligopeptide ABC transporter ATP-binding protein	glutamine amidotransferase	LysR family transcriptional regulator	glutamine ABC transporter ATP-binding protein	polar amino acid ABC transporter permease	polar amino acid ABC transporter permease	polar amino acid ABC transporter substrate-binding protein	polar amino acid ABC transporter substrate-binding protein	(EC:2.7.3)	LysR family transcriptional regulator	metallo-beta-lactamase	aspartate kinase (EC:2.7.2.4)	phosphoenolpyruvate-protein phosphotransterase (EC:2.7.3.9)	EC:1.1.1.49)	phosphogluconate dehydratase (EC:4.2.1.12)	(EC:4.1.2.14 4.1.3.16)
1.31	-3.18	-26.30	-11.90	1.22	х	1.63	2.63	0.97	2.10	1.40	-1.56	1.30	1.07	-1.22	0.95	Х	1.14	1.13	Х	-1.40	-1.00	-1.06	-1.37	х	-2.00	1.14	1.21	-1.20	-1.56	0.99	1.10	-1.19	1.10	1.31	
0.65	0.03	0.01	0.01	0.78	×	0.00	0.01	0.91	0.04	0.66	0.14	0.05	0.97	0.22	0.14	×	0.96	Х	×	0.54	0.96	×	0.17	×	0.53	0.95	0.53	0.74	0.10	0.42	0.72	0.24	0.86	0.28	
-1.07	1.40	-3.92	-1.16	1.21	-1.34	1.49	-1.14	1.83	1.44	1.58	1.30	-2.27	-1.20	-1.53	1.01	1.12	-1.33	-1.11	1.12	-1.79	-1.04	-1.11	1.01	-1.15	-1.14	-1.12	1.20	1.19	-1.40	-1.42	1.68	1.84	1.29	1.17	
0.84	0.24	0.08	0.39	0.60	×	0.03	0.50	0.13	0.27	0.13	0.05	0.01	0.72	0.01	0.93	0.40	0.23	0.36	0.64	0.00	0.90	0.85	0.98	0.39	0.81	0.95	0.18	0.73	0.00	0.10	0.07	0.08	0.34	0.75	
0.12	-0.89	-15.11	-6.53	1.22	N/A	1.56	0.75	1.40	1.77	1.49	-0.13	-0.49	-0.06	-1.38	0.98	N/A	-0.10	0.01	N/A	-1.60	-1.02	-1.09	-0.18	N/A	-1.57	0.01	1.21	-0.01	-1.48	-0.22	1.39	0.33	1.20	1.24	
1.19	2.29	11.19	5.37	0.01	N/A	0.07	1.89	0.43	0.33	0.09	1.43	1.79	1.14	0.16	0.03	N/A	1.24	1.12	N/A	0.19	0.02	0.03	1.19	N/A	0.43	1.13	0.01	1.20	0.08	1.20	0.29	1.52	0.10	0.07	
1.05	-2.15	-9.57	-1.55	1.94	х	1.64	1.83	2.64	2.27	1.58	1.08	-1.16	-1.20	1.36	1.38	-0.88	-2.42	-1.45	1.25	-2.26	-0.97	-1.15	-1.43	х	-1.20	-1.71	-1.29	-1.11	-0.99	-1.97	4.03	-1.37	0.84	0.95	
0.81	0.08	0.00	0.02	0.07	x	0.01	0.00	0.63	0.02	0.19	0.46	0.22	0.73	0.16	0.10	0.10	0.67	0.81	0.34	0.02	0.67	0.85	0.31	x	1.00	0.85	0.36	0.85	0.34	0.03	0.02	0.30	0.41	0.06	
1.02	-1.59	-2.13	1.05	1.27	-1.26	1.53	1.08	1.20	1.46	1.55	-1.45	-1.18	-1.25	1.10	0.98	-1.16	1.33	1.24	-1.30	1.29	-1.18	1.14	-1.25	-1.07	1.08	1.11	1.42	1.28	1.25	1.13	1.00	1.05	2.95	3.26	
0.99	0.12	0.03	0.77	0.25	х	0.04	0.28	0.59	0.06	0.05	0.04	0.15	0.49	0.80	0.48	0.71	0.21	0.46	x	0.14	0.49	0.83	0.56	х	0.89	0.94	0.13	0.45	0.03	0.49	0.68	0.83	0.03	0.01	
1.04	-1.87	-5.85	-0.25	1.61	N/A	1.59	1.46	1.92	1.87	1.57	-0.19	-1.17	-1.23	1.23	1.18	-1.02	-0.55	-0.11	-0.03	-0.49	-1.07	-0.01	-1.34	N/A	-0.06	-0.30	0.06	0.09	0.13	-0.42	2.52	-0.16	1.90	2.11	
0.02	0.28	3.72	1.30	0.34	N/A	0.05	0.38	0.72	0.41	0.02	1.27	0.01	0.03	0.13	0.20	0.14	1.88	1.35	1.28	1.78	0.11	1.15	0.09	N/A	1.14	1.41	1.36	1.20	1.12	1.55	1.52	1.21	1.05	1.16	
-1.09	1.56	1.33	1.29	1.20	1.05	-1.41	-1.22	-1.10	-1.35	0.99	1.16	1.82	1.42	1.09	1.45	-0.96	-1.11	-1.28	-1.17	1.25	-1.32	-1.19	-1.34	1.08	-1.33	-1.43	1.08	1.19	-1.42	1.61	-1.42	1.51	-1.31	1.12	
0.80	0.06	0.08	0.15	0.22	х	0.01	0.20	0.44	0.02	0.55	0.15	0.02	0.43	0.73	0.02	0.75	0.71	0.19	0.60	0.27	0.37	0.83	0.48	0.14	0.73	0.76	0.60	0.54	0.02	0.01	0.16	0.20	0.56	0.20	
-1.59	1.21	1.64	-0.99	1.03	х	-1.25	-1.10	-1.58	-1.28	1.07	-1.29	1.48	1.47	1.19	1.39	1.14	-1.03	X	-1.06	1.41	-1.58	1.46	-1.11	1.21	1.32	1.71	-1.13	-1.12	-1.70	1.62	-1.93	-1.33	-1.21	1.67	
0.36	0.16	0.07	0.82	0.79	Х	0.39	0.91	0.39	0.17	0.75	0.17	0.02	0.38	0.57	0.08	0.29	0.60	X	х	0.02	0.22	х	0.95	х	0.32	х	0.67	0.97	0.02	0.05	0.01	0.47	0.63	0.57	
-1.34	1.39	1.49	0.15	1.12	N/A	-1.33	-1.16	-1.34	-1.32	1.03	-0.07	1.65	1.45	1.14	1.42	0.09	-1.07	N/A	-1.12	1.33	-1.45	0.14	-1.23	1.15	-0.01	0.14	-0.02	0.03	-1.56	1.62	-1.68	0.09	-1.26	1.40	
0.25	0.18	0.16	1.14	0.09	N/A	0.08	0.06	0.24	0.04	0.04	1.23	0.17	0.03	0.05	0.03	1.05	0.04	N/A	0.05	0.08	0.13	1.33	0.12	0.06	1.33	1.57	1.11	1.16	0.14	0.01	0.26	1.42	0.05	0.28	

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### Chapter 10: Appendix

PO3104	PO3103	PO3102	PO3101	PO3100	PO3099	PO3098	PO3097	PO3096	PO3095	PO3094	PO3093	PO3092	PO3091	PO3090	PO3089	PO3088	PO3087	PO3086	PO3085	PO3084	PO3083	PO3082	PO3081	PO3080	PO3079	PO3078	PO3077	PO3076	PO3075	PO3074	PO3073	PO3072	PO3071	PO3070	PO3069	PO3068
	fhs-2						mmsB-2														dxnH	dxnG			topA			ctaC	cyoE	ctaG	ctaE		thrC			
MOSC domain-containing protein	formatetetrahydrofolate ligase (EC:6.3.4.3)	chorismate mutase	bifunctional 5,10-methylene-tetrahydrofolate dehydrogenase/5,10-methylene-tetrahydrofolate cyclohydrolase	hypothetical protein	hypothetical protein	DEAD/DEAH box helicase	3-hydroxyisobutyrate dehydrogenase (EC:1.1.1.31)	hypothetical protein	hypothetical protein	lipoprotein	hypothetical protein	TadC family type II/IV secretion system protein	TadC family type II/IV secretion system protein	type II/IV secretion system protein	ATPase	OmpA family protein	type II/III secretion system protein	CpaB family protein	hypothetical protein	Slt family transglycosylase	(EC:2.8.3.6)	3-oxoadipate CoA-succinyl transferase subunit alpha (EC:2.8.3.6)	glyoxalase	TetR family transcriptional regulator	DNA topoisomerase I (EC:5.99.1.2)	DNA processing protein DprA	TldD/PmbA family protein	cytochrome c oxidase subunit II (EC:1.9.3.1)	protoheme IX farnesyltransferase (EC:2.5.1)	cytochrome C oxidase assembly protein	cytochrome c oxidase subunit III (EC:1.9.3.1)	SURF1 family protein	threonine synthase (EC:4.2.3.1)	M16 family peptidase	ribosomal-protein-alanine acetyltransferase	hypothetical protein
1.26	12.00	×	×	1.21	-1.32	-2.07	1.05	1.62	-2.21	-2.18	-1.47	1.62	1.73	1.76	1.31	1.13	-1.60	1.62	1.28	-2.55	1.58	1.42	-1.28	-1.23	-1.18	1.21	1.04	4.28	4.94	3.15	2.48	-0.98	1.17	1.83	1.04	-1.43
0.61	0.00	x	×	0.01	0.25	0.01	0.81	0.52	0.00	0.01	0.02	0.51	0.66	0.50	0.89	0.85	0.01	0.56	0.84	0.01	0.21	0.77	0.36	0.62	0.71	0.60	0.97	0.03	0.06	0.02	0.01	0.93	0.10	0.07	0.06	0.61
1.00	2.10	х	×	1.14	1.23	-1.37	-1.12	-1.02	-1.61	-2.10	-1.57	1.20	1.27	1.16	1.03	-1.23	-1.12	-1.19	1.59	-1.13	1.56	1.06	-1.36	1.25	-1.07	1.99	-1.26	2.03	1.50	1.12	1.17	1.39	1.00	1.15	1.13	1.03
0.81	0.04	x	×	0.35	0.27	0.23	0.16	0.45	0.14	0.08	0.15	0.55	0.61	0.30	0.92	0.17	0.29	0.38	0.11	0.48	0.21	0.80	0.29	0.44	0.74	0.06	0.33	0.08	0.05	0.61	0.18	0.08	0.60	0.43	0.73	0.83
1.13	7.05	N/A	N/A	1.18	-0.05	-1.72	-0.04	0.30	-1.91	-2.14	-1.52	1.41	1.50	1.46	1.17	-0.05	-1.36	0.22	1.44	-1.84	1.57	1.24	-1.32	0.01	-1.13	1.60	-0.11	3.16	3.22	2.14	1.83	0.21	1.09	1.49	1.09	-0.20
0.13	4.95	N/A	N/A	0.04	1.28	0.35	1.09	1.32	0.30	0.04	0.05	0.21	0.23	0.30	0.14	1.18	0.24	1.41	0.16	0.71	0.01	0.18	0.04	1.24	0.05	0.39	1.15	1.13	1.72	1.02	0.66	1.18	0.09	0.34	0.04	1.23
2.63	1.10	Х	x	0.99	1.37	-2.68	-2.95	-2.22	-1.89	-2.41	-2.65	-1.71	-2.13	-2.62	-3.21	1.04	-2.56	-1.77	0.68	-3.04	3.49	2.91	1.10	1.96	3.92	-3.63	2.13	8.56	13.10	5.20	2.82	3.97	3.98	4.93	-2.54	-1.43
0.16	0.63	x	×	0.95	0.04	0.00	0.00	0.19	0.01	0.01	0.01	0.20	0.36	0.22	0.01	0.02	0.00	0.21	0.00	0.00	0.01	0.03	0.52	0.62	0.08	0.00	0.01	0.01	0.00	0.02	0.00	0.03	0.00	0.01	0.00	0.71
1.45	-2.47	x	×	-1.41	-1.35	1.13	-1.17	-1.12	-1.10	-1.12	1.09	-1.03	1.01	1.03	0.99	-1.47	-1.77	1.02	1.32	1.07	1.59	2.12	-2.01	-1.21	-1.07	3.76	0.99	1.11	0.98	1.16	1.20	-1.46	-1.70	-1.14	1.01	2.36
0.25	0.01	х	×	0.09	0.08	0.28	0.12	0.36	0.08	0.31	0.90	0.87	0.90	0.95	0.74	0.02	0.02	0.87	0.40	0.66	0.01	0.06	0.04	0.46	0.49	0.01	0.65	0.80	0.26	0.63	0.24	0.11	0.02	0.35	0.95	0.14
2.04	-0.69	N/A	N/A	-0.21	0.01	-0.78	-2.06	-1.67	-1.50	-1.77	-0.78	-1.37	-0.56	-0.80	-1.11	-0.22	-2.17	-0.38	1.00	-0.99	2.54	2.52	-0.46	0.38	1.43	0.06	1.56	4.84	7.04	3.18	2.01	1.26	1.14	1.90	-0.77	0.47
0.59	1.79	N/A	N/A	1.20	1.36	1.91	0.89	0.55	0.39	0.65	1.87	0.34	1.57	1.83	2.10	1.26	0.40	1.40	0.32	2.06	0.95	0.40	1.56	1.59	2.50	3.70	0.57	3.73	6.06	2.02	0.81	2.72	2.84	3.04	1.78	1.90
1.03	-1.53	х	×	-1.01	1.22	-1.39	-2.46	-1.80	-1.17	-1.18	-1.70	-1.21	-1.47	-1.48	-1.98	-1.63	-1.22	-1.34	1.20	1.03	1.43	1.02	-1.40	1.31	-1.26	-1.19	1.52	1.20	1.08	-1.96	-1.29	-1.25	-1.85	-1.34	-1.14	1.68
0.98	0.11	x	×	0.44	0.26	0.16	0.00	0.09	0.26	0.18	0.03	0.56	0.24	0.12	0.02	0.02	0.18	0.06	0.57	0.75	0.01	0.67	0.03	0.24	0.37	0.32	0.02	0.13	0.53	0.01	0.00	0.27	0.02	0.04	0.41	0.30
1.02	-1.77	×	×	-1.20	-1.21	1.15	-2.64	-2.38	-1.62	0.88	1.09	-1.07	1.03	0.97	-1.26	-1.74	-1.38	1.49	1.20	-1.20	2.27	2.41	-2.61	1.12	-1.48	-1.21	1.41	1.63	0.95	-1.60	0.88	-2.23	-2.30	-1.65	1.16	1.84
0.91	0.25	x	×	0.09	0.72	0.26	0.01	0.03	0.03	0.03	0.81	0.68	0.92	0.70	0.05	0.01	0.14	0.44	0.92	0.58	0.07	0.00	0.01	0.45	0.15	0.76	0.46	0.13	0.03	0.11	0.04	0.03	0.01	0.10	0.68	0.29
1.03	-1.65	N/A	N/A	-1.11	0.01	-0.12	-2.55	-2.09	-1.40	-0.15	-0.31	-1.14	-0.22	-0.25	-1.62	-1.69	-1.30	0.08	1.20	-0.09	1.85	1.72	-2.01	1.22	-1.37	-1.20	1.47	1.42	1.02	-1.78	-0.21	-1.74	-2.08	-1.50	0.01	1.76
0.01	0.12	N/A	N/A	0.10	1.22	1.27	0.09	0.29	0.23	1.03	1.40	0.07	1.25	1.23	0.36	0.06	0.08	1.42	0.00	1.12	0.42	0.70	0.61	0.10	0.11	0.01	0.06	0.21	0.06	0.18	1.08	0.49	0.22	0.16	1.15	0.08

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SPO3142	SPO3141	SPO3140	SPO3139	SPO3138	SPO3137	SPO3136	SPO3135	SPO3134	SPO3133	SPO3132	SPO3131	SPO3130	SPO3129	SPO3128	SPO3127	SPO3126	SPO3125	SPO3124	SPO3123	SPO3122	SPO3121	SPO3120	SPO3119	SPO3118	SPO3117	SPO3116	SPO3115	SPO3114	SPO3113	SPO3112	SPO3111	SPO3110	SPO3109	SPO3108	SPO3107	SPO3106	SPO3105
						ileS		ogt			pcs	xerC		fsaB	priA										ruvC	ruvA	ruvB			tolQ			tolB	pal			ftsH
hypothetical protein	acetyltransferase	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	isoleucyl-tRNA synthetase (EC:6.1.1.5)	T4 family peptidase	methylated-DNAprotein-cysteine methyltransferase (EC:2.1.1.63)	hypothetical protein	hypothetical protein	phosphatidylcholine synthase (EC:2.7.8.24)	site-specific tyrosine recombinase XerC	hypothetical protein	translaldolase (EC:2.2.1.2)	primosome assembly protein PriA	sigma factor regulator PrtR	ECF subfamily RNA polymerase sigma factor	hypothetical protein	hypothetical protein	hypothetical protein	MATE efflux family protein	50S ribosomal protein L11 methyltransferase	hypothetical protein	hypothetical protein	Holliday junction resolvase (EC:3.1.22.4)	Holliday junction DNA helicase RuvA	EC:3.1.22.4)	hypothetical protein	hypothetical protein	proton transporter TolQ	biopolymer ExbD/TolR family transporter	TonB domain-containing protein	translocation protein TolB	peptidoglycan-associated lipoprotein	hypothetical protein	PP-loop family protein	ATP-dependent metalloprotease FtsH (EC:3.4.24)
1.33	-1.27	-1.16	-1.72	-1.16	1.20	1.26	1.40	-1.45	1.07	-1.10	-1.09	-1.40	-1.74	1.34	-1.07	-1.07	-1.42	1.33	х	-1.19	х	-1.41	2.20	7.25	-1.56	-1.18	-1.28	-1.28	-1.28	1.02	-1.30	1.13	1.38	0.98	1.29	-1.66	1.44
0.18	0.68	0.30	0.29	0.88	0.13	0.27	×	0.10	0.93	0.90	0.71	0.20	0.29	0.03	0.88	0.87	0.87	0.06	×	0.68	×	0.01	0.02	0.00	0.06	0.35	0.67	0.43	0.82	0.68	0.04	0.70	0.08	0.13	0.71	0.22	0.05
-1.83	-1.61	-1.31	-1.30	1.82	-1.28	1.40	1.51	-1.12	1.09	1.04	-1.07	-1.14	-1.07	1.29	-1.04	-1.05	-1.06	-1.08	×	1.09	1.30	1.46	1.60	2.56	1.03	1.11	1.19	1.39	1.12	1.09	-1.26	-1.31	-1.38	-1.81	-1.45	-1.07	-1.43
0.04	0.12	0.82	0.08	0.46	0.50	0.12	0.22	0.72	0.09	0.94	0.76	0.31	0.17	0.05	0.92	0.90	0.89	0.54	×	0.69	0.02	0.02	0.02	0.01	0.59	0.34	0.43	0.67	0.74	0.72	0.19	0.32	0.06	0.16	0.10	0.77	0.11
-0.25	-1.44	-1.24	-1.51	0.33	-0.04	1.33	1.46	-1.29	1.08	-0.03	-1.08	-1.27	-1.41	1.32	-1.06	-1.06	-1.24	0.13	N/A	-0.05	N/A	0.03	1.90	4.91	-0.27	-0.03	-0.05	0.05	-0.08	1.06	-1.28	-0.09	0.00	-0.41	-0.08	-1.37	0.01
1.58	0.17	0.08	0.21	1.49	1.24	0.07	0.06	0.16	0.01	1.07	0.01	0.13	0.34	0.03	0.02	0.01	0.18	1.21	N/A	1.14	N/A	1.44	0.30	2.35	1.30	1.15	1.24	1.34	1.20	0.04	0.02	1.22	1.38	1.40	1.37	0.30	1.44
1.65	-1.23	1.08	-3.10	1.37	2.54	4.27	3.34	-1.62	1.39	1.25	2.45	1.27	0.91	4.57	-1.75	-1.01	-1.89	1.14	Х	1.61	2.39	-1.08	-2.93	7.21	-1.56	1.46	-1.23	1.94	-1.45	2.21	1.95	0.98	1.55	-1.47	1.09	-1.85	2.13
0.03	0.76	0.02	0.03	0.77	0.01	0.02	0.00	0.11	0.72	0.54	0.02	0.23	0.74	0.01	0.30	0.24	0.78	0.39	x	0.10	0.01	0.43	0.01	0.00	0.16	0.16	0.78	0.08	0.18	0.04	0.01	0.71	0.09	0.01	0.45	0.20	0.01
-1.26	-1.38	-1.39	1.05	-1.05	-1.56	-2.34	-1.34	1.10	1.64	-0.98	1.05	-1.35	-1.11	-1.60	-1.29	1.11	1.09	-1.08	х	-1.66	-1.46	-1.41	1.47	1.48	-1.89	-1.38	-1.63	-1.54	1.11	0.99	1.00	-1.12	-1.23	-1.30	-1.14	-1.06	1.29
0.17	0.04	0.02	1.00	0.93	0.12	0.02	0.08	0.66	0.03	0.97	0.95	0.08	0.25	0.07	0.23	0.44	0.65	0.73	x	0.03	0.09	0.03	0.06	0.05	0.04	0.08	0.06	0.56	0.30	0.27	0.58	0.16	0.00	0.09	0.18	0.87	0.06
0.20	-1.31	-0.16	-1.03	0.16	0.49	0.97	1.00	-0.26	1.52	0.14	1.75	-0.04	-0.10	1.49	-1.52	0.05	-0.40	0.03	N/A	-0.02	0.47	-1.25	-0.73	4.35	-1.73	0.04	-1.43	0.20	-0.17	1.60	1.47	-0.07	0.16	-1.39	-0.02	-1.46	1.71
1.46	0.08	1.24	2.08	1.21	2.05	3.31	2.34	1.36	0.13	1.11	0.70	1.31		3.09	0.23	1.06	1.49	1.11	N/A	1.64	1.93	0.16	2.20	2.87	0.17	1.42	0.20	1.74	1.28	0.61	0.48	1.05	1.39	0.09		0.40	0.42
1.34	1.57	1.90	-1.52	1.11	-1.60	-1.06	-1.06	1.24	1.36	1.24	-1.39	1.10	1.19	-1.20	-1.36	1.42	1.34	-1.01	×	-1.34	1.10	-1.11	-1.05	-1.71	1.16	1.35	1.48	1.34	1.09	1.57	1.03	-1.28	-1.10	-1.22	1.25	1.50	1.33
0.09	0.09	0.01	2 0.32	0.69	0 0.13	6 0.56	6 0.52	0.07	5 <u>0.02</u>	1 0.79	9 0.17	) 0.65	0.28	0 0.14	6 0.07	0.06	0.05	0.88	x	4 0.06	0.52	0.24	5 0.41		5 0.14	5 0.01	3 0.16	0.60	0.47	7 0.01	3 0.74	8 0.16	0 0.07	2 0.18			3 0.04
1.26	1.27	1.35	2.05	1.38	-2.48	-1.95	-2.07	1.87	1.42	1.23	-1.35	-2.01		-1.65	-1.28	1.23	1.31	-1.61	x	-1.18	-2.02	-1.46	-1.48	-2.39	-1.00	-1.06	1.20	1.29	-1.03	1.24	-1.45	5 1.05	-1.15	-1.40		-1.32	1.94
5 0.07	0.22	5 0.06	5 0.14	» ×	8 0.00	5 0.06	7 0.03	7 0.25	0.13	3 0.09	5 0.13	1 0.02	0 0.11	5 0.07	8 0.68	3 0.62	0.23	1 0.02	x	8 0.74	2 0.15	6 0.20	8 0.10	9 0.14	0 0.48	6 0.60	) 0.51	0.75	3 0.84	0.14	5 0.04	5 0.77	5 0.04	0 0.00		2 0.33	4 0.07
7 1.30	2 1.42	6 1.63	4 0.27	1.25	0 -2.04	6 -1.51	-1.57	5 1.56	3 1.39	9 1.24			_	7 -1.43	8 -1.32	2 1.33	3 1.33	2 -1.31	N/A	4 -1.26	5 -0.46	0 -1.29	0 -1.27		8 0.08	_	1 1.34	5 1.32	4 0.03	4 1.41	4 -0.21		4 -1.13	0 -1.31	4 1.12	3 0.09	7 1.64
0 0.04	2 0.15	3 0.28	7 1.79	5 0.13	0.44	0.45	0.51	6 0.31	9 0.03	4 0.01	0.02	-6 1.56	1.25	-3 0.23	0.04	3 0.10	3 0.02	0.30	A N/A	0.08	-6 1.56	0.17	0.21	0.34	8 1.08	5 1.21	4 0.14	2 0.03	3 1.06	1 0.17	1 1.24	2 1.17	3 0.02	0.09	2 0.13	9 1.41	4 0.31
4	S	s	ç	3	4	S		_	3		2	5	51	3	4	5	2	J	-	æ	5	7	_	4	×	_	-+	3	J	7	4	7	2	ę			

#### SPO3178 SPO3176 SPO3172 SPO3170 SPO3169 SPO3162 SPO3161 SPO3160 SPO3157 SPO3151 SPO3148 SPO3147 SPO3174 SPO3173 SPO3171 SPO3167 SPO3165 SPO3154 SPO3152 SPO3149 SPO3143 SPO3155 SPO3145 SPO3144 SPO3180 SPO3179 SPO3177 SPO3166 SPO3164 SPO3163 SPO3159 SPO3158 SPO3156 SPO3150 SPO3146 SPO3175 SPO3168 SPO3153 ppaChisC atpA atpG atpDatpC truA rpiB clpAatpH prsnahD ribFClpA cyclohexadienyl dehydrogenase (EC:1.3.1.12) ATP-dependent Clp protease ATP-binding subunit riboflavin biosynthesis protein RibF (EC:2.7.1.26 2.7.7.2) manganese-dependent inorganic pyrophosphatase (EC:3.6.1.1) ribose 5-phosphate isomerase B (EC:5.3.1.6) hypothetical protein histidinol-phosphate aminotransferase (EC:2.6.1.9) hypothetical protein hypothetical protein cation transport protein ChaC hypothetical protein hypothetical protein M24/M37 family peptidase hydroxyacylglutathione hydrolase hypothetical protein hypothetical protein ATP synthase subunit delta (EC:3.6.3.14) ATP synthase F0F1 subunit alpha (EC:3.6.3.14) ATP synthase F0F1 subunit gamma (EC:3.6.3.14) ATP synthase F0F1 subunit beta (EC:3.6.3.14) ATP synthase F0F1 subunit epsilon (EC:3.6.3.14) hypothetical protein ribose-phosphate pyrophosphokinase (EC:2.7.6.1) 2-hydroxychromene-2-carboxylate isomerase alpha/beta hydrolase hypothetical protein acyl dehydratase MaoC AraC family transcriptional regulator hypothetical protein hypothetical protein GumN family protein GumN family protein hypothetical protein tRNA pseudouridine synthase A (EC:5.4.99.12) AsnC family transcriptional regulator L-threonine aldolase monooxygenase HAD family hydrolase 0.99 0.96 2.17 1.25 1.02 1.58 1.78 2.01 2.05 1.15 1.83 1.80 1.72 2.28 1.16 -0.96 2.62 1.88 1.48 1.13 1.61 1.29 .33 1.25 3.74 .15 0.57 0.93 0.62 0.40 0.11 0.39 0.01 0.76 0.39 0.37 0.56 0.01 0.660.04 0.100.01 0.28 0.07 0.20 0.07 0.25 0.230.80 X 0.25 0.16 0.010.01 0.02 0.02 0.240.88 0.08 0.00 1.00 1.00 0.00 0.01 2.15 0.990.96 1.021.12 -1.24 -1.320.99 1.09 2.121.141.58 1.46 1.37 1.00 1.25 1.39 1.05 1.30 1.04 1.37 1.64 1.1 1.23 1.22 1.27 1.23 1.23 1.01 1.36 1.100.18 0.01 0.69 0.990.00 0.680.41 0.73 0.40 0.61 0.01 0.140.77 0.61 0.70 0.68 0.74 0.02 0.100.22 0.44 0.30 0.52 0.05 0.18 0.05 0.40 0.21 0.97 0.74 0.37 0.01 0.60 0.86 0.810.59 0.02 1.00 0.04 1.75 0.12 0.09 2.39 0.42 -0.13-1.2 -0.19 2.09 1.48 -0.03 -0.2 -1.25 0.28 1.31 1.38 1.57 1.55 1.74 1.46 1.07 1.42 1.40 1.43 1.06 1.13 1.26 1.26 1.09 1.26 .40 òœ 0.17 0.05 0.27 0.40 0.11 0.24 0.53 0.081.14 1.14 0.04 0.17 0.60 0.46 0.04 0.100.35 0.03 0.42 2.49 0.08 0.04 0.24 1.31 0.82 0.06 0.07 0.09 1.38 1.05 0.07 1.15 1.15 1.22 1.331.05 1.47 1.21 5.60 6.44 6.51 9.45 2.930.98 2.302.99 -1.22 -1.43 1.211.37 1.145.33 1.30 1.74 2.81 3.36 2.89 1.41 3.87 1.50 7.22 1.21 3.83 1.4 1.94 0.53 0.01 0.32 0.87 0.27 0.09 0.15 0.01 0.20 0.08 0.96 0.10 0.01 0.05 0.01 0.00 0.02 0.060.01 0.54 0.20 0.00 0.02 0.03 0.00 0.00 0.01 0.00 0.00 0.230.05 0.02 0.00 0.00 0.01 1.00 0.00 0.04 -1.25 -1.35 -1.35 Ė -1.37 1.04 1.41 1.90 -1.26 1.12 1.20 1.22 1.19 1.00 1.13 1.38 1.00 1.6] 1.29 0.04 0.06 0.160.07 0.25 0.19 0.54 0.06 0.18 0.20 0.35 0.01 0.13 0.03 0.07 0.06 0.06 0.14 0.62 0.66 0.410.200.15 0.20 0.64 0.17 0.01 0.86 0.98 0.69 0.05 0.12 0.090.46 0.15 0.50 0.87 0.63 0.69 -1.73 -1.29 -1.39 0.050.03 -0.35 -0.20 -0.0 -1.74 -0.02 2.04 2.48 0.01 0.31 0.85 0.15 0.66 -0.10 -1.19 0.213.00 0.52 -0.32 1.99 2.58 3.88 2.28 1.17 1.90 1.24 1.7 1.27 0.04 0.34 0.54 0.19 2.300.17 0.35 0.07 1.16 1.35 0.03 1.923.62 3.30 3.97 3.94 5.57 0.91 2.04 2.63 2.07 0.73 2.27 1.30 4.22 2.56 1.21 1.39 1.42 1.30 1.09 1.26 1.07 1.78 1.23 1.45 1.42 .1 4 1.331.50 1.15 1.44 1.04 1.61 1.42 1.06 1.15 1.60 1.06 4 1.07 1.16 1.06 .4 0.17 0.78 0.74 0.10 0.230.02 0.900.390.37 0.29 0.48 0.140.10 0.30 0.94 0.08 0.07 0.92 0.80 0.06 0.25 0.20 0.55 0.01 0.86 0.0] 0.02 0.04 0.35 0.03 0.01 0.06 0.23 0.65 0.06 0.32 0.03 0.05 -1.4 2.45 1.40 2.10 0.98 2.16 1.85 1.84 1.53 1.01 1.82 1.681.22 1.91 1.10 1.341.43 5 in 0.00 0.090.05 0.05 0.02 0.59 0.18 0.03 0.30 0.05 0.26 0.06 0.07 0.10 0.06 0.05 0.77 0.28 0.00 0.80 0.30 0.010.010.05 0.03 0.80 0.83 0.51 0.12 0.02 0.08 0.14 0.06 0.84 0.01 0.01 0.02 0.01 0.160.07 0.02 0.28 0.05 0.33 0.09 1.48 1.45 1.63 1.65 1.61 1.89 1.16 1.71 1.23 .45 .29 .80 .25 0.48 0.05 0.19 0.56 0.01 1.18 1.18 1.41 0.39 0.21 0.24 1.24 0.360.44 1.18 0.12 0.14 0.13 0.13 0.15 0.100.72 0.42 0.20 0.23 0.07 0.100.110.47 1.08 0.05 0.56 1.46 1.091.58 1.25 1.34

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SPO3217	SPO3216	SPO3215	SPO3214	SPO3213	SPO3212	SPO3211	SPO3210	SPO3209	SPO3208	SPO3207	SPO3206	SPO3205	SPO3204	SPO3203	SPO3202	SPO3201	SPO3200	SPO3199	SPO3198	SPO3197	SPO3196	SPO3195	SPO3194	SPO3193	SPO3192	SPO3191	SPO3190	SPO3189	SPO3188	SPO3187	SPO3186	SPO3185	SPO3184	SPO3183	SPO3182	SPO3181
def-1	fmt				rnhA					ispH		folK	rpoZ			pdxJ	acpS	lepB	rnc		recO									comC	opuD					rpsD
peptide deformylase (EC:3.5.1.88)	methionyl-tRNA formyltransferase (EC:2.1.2.9)	hypothetical protein	hypothetical protein	hypothetical protein	ribonuclease H	hypothetical protein	hypothetical protein	amino acid transporter LysE	glutathione S-transferase	4-nyuroxymeutyiour-z-enyi uipnospnate reductase (EC:1.17.1.2)	hypothetical protein	2-ammo-4-hydroxy-o- hydroxymethyldihydropteridine pyrophosphokinase (EC:2.7.6.3)	EC:2.7.7.6)	guanosine-3,5'-bis(diphosphate) 3'- pyrophosphohydrolase porto area of pyrophosphohydrolase	hypothetical protein	pyridoxine 5'-phosphate synthase	4'-phosphopantetheinyl transferase (EC:2.7.8.7)	signal peptidase I (EC:3.4.21.89)	ribonuclease III (EC:3.1.26.3)	hypothetical protein	DNA repair protein RecO	histone deacetylase	ribonucleoside-diphosphate reductase	hypothetical protein	LysR family transcriptional regulator	aldehyde dehydrogenase	renal dipeptidase	hypothetical protein	hypothetical protein	(2R)-3-sulfolactate dehydrogenase	glycine betaine transporter	GntR family transcriptional regulator	acetyltransferase	molybdopterin-binding oxidoreductase	hypothetical protein	30S ribosomal protein S4
1.47	1.02	1.27	1.39	-1.54	-1.63	-1.32	1.01	-1.05	-1.31	-1.42	-1.47	-1.41	1.27	1.09	-1.21	1.54	1.64	2.43	1.52	1.00	1.07	1.40	1.78	-1.09	-3.99	-2.06	-2.17	-1.69	1.25	-1.75	-1.47	-1.85	2.16	-1.10	-2.11	-2.09
0.09	0.90	0.86	0.15	0.01	0.03	0.76	0.97	0.16	0.09	0.36	0.49	0.12	0.10	0.71	0.11	0.02	0.01	0.05	0.21	0.61	0.79	0.50	0.32	0.26	0.01	0.17	0.02	0.57	0.94	0.23	0.10	0.04	0.08	0.33	0.00	0.04
-1.09	0.99	-1.05	1.26	-1.40	1.41	1.23	1.07	1.03	-1.13	1.01	0.99	-1.17	-1.47	1.11	1.13	1.14	1.02	1.04	1.24	1.63	-1.11	1.32	1.32	1.08	-1.54	-1.41	-1.54	-1.25	-1.27	-1.53	-1.44	-1.10	1.04	1.07	-1.09	1.46
0.48	0.95	0.69	0.25	0.21	0.05	0.33	0.85	0.95	0.04	0.89	0.90	0.45	0.04	0.66	0.42	0.36	0.75	0.92	0.33	0.13	0.28	0.16	0.14	0.69	0.07	0.05	0.06	0.19	0.55	0.19	0.15	0.24	0.90	0.39	0.61	0.07
0.19	1.00	0.11	1.33	-1.47	-0.11	-0.05	1.04	-0.01	-1.22	-0.21	-0.24	-1.29	-0.10	1.10	-0.04	1.34	1.33	1.74	1.38	1.31	-0.02	1.36	1.55	-0.01	-2.77	-1.74	-1.86	-1.47	-0.01	-1.64	-1.46	-1.48	1.60	-0.02	-1.60	-0.32
1.28	0.02	1.16	0.06	0.07	1.52	1.28	0.03	1.04	0.09	1.22	1.23	0.12	1.37	0.01	1.17	0.20	0.31	0.70	0.14	0.32	1.09	0.04	0.23	1.09	1.23	0.33	0.32	0.22	1.26	0.11	0.02	0.38	0.56	1.09	0.51	1.78
2.66	1.19	1.63	1.86	-2.05	-1.05	1.61	1.41	1.10	2.21	1.72	2.26	1.13	4.17	2.38	1.91	2.62	2.64	5.31	4.14	1.06	-1.06	-1.13	3.11	1.08	-1.21	-1.55	-1.50	-1.66	1.57	-1.48	-2.56	-1.97	-1.00	-2.30	-1.42	2.37
0.01	0.20	0.13	0.07	0.02	0.84	0.14	0.17	0.10	0.02	0.22	0.06	0.25	0.00	0.01	0.02	0.03	0.00	0.03	0.00	0.93	0.76	0.81	0.09	0.86	0.84	0.44	0.15	0.70	0.81		0.02		0.56	_		0.01
-1.08	-1.22	1.13	-1.03	1.09	-1.45	1.07	1.01	1.10	-1.59	-1.19	0.98	-1.12	1.37	1.29	1.13	1.35	1.21	-1.06	-1.28	-1.33	1.36	1.31	-1.18	2.12	1.18	1.08	1.54	1.42	1.18	1.51	1.47	-1.04	1.18	-1.49	-2.04	-1.52
0.26	0.44	0.38	0.71	0.67	0.09	0.75	0.72	0.75	0.02	0.23	0.70	0.10	0.30	0.07	0.57	0.00	0.25	0.60	0.01	0.55	0.10	0.09	0.55	0.01	0.10	0.63	0.07	0.05	0.21	0.37	0.03	0.78	0.55	0.07	0.02	0.08
0.79	-0.02	1.38	0.42	-0.48	-1.25	1.34	1.21	1.10	0.31	0.27	1.62	0.00	2.77	1.84	1.52	1.99	1.93	2.13	1.43	-0.14	0.15	0.09	0.97	1.60	-0.02	-0.24	0.02	-0.12	1.38	0.02	-0.55	-1.51	0.09	-1.90	-1.73	0.43
1.87	1.21	0.25	1.45	1.57	0.20	0.27	0.20	0.00	1.90	1.46	0.64	1.13	1.40	0.55	0.39	0.64	0.72	3.19	2.71	1.20	1.21	1.22	2.15	0.52	1.20	1.32	1.52	1.54	0.20	1.50	2.02	0.47	1.09	0.41	0.31	1.95
1.18		1.02	1.40	1.03		-1.12	1.58	1.03	-1.05	1.31	1.44	-1.57	-1.36	0.99		1.03		-	1.26	1.35		1.11		1.57							1.51		-			1.06
8 0.21		2 0.93	0 0.03		3 0.61	2 0.42	8 0.05	3 0.93	5 0.54	1 0.28	4 0.31	7 0.03	6 0.17	9 0.34	6 0.04	3 0.70	7 0.05	3 0.67	6 0.05	5 0.41	4 0.76	1 0.68	9 0.55	7 0.18	0.75	6 0.77	7 0.64	8 0.74	3 0.94	0 0.96	1 0.02		5 0.56			6 0.83
			3 1.05	4 2.61		2 -1.10	5 1.38				1 1.46		7 0.89	4 0.98	i.		5 1.17		5 -1.38	1 -1.06				8 2.86		7 -0.98	-0.92		4 1.54		2 1.79		6 -0.93			3 -1.54
1.16 0.		-1.27 0.			1.43 0.			1.28 0.	1.52 0.	1.25 0.		1.55 0.			1.36 0.	1.44 0.					1.84 0.				-1.24 0.			1.23 0.								
0.14 0.	-	0.12 -0	0.87 1.	0.06 1.	0.10 -1	0.70 -1	0.44 1.	0.37 1.	0.03 -1	0.33 1.	0.37 1.		0.23 -0	0.29 0.	0.13 -1	0.04 1.	0.77 -0	0.37 -1	0.04 -0	0.95 0.	0.01 1.		0.61 0.	0.04 2.	0.41 -1	0.72 -1	0.42 0.	0.37 0.	0.33 1.	0.41 -0	0.06 1.	-	0.56 -1			0.05 -0
0.01		-0.13	1.23 (	1.82 (	-1.23 (	-1.11 (	1.48 (	1.16 (	-1.29 (	1.28 (	1.45 (	-1.56 (	-0.24	0.98 (	-1.41	1.24 (	-0.10		-0.06	0.15	1.44 (	1.10 (	0.07	2.22 (		-1.02 (	0.07	0.13	1.29 (		1.65 (		-1.04 (			-0.24
1.17	0.12	1.15	0.17	0.79	0.20	0.01	0.10	0.13	0.23	0.03	0.01	0.01	1.12	0.01	0.05	0.21	1.27	0.06	1.32	1.21	0.40	0.02	1.22	0.65	0.09	0.04	1.00	1.10	0.25	0.03	0.14	1.37	0.11	1.35	0.27	1.30

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SPO3256 rpmE	SPO3255 rimM	SPO3254 bluB	SPO3253 rpsP	SPO3252	SPO3251	SPO3250	SPO3249	SPO3248	SPO3247	SPO3246 ffh	SPO3245 nadC	SPO3244	SPO3243 nadA	SPO3242	SPO3241	SPO3240	SPO3239	SPO3238	SPO3237 atpl	SPO3236 atpB	SPO3235 atpE	SPO3234	SPO3233 atpF	SPO3232	SPO3231 pdhR	SPO3230	SPO3229	SPO3228	SPO3227	SPO3226	SPO3225 cobD	SPO3224 cobC	SPO3223	SPO3222	SPO3221	SPO3220	SPO3219 def-3	SPO3218 def-2
50S ribosomal protein L31	16S rRNA-processing protein RimM	cobalamin biosynthesis protein BluB	30S ribosomal protein S16	chorismate mutase (EC:5.4.99.5)	hypothetical protein	acetyltransferase	acetyltransferase	acetyltransferase	acetyltransferase	signal recognition particle protein	nicotinate-nucleotide pyrophosphorylase (EC:2.4.2.19)	L-aspartate oxidase (EC:1.4.3.16)	quinolinate synthetase	prolyl-tRNA synthetase	Asp/Glu/Hydantoin racemase	LysR family transcriptional regulator	hypothetical protein	ArsR family transcriptional regulator	ATP synthase F0 subunit I (EC:3.6.3.14)	ATP synthase F0F1 subunit A (EC:3.6.3.14)	ATP synthase F0F1 subunit C (EC:3.6.3.14)	ATP synthase F0F1 subunit B' (EC:3.6.3.14)	ATP synthase F0F1 subunit B (EC:3.6.3.14)	hypothetical protein	pyruvate dehydrogenase complex repressor	class I and II aminotransferase	lipoprotein	SMC protein	hypothetical protein	hypothetical protein	cobalamin biosynthesis protein	cobalamin biosynthetic protein CobC	response regulator	hypothetical protein	hypothetical protein	class I and II aminotransferase	peptide deformylase (EC:3.5.1.88)	peptide deformylase (EC:3.5.1.88)

-1.42	1.07	-1.09	1.04	1.06	-1.30	-1.31	-1.38	-1.38	-1.06	-1.14	1.94	1.58	1.26	2.66	1.19	1.22	1.26	-1.68	1.09	2.21	3.36	2.83	2.45	-1.11	1.10	1.77	1.49	-1.85	-1.66	-1.66	1.52	-1.08	-1.82	-1.59	0.95	1.42	-1.06	1.45
0.32	0.60	0.76	0.79	1.00	0.01	х	Х	0.72	0.46	0.67	0.17	0.85	0.67	0.05	0.60	0.70	0.90	0.02	0.80	0.01	0.01	0.02	0.01	0.44	0.98	0.75	0.41	0.04	0.42	0.04	0.08	0.93	0.08	0.81	0.08	0.03	0.45	0.03
1.48	1.48	1.17	1.15	-1.19	0.99	1.17	1.28	1.13	1.59	1.17	-1.53	-1.73	-1.91	1.24	-1.06	-1.02	-1.13	-1.38	0.98	-1.04	1.90	1.19	1.32	1.23	-1.32	-1.38	×	1.39	-1.19	-1.35	1.27	1.23	1.51	-1.53	1.21	-1.17	-1.42	0.99
0.01	0.44	0.38	0.06	0.35	0.09	0.75	0.06	0.70	0.01	0.08	0.01	0.21	0.07	0.24	0.36	0.95	0.85	0.08	0.65	0.83	0.00	0.29	0.25	0.47	0.42	0.66	×	0.20	0.73	0.20	0.04	0.50	0.01	0.36	0.04	0.02	0.08	0.92
0.03	1.28	0.04	1.10	-0.06	-0.15	-0.07	-0.05	-0.13	0.27	0.02	0.21	-0.08	-0.33	1.95	0.06	0.10	0.07	-1.53	1.03	0.59	2.63	2.01	1.89	0.06	-0.11	0.20	N/A	-0.23	-1.43	-1.51	1.40	0.08	-0.16	-1.56	1.08	0.13	-1.24	1.22
1.45	0.21	1.13	0.05	1.13	1.15	1.24	1.33	1.26	1.33	1.16	1.74	1.66	1.59	0.71	1.13	1.12	1.20	0.15	0.06	1.63	0.73	0.82	0.57	1.17	1.21	1.58	N/A	1.62	0.24	0.16	0.13	1.16	1.67	0.03	0.13	1.30	0.18	0.23
5.13	2.45	1.94	2.32	1.78	-1.59	1.23	2.38	2.26	2.87	1.98	6.42	2.45	3.65	4.26	1.00	2.10	2.31	1.14	7.24	9.39	11.40	13.60	11.70	1.34	1.66	1.20	-2.43	-1.26	-1.88	2.15	1.69	1.78	-2.16	-1.50	-2.70	2.76	1.90	3.36
0.00	0.08	0.01	0.03	0.03	0.03	0.84	0.00	0.35	0.00	0.01	0.04	0.64	0.08	0.03	0.52	0.14	0.63	0.03	0.02	0.00	0.00	0.00	0.00	0.33	0.44	0.91	0.07	0.44	0.08	0.03	0.00	0.67	0.02	0.84	0.00	0.01	0.02	0.00
-1.14	-1.51	-2.46	-3.18	-2.83	-2.45	-2.67	-2.62	-2.60	-2.00	-2.94	-1.12	1.16	-1.39	-1.37	1.32	-1.19	1.04	1.18	-1.52	-1.50	-1.29	-1.24	-1.64	1.61	1.58	-1.27	х	1.37	1.27	-1.76	-1.06	-1.21	1.72	1.07	1.25	1.60	-1.58	-1.18
0.29	0.49	0.07	0.01	0.01	0.01	0.17	0.00	0.10	0.01	0.01	0.34	0.78	0.06	0.04	0.04	0.59	0.96	0.42	0.01	0.04	0.11	0.09	0.02	0.10	0.27	0.51	×	0.08	0.81	0.03	0.63	0.30	0.04	0.83	0.17	0.04	0.02	0.24
2.00	0.47	-0.26	-0.43	-0.53	-2.02	-0.72	-0.12	-0.17	0.44	-0.48	2.65	1.81	1.13	1.45	1.16	0.46	1.68	1.16	2.86	3.95	5.06	6.18	5.03	1.48	1.62	-0.04	N/A	0.06	-0.31	0.20	0.32	0.29	-0.22	-0.22	-0.73	2.18	0.16	1.09
3.14	1.98	2.20	2.75	2.31	0.43	1.95	2.50	2.43	2.44	2.46	3.77	0.65	2.52	2.82	0.16	1.65	0.64	0.02	4.38	5.45	6.35	7.42	6.67	0.14	0.04	1.24	N/A	1.32	1.58	1.96	1.38	1.50	1.94	1.29	1.98	0.58	1.74	2.27
-1.16	1.10	-1.05	1.09	-2.59	-1.81	-1.49	-1.32	-1.65	-1.11	-1.19	0.99	-1.52	-1.16	-1.00	1.02	1.60	1.45	-1.40	-1.64	-1.52	1.42	1.08	1.65	1.42	1.03	-1.30	x	1.78	1.09	1.07	-1.31	-1.18	1.07	-1.52	1.72	0.98	-1.18	-1.02
0.14	0.88	0.68	0.60	0.00	0.04	0.50	0.03	0.26	0.50	0.23	0.78	0.12	0.44	0.91	0.97	0.28	0.41	0.22	0.01	0.00	0.18	0.90	0.04	0.10	0.99	0.55	×	0.03	0.71	0.61	0.08	0.40	0.33	0.21	0.00	0.55	0.17	0.45
-1.81	-1.78	-1.33	-1.45	-3.69	-1.64	-2.10	-2.07	-1.77	-2.02	-1.15	0.93	-1.26	-1.20	-1.77	1.14	1.13	1.30	-1.05	-1.66	-1.35	0.92	-1.37	0.94	1.21	1.45	-1.18	x	1.37	2.25	-1.16	-1.58	-1.54	1.11	-1.58	1.88	1.59	-1.52	-1.39
0.01	0.42	0.60	0.01	0.01	0.05	0.23	0.01	0.04	0.01	0.43	0.14	0.25	0.07	0.05	0.11	0.72	0.63	0.81	0.10	0.14	0.21	0.12	0.26	0.50	0.56	0.87	×	0.09	0.10	0.23	0.16	0.25	0.09	0.09	0.01	0.02	0.04	0.04
-1.49	-0.34	-1.19	-0.18	-3.14	-1.73	-1.80	-1.70	-1.71	-1.57	-1.17	0.96	-1.39	-1.18	-1.38	1.08	1.37	1.38	-1.23	-1.65	-1.44	1.17	-0.15	1.30	1.32	1.24	-1.24	N/A	1.58	1.67	-0.04	-1.45	-1.36	1.09	-1.55	1.80	1.28	-1.35	-1.21
0.33	1.44	0.14	1.27	0.55	0.09	0.31	0.38	0.06	0.46	0.02	0.03	0.13	0.02	0.39	0.06	0.24	0.08	0.17	0.01	0.09	0.25	1.23	0.35	0.11	0.21	0.06	N/A	0.20	0.58	1.12	0.14	0.18	0.02	0.03	0.08	0.31	0.17	0.18

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SPO3299	SPO3298	SPO3297	SPO3296	SPO3295	SPO3294	SPO3293	SPO3292	SPO3291	SPO3290	SPO3289	SPO3288	SPO3287	SPO3286	SPO3285	SPO3284	SPO3282	SPO3281	SPO3280	SPO3279	SPO3278	SPO3277	SPO3276	SPO3275	SPO3274	SPO3273	SPO3272	SPO3271	SPO3270	SPO3263	SPO3262	SPO3261	SPO3260	SPO3259	SPO3258	SPO3257
													atsA		rpmJ					pyrF		clpB											trmD		rplS
TetR family transcriptional regulator	DNA-binding response regulator	sensory box histidine kinase/response regulator	AMP-binding protein	branched-chain amino acid ABC transporter ATP- binding protein	permease	hypothetical protein branched-chain amino acid ABC transporter	permease	substrate-binding protein branched-chain amino acid ABC transporter	binding protein branched chain amino acid ABC transnorter	phenylacetate-CoA ligase branched-chain amino acid ABC transporter ATP-	ferric iron ABC transporter permease	protein	arylsulfatase (EC:3.1.6.1)	hypothetical protein	50S ribosomal protein L36	N-formylglutamate amidohydrolase	DinB family protein	DNA polymerase IV (EC:2.7.7.7)	NUDIX domain-containing protein	orotidine 5'-phosphate decarboxylase (EC:4.1.1.23)	hypothetical protein	A IP-dependent Clp protease, A IP-binding subunit ClpB	hypothetical protein	hypothetical protein	hypothetical protein	glutamate synthase	sulfite oxidase subunit YedY	sulfite oxidase subunit YedZ	hypothetical protein	MarR family transcriptional regulator	glutathione S-transferase	methyltransferase-like protein	tRNA (guanine-N(1)-)-methyltransferase	glycosyl hydrolase	50S ribosomal protein L19
Х	4.80	2.53	-1.94	-1.48	-2.06	-1.48	-1.24	-1.90	-1.10	1.07	-1.30	1.41	3.36	1.84	-1.30	-1.45	1.97	1.02	1.18	1.43	1.52	1.04	-1.98	2.37	-1.75	-1.08	1.26	х	-1.16	-1.41	-1.20	-1.39	-1.08	1.09	-1.42
х	0.00	0.00	0.55	0.07	0.00	0.63	0.72	0.01	0.21	0.91	0.09	0.08	0.15	0.07	0.04	0.04	0.00	0.78	0.57	0.28	0.13	0.40	0.01	0.00	0.09	0.09	0.40	×	0.76	0.32	0.05	0.61	0.48	0.83	0.04
1.21	1.39	1.43	1.03	1.16	1.13	1.32	1.07	-1.41	-1.06	-1.09	1.09	1.96	1.23	1.50	1.41	1.06	1.10	1.43	-1.11	-1.12	-1.11	-2.24	-1.00	-1.31	-1.03	1.04	1.45	1.52	-1.23	-1.32	-1.45	1.07	-1.30	-1.63	-1.42
0.09	0.50	0.19	0.94	0.58	0.77	0.01	0.86	0.00	0.83	0.52	0.82	0.32	0.20	0.28	0.15	0.57	0.63	0.12	0.61	0.27	0.55	0.03	0.85	0.46	0.85	0.92	0.13	0.03	0.75	0.51	0.25	0.44	0.09	0.01	0.18
N/A	3.10	1.98	-0.46	-0.16	-0.47	-0.08	-0.09	-1.66	-1.08	-0.01	-0.11	1.69	2.30	1.67	0.05	-0.20	1.54	1.23	0.03	0.16	0.21	-0.60	-1.49	0.53	-1.39	-0.02	1.36	N/A	-1.20	-1.37	-1.33	-0.16	-1.19	-0.27	-1.42
N/A	1.71	0.55	1.49	1.32	1.60	1.40	1.16	0.25	0.02	1.08	1.20	0.27	1.07	0.17	1.36	1.26	0.44	0.20	1.15	1.28	1.32	1.64	0.49	1.84	0.36	1.06	0.10	N/A	0.04	0.04	0.13	1.23	0.11	1.36	0.00
-0.89	2.90	1.65	-4.03	-3.43	-4.14	-2.96	-1.46	-6.52	-2.91	-1.59	-1.06	2.26	-1.69	-1.87	8.20	-1.38	2.17	1.53	1.92	4.24	1.25	1.38	3.12	-2.61	-1.96	-1.21	1.07	1.27	-1.03	-2.75	-1.48	-1.36	2.35	4.76	5.94
0.14	0.03	0.03	0.29	0.03	0.00	0.03	0.67	0.00	0.06	0.23	0.79	0.05	0.44	0.09	0.00	0.10	0.01	0.16	0.04	0.01	0.27	0.16	0.33	0.02	0.24	0.15	0.76	×	0.50	0.03	0.04	0.78	0.00	0.05	0.00
-1.92	1.11	1.14	1.39	1.19	-1.15	1.04	1.02	1.49	1.49	1.18	1.02	1.23	1.33	-1.13	-1.37	-1.30	-0.99	-1.31	-1.18	-1.36	1.43	-1.80	1.43	-1.00	-1.20	1.61	1.29	-1.18	-1.29	-1.09	-2.76	-1.13	-1.11	-1.21	-1.33
0.07	0.80	0.12	0.46	0.69	0.52	0.56	0.98	0.06	0.19	0.51	0.97	0.49	0.28	0.68	0.13	0.22	0.95	0.21	0.37	0.18	0.12	0.02	0.07	0.92	0.20	0.01	0.26	×	0.77	0.30	0.00	0.56	0.17	0.27	0.06
-1.40	2.01	1.40	-1.32	-1.12	-2.65	-0.96	-0.22	-2.52	-0.71	-0.21	-0.02	1.75	-0.18	-1.50	3.42	-1.34	0.59	0.11	0.37	1.44	1.34	-0.21	2.28	-1.81	-1.58	0.20	1.18	0.05	-1.16	-1.92	-2.12	-1.25	0.62	1.78	2.31
0.52	0.90	0.26	2.71	2.31	1.50	2.00	1.24	4.01	2.20	1.39	1.04	0.52	1.51	0.37	4.79	0.04	1.58	1.42	1.55	2.80	0.09	1.59	0.85	0.81	0.38	1.41	0.11	1.23	0.13	0.83	0.64	0.12	1.73	2.99	3.64
1.53	1.11	1.17	1.11	-1.16	-1.12	-1.26	-1.03	1.25	1.28	-1.16	-1.43	-1.54	1.13	1.04	-1.19	2.09	-1.82	1.52	1.21	-1.31	-1.30	1.55	1.49	-1.15	1.11	1.66	1.25	1.65	1.07	-1.24	-1.12	1.09	-1.06	1.34	-1.52
0.02	0.54	0.48	0.76	0.77	0.09	0.21	0.90	0.24	0.43	0.33	0.17	0.20	0.32	0.83	0.30	0.01	0.02	0.02	0.13	0.02	0.04	0.09	0.02	0.57	0.52	0.01	0.30	0.26	0.83	0.33	0.21	0.70	0.34	0.07	0.04
-1.36	1.38	-1.10	1.19	1.13	1.22	-1.20	1.11	2.60	1.31	1.08	-1.03	-1.78	1.30	-1.27	-2.50	1.30	-1.86	-1.31	-0.99	-1.54	-1.56	1.34	-1.08	-1.56	1.11	1.71	-1.27	x	-0.94	-1.20	-1.44	-1.22	1.11	1.30	-2.30
0.18	0.52	0.52	0.64	0.70	0.43	0.26	0.66	0.03	0.39	0.44	0.97	х	0.38	0.32	0.01	0.06	0.03	0.07	0.75	0.11	0.03	0.09	0.79	0.07	0.62	0.14	0.57	×	0.44	0.51	0.09	0.19	0.55	0.05	0.02
0.09	1.25	0.03	1.15	-0.02	0.05	-1.23	0.04	1.93	1.30	-0.04	-1.23	-1.66	1.22	-0.12	-1.85	1.70	-1.84	0.11	0.11	-1.43	-1.43	1.45	0.21	-1.36	1.11	1.69	-0.01	N/A	0.07	-1.22	-1.28	-0.06	0.03	1.32	-1.91
1.45	0.13	1.14	0.04	1.15	1.17	0.03	1.07	0.68	0.02	1.12	0.20	0.12	0.09	1.16	0.66	0.40	0.02	1.42	1.10	0.12	0.13	0.11	1.29	0.21	0.00	0.03	1.26	N/A	1.00	0.02	0.16	1.16			0.39

SP03300       bypothetical protein         SP03301       Lysk family transcriptional regulator         SP03302       GDSL-like lipase/acythydrolase         SP03303       GDSL-like lipase/acythydrolase         SP03304       phosphoglycerate mutase         SP03305       glutathione S-transferase         SP03306       saccharopine dehydrogenase         SP03307       saccharopine dehydrogenase         SP03308       hypothetical protein         SP03309       saccharopine dehydrogenase         SP03310       AD regulatory protein         SP03311 <i>ihvD</i> AD regulatory protein         SP03312       DNA-binding protein         SP03313 <i>ihvD</i> SP03310 <i>ihvD</i> SP03311 <i>ihvD</i> SP03312       DNA-binding protein         SP03313 <i>ihvD</i> SP03314 <i>ihvD</i> SP03320       hypothetical protein         SP03321       bypothetical protein         SP03322       bypothetical protein         SP03323 <i>qubD-2</i> SP03324       globin domain-containing protein         SP03325       globin domain-containing protein         SP03326	2,3,4,5-tetrahydropyridine-2,6-carboxylate N succinyltransferase (EC:2.3.1.117)	dapD	SPO3337
ihD gabD-2 dapE	hypothetical protein		SPO3336
ihD gabD-2 dapE	glutamine ABC transporter substrate-bindi protein		SPO3335
ihD gabD-2 dapE	hypothetical protein		SPO3334
ihD gabD-2	hypothetical protein		SPO3333
ihD gabD-2	nyporneucat protem succinyI-diaminopimelate desuccinylase (EC:3.5.1.18)	dapE	SPO3332
ihr gabD-2	ribonuclease R (EC:3.1)	rnr	SPO3330
iivD	globin domain-containing protein		SPO3329
ibD	creatinase (EC:3:3:3:3) succinate-semialdehyde dehydrogenase (EC:1.2.1.16)	gabD-2	SPO3328
	hypothetical protein		SPO3326
iv	hypothetical protein		SPO3325
ivD	hypothetical protein		SPO3324
iv D	hypothetical protein		SPO3323
ibD	hypothetical protein		SPO3322
inD	hypothetical protein		SPO3321
Gui	hypothetical protein		SPO3320
<i>United</i>	EpsK domain-containing protein		SPO3319
ihib	hypothetical protein		SPO3318
ΰνĐ	transcriptional regulator		SPO3317
iwD	hypothetical protein		SPO3316
twb	LysR family transcriptional regulator		SPO3315
	dihydroxy-acid dehydratase (EC:4.2.1.9)	ilvD	SPO3314
	DNA-binding protein		SPO3313
	hypothetical protein		SPO3312
	ADA regulatory protein		SPO3311
	hypothetical protein		SPO3310
	saccharopine dehydrogenase		SPO3309
	hypothetical protein		SPO3308
	saccharopine dehydrogenase		SPO3307
	glutathione S-transferase		SPO3306
	phosphoglycerate mutase		SPO3305
	allantoate amidohydrolase		SPO3304
	GDSL-like lipase/acylhydrolase		SPO3303
	hypothetical protein		SPO3302
-	LysR family transcriptional regulator		SPO3301
	hypothetical protein		SPO3300

1.97	-1.38	Х	-2.21	-2.35	-1.81	1.11	1.58	1.61	1.10	1.73	-1.45	1.35	-1.24	1.49	1.57	1.93	1.48	-1.62	1.62	-1.23	-1.85	0.91	-2.58	-1.93	1.13	-1.49	-2.38	1.51	2.06	1.52	-1.38	-1.07	2.07	1.96	Х	-1.29	-1.54
0.34	0.02	х	0.04	0.14	0.01	1.00	х	0.02	0.95	0.74	0.02	0.59	0.09	0.41	0.01	0.01	0.52	0.52	0.05	0.05	0.57	0.86	0.51	0.07	0.23	0.02	0.03	0.47	0.02	0.02	0.06	0.90	0.52	0.03	x	0.74	0.09
1.35	1.40	1.66	-1.59	-1.72	1.27	1.03	1.17	1.66	1.04	1.45	1.34	-1.20	-1.27	-1.34	-1.07	-1.26	1.18	-1.37	-1.15	-1.12	1.40	-1.14	1.06	-1.28	1.05	-1.23	1.71	-1.07	1.05	1.18	-1.09	1.10	-1.29	1.54	-1.02	-1.34	-1.04
0.09	0.46	0.02	0.07	0.07	0.66	0.97	0.25	0.05	0.94	0.38	0.08	0.16	0.34	0.09	0.21	0.49	0.60	0.21	0.14	0.71	0.54	0.79	0.77	0.28	0.43	0.16	0.02	0.59	0.89	0.06	0.59	0.58	0.11	0.05	0.92	0.50	0.51
1.66	0.01	N/A	-1.90	-2.04	-0.27	1.07	1.38	1.64	1.07	1.59	-0.05	0.08	-1.26	0.08	0.25	0.34	1.33	-1.50	0.24	-1.18	-0.23	-0.11	-0.76	-1.61	1.09	-1.36	-0.34	0.22	1.56	1.35	-1.24	0.02	0.39	1.75	N/A	-1.32	-1.29
0.31	1.39	N/A	0.31	0.31	1.54	0.04	0.21	0.02	0.03	0.14	1.40	1.28	0.02	1.42	1.32	1.60	0.15	0.13	1.39	0.05	1.63	1.03	1.82	0.33	0.04	0.13	2.05	1.29	0.50	0.17	0.15	1.09	1.68	0.21	N/A	0.03	0.25
3.10	-1.17	1.88	-2.49	-2.88	-1.16	1.11	2.56	-6.40	-1.14	1.26	-1.21	1.57	1.36	1.11	1.16	1.28	1.11	1.77	1.15	1.57	-1.26	-2.13	-1.92	-1.52	-1.17	-1.76	1.13	1.51	2.54	2.02	1.76	2.92	1.38	-1.85	-1.46	1.89	-1.17
0.02	0.27	0.04	0.09	0.14	0.33	0.99	0.03	0.00	0.16	0.83	0.05	0.21	0.28	1.00	0.61	0.15	0.91	0.37	0.59	0.01	0.97	0.54	0.53	0.24	0.94	0.03	0.68	0.65	0.00	0.01	0.08	0.11	0.12	0.03	0.21	0.43	0.67
1.07	-1.18	-1.16	-1.74	-1.58	-1.63	1.03	-1.35	1.63	1.20	-0.99	1.03	1.39	1.03	1.09	-1.06	1.15	-1.05	-1.15	-1.28	1.05	3.91	-1.05	-1.56	-1.10	1.37	1.23	1.06	-1.10	0.99	-1.14	0.99	-1.11	-1.23	-1.15	1.15	-1.06	-1.25
0.91	0.11	0.27	0.03	0.06	0.24	0.96	0.09	0.09	0.32	0.97	0.88	0.05	0.91	0.76	0.40	0.17	0.64	0.36	0.12	0.78	0.16	0.91	0.09	0.59	0.05	0.59	0.71	0.41	0.43	0.18	0.58	0.37	0.42	0.14	0.42	0.70	0.25
2.09	-1.18	0.36	-2.12	-2.23	-1.40	1.07	0.61	-2.39	0.03	0.14	-0.09	1.48	1.20	1.10	0.05	1.22	0.03	0.31	-0.07	1.31	1.33	-1.59	-1.74	-1.31	0.10	-0.27	1.10	0.21	1.76	0.44	1.37	0.91	0.08	-1.50	-0.16	0.42	-1.21
1.02	0.01	1.52	0.37	0.65	0.23	0.04	1.96	4.02	1.17	1.13	1.12	0.09	0.17	0.01	1.11	0.07	1.08	1.46	1.22	0.26	2.59	0.54	0.18	0.21	1.27	1.50	0.03	1.31	0.78	1.58	0.39	2.02	1.31	0.35	1.31	1.48	0.04
1.02	1.23	-1.14	-1.23	-1.35	1.84	-1.01	-1.16	-1.00	1.04	1.09	1.49	1.58	1.56	1.39	1.35	-1.39	1.48	-1.12	-1.34	-1.18	-1.08	1.24	1.09	-1.49	1.27	-1.25	1.20	-1.87	-2.20	-1.45	-2.04	-1.50	1.29	1.14	-1.14	1.10	1.23
0.81	0.14	0.38	0.21	0.28	0.01	0.99	0.05	0.37	0.93	0.82	0.27	0.09	0.13	0.19	0.05	0.02	0.02	0.44	0.05	0.34	0.95	0.61	0.70	0.10	0.20	0.02	0.19	0.03	0.58	0.10	0.01	0.09	0.32	0.21	0.79	0.38	0.17
1.17	1.24	-1.46	-1.14	-1.56	1.37	1.62	-1.68	х	1.02	-1.18	1.05	2.20	1.65	1.72	1.21	-1.11	1.41	-1.17	-1.51	-1.46	1.34	2.01	0.96	2.05	-1.03	-1.38	-1.29	-1.52	-1.86	-1.22	-1.84	-1.97	1.37	-1.15	Х	1.19	-1.03
0.23	0.42	0.07	0.36	0.13	0.04	0.45	0.02	×	0.96	0.91	0.65	0.04	0.01	0.06	0.42	0.25	0.14	0.63	0.03	0.09	0.65	0.34	0.74	0.17	0.70	0.03	0.16	0.10	0.01	0.09	0.01	0.02	0.39	0.29	X	0.35	0.90
1.10	1.24	-1.30	-1.19	-1.46	1.61	0.31	-1.42	N/A	1.03	-0.04	1.27	1.89	1.61	1.56	1.28	-1.25	1.45	-1.15	-1.43	-1.32	0.13	1.63	1.02	0.28	0.12	-1.32	-0.05	-1.70	-2.03	-1.34	-1.94	-1.74	1.33	-0.01	N/A	1.15	0.10
0.08	0.01	0.16	0.05	0.11	0.24	1.32	0.26	N/A	0.01	1.14	0.22	0.31	0.04	0.17	0.07	0.14	0.04	0.02	0.09	0.14	1.21	0.38	0.07	1.77	1.15	0.06	1.25	0.18	0.17	0.12	0.10	0.24	0.04	1.15	N/A	0.04	1.13

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M. Kirkwood

# Chapter 10: Appendix

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SPO3374	SPO3373	SPO3372	SPO3371	SPO3370	SPO3369	SPO3368	SPO3367	SPO3366	SPO3365	SPO3364	SPO3363	SPO3362	SPO3361	SPO3360	SPO3359	SPO3358	SPO3357	SPO3356	SPO3355	SPO3354	SPO3353	SPO3352	SPO3351	SPO3350	SPO3349	SPO3348	SPO3347	SPO3346	SPO3345	SPO3344	SPO3343	SPO3342	SPO3341	SPO3340	SPO3339	SPO3338
purH	lspA						deoC					mutL		kbl	tdh				serA		serB-2															bioB
phosphoribosylaminoimidazolecarboxamide formyltransferase/IMP cyclohydrolase (EC:2.1.2.3	lipoprotein signal peptidase (EC:3.4.23.36)	hypothetical protein	M16 family peptidase	M16 family peptidase	short chain dehydrogenase	aldehyde dehydrogenase	deoxyribose-phosphate aldolase (EC:4.1.2.4)	zinc/manganese/iron ABC transporter substrate- binding protein	zinc/manganese/iron ABC transporter A1P-binding protein	zinc/manganese/iron ABC transporter permease	zinc/manganese/iron ABC transporter permease	DNA mismatch repair protein	RmuC domain-containing protein	2-amino-3-ketobutyrate CoA ligase (EC:2.3.1.29)	L-threonine 3-dehydrogenase (EC:1.1.1.103)	serine/threonine protein phosphatase	LysR family transcriptional regulator	hypothetical protein	EC:1.1.1.95)	D-3 phosphostering dehydrogenese	phosphoserine phosphatase (EC:3.1.3.3)	ArsR family transcriptional regulator	Aha1 domain-containing protein	hypothetical protein	L-asparaginase	hypothetical protein	hypothetical protein	radical SAM protein	hypothetical protein	cystathionine gamma-lyase (EC:4.4.1.1)	hypothetical protein	decarboxylase	threonine dehydratase (EC:4.3.1.19)	GntR family transcriptional regulator	bioY family protein	biotin synthase (EC:2.8.1.6)
-1.19	1.82	0.99	1.56	-1.13	1.07	0.96	-1.23	2.76	2.96	2.35	2.56	1.75	1.21	-1.23	-1.03	-1.16	-1.50	-2.59	-1.12	1.19	-1.05	-2.80	-1.52	-1.78	1.09	-1.25	-1.37	-1.08	1.53	-1.26	-1.59	1.25	-1.06	1.24	1.07	-1.05
0.17	0.03	0.68	0.17	0.01	0.79	0.48	0.01	0.01	0.11	0.32	0.00	0.17	0.09	0.57	0.93	0.33	0.09	0.14	0.67	1.00	0.62	0.47	0.04	0.16	0.10	0.15	0.44	0.46	0.12	0.17	0.75	0.05	0.59	Х	0.61	×
1.90	1.71	1.14	-1.46	-1.32	-1.04	-1.23	-1.55	-1.21	-1.16	1.21	1.15	-1.10	-1.38	-1.56	-1.17	1.25	-1.11	-1.00	1.50	0.97	1.69	-1.19	-1.04	1.05	-1.47	-1.13	-1.18	-1.02	1.59	1.08	-1.01	1.37	1.10	-1.17	1.10	1.01
0.00	0.07	0.34	0.23	0.04	0.92	0.08	0.02	0.06	0.34	0.56	0.48	0.17	0.05	0.03	0.27	0.12	0.33	0.97	0.20	0.29	0.15	0.57	0.28	0.92	0.01	0.30	0.64	0.28	0.08	0.61	1.00	0.22	0.63	0.09	0.73	0.98
0.36	1.77	1.06	0.05	-1.23	0.02	-0.13	-1.39	0.78	0.90	1.78	1.86	0.33	-0.09	-1.40	-1.10	0.05	-1.31	-1.79	0.19	1.08	0.32	-2.00	-1.28	-0.37	-0.19	-1.19	-1.28	-1.05	1.56	-0.09	-1.30	1.31	0.02	0.04	1.09	-0.02
1.55	0.06	0.08	1.51	0.10	1.06	1.10	0.16	1.99	2.06	0.57	0.71	1.43	1.30	0.17	0.07	1.21	0.19	0.80	1.31	0.11	1.37	0.81	0.24	1.42	1.28	0.06	0.10	0.03	0.03	1.17	0.29	0.06	1.08	1.21	0.02	1.03
1.63	3.54	1.28	3.92	2.06	-2.91	-1.28	-1.33	1.17	1.14	1.28	1.13	1.89	1.02	17.20	23.70	1.45	-2.03	-1.78	2.80	2.64	2.37	-2.36	-1.58	-1.39	2.38	2.30	1.75	2.56	-1.80	-1.61	-4.18	2.21	2.04	1.99	1.30	-1.19
0.01	0.00	0.22	0.02	0.01	0.01	0.19	0.04	0.03	0.37	0.97	0.89	0.11	0.50	0.00	0.00	0.04	0.07	0.33	0.13	0.11	0.05	0.59	0.06	0.13	0.00	0.03	0.49	0.01	0.03	0.04	0.28	0.00	0.05	0.02	0.11	0.90
-1.15	-1.28	-1.06	1.15	1.13	-1.15	-1.20	-1.08	-2.18	-1.86	-1.92	-2.63	1.15	-1.16	1.04	1.19	-1.62	-1.57	-1.13	-1.24	-1.13	-1.37	-1.42	1.39	1.14	1.04	1.43	1.22	-1.85	1.22	-1.02	-1.14	-1.36	1.29	-1.10	1.09	1.40
0.20	0.23	0.53	0.67	0.49	0.38	0.28	0.27	0.02	0.02	0.12	0.02	0.05	0.31	0.73	0.32	0.01	0.09	0.80	0.58	0.58	0.45	0.40	0.02	0.75	0.93	0.29	0.71	0.10	0.19	0.54	0.94	0.07	0.21	0.50	0.65	0.71
0.24	1.13	0.11	2.54	1.60	-2.03	-1.24	-1.21	-0.51	-0.36	-0.32	-0.75	1.52	-0.07	9.12	12.45	-0.09	-1.80	-1.46	0.78	0.76	0.50	-1.89	-0.10	-0.13	1.71	1.87	1.49	0.36	-0.29	-1.32	-2.66	0.43	1.67	0.45	1.20	0.11
1.39	2.41	1.17	1.39	0.47	0.88	0.04	0.13	1.68	1.50	1.60	1.88	0.37	1.09	8.08	11.26	1.54	0.23	0.32	2.02	1.89	1.87	0.47	1.49	1.27	0.67	0.44	0.27	2.21	1.51	0.30	1.52	1.79	0.38	1.55	0.11	1.30
1.06	-1.20	-1.21	1.13	1.29	1.29	-1.01	-1.12	-1.01	-1.06		1.31	-1.00	1.07		5 1.08	-1.37	-1.25	1.08	-1.68	-1.09	1.07	-1.18	1.05	1.41	-1.32	1.54	1.29	-1.23	1.56	-1.18	-1.09	1.09	-1.12	1.27	-1.10	-1.00
0.81		0.16		0.22	0.15	0.88	0.18		5 0.67	0.41	0.16	0.93	0.50		0.38	0.22	0.25	0.69	0.31	0.58	0.89	3 0.63	0.84	0.33	2 0.01	. 0.08		3 0.17	0.05	3 0.18	0.97	0.34	0.65	0.06	0.71	
-1.16	-2.47	-1.37	1.41	1.34					-1.74	-1.12	-1.42	1.18	1.19			-1.75	-1.21	-0.93	-1.57	1.11	1.04		1.01	1.24	-1.28		1.12			1.24	1.69		-1.13	-1.16		-1.16
6 0.10		7 0.13	0.31	4 0.08	5 0.01	8 0.17			4 0.01		2 0.19	3 0.33		5 0.31	€ 0.40	5 0.06	1 0.37	3 0.32	7 0.40	0.85	4 0.96		0.71	4 0.59	8 0.11	0.24	0.93	1 0.04	5 0.03	4 0.35	X	9 0.12	3 0.70			6 0.90
0 -0.05					1 2.07	7 -1.15	3 -1.11		-1.40		9 -0.05					6 -1.56		2 0.08		5 0.01				9 1.33		4 1.42	3 1.21	4 -1.32				2 -0.20	0 -1.13	-		0 -1.08
)5 1.11	0.64	0.08	7 0.14	2 0.03			.1 0.02		0 0.34	8 1.20	1.37	9 1.09	3 0.06		4 0.05	0.19	0.02	8 1.01	0.05	1 1.10	6 0.02	0.16	3 0.02	3 0.09	0 0.02	2 0.12	1 0.09	0.09	1 0.06	3 1.21	0 1.39	1.29	.3 0.00	6 1.22	1 1.11	0.08
	4	æ	-+	3	æ	4+	5	_	++	0	7	ę	5	-	л	¢	5	1	5	0	2	5	5	ę	2	5	Ŷ	Ŷ	5	-	ć	Ŷ	5	5	_	~

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SPO3412	SPO3411	SPO3410	SPO3409	SPO3408	SPO3407	SPO3406	SPO3405	SPO3404	SPO3403	SPO3402	SPO3401	SPO3400	SPO3399	SPO3398	SPO3397	SPO3396	SPO3395	SPO3394	SPO3393	SPO3392	SPO3391	SPO3390	SPO3389	SPO3388	SPO3387	SPO3386	SPO3385	SPO3384	SPO3383	SPO3382	SPO3381	SPO3380	SPO3379	SPO3378	SPO3377	SPO3376	SPO3375	
																	rarD-2			rluA			pnp															
hypothetical protein	gamma-glutamyltranspeptidase	anti-sigma B factor	anti-anti-sigma factor	acetyl-CoA C-acetyltransferase	hypothetical protein	TetR family transcriptional regulator	hypothetical protein	hypothetical protein	DNA-binding protein	amino acid transporter LysE	hypothetical protein	glycine cleavage system protein T	alcohol dehydrogenase (EC:1.1.1.1)	homocysteine S-methyltransferase	AraC family transcriptional regulator	FAD-dependent oxidoreductase	protein RarD	GDSL-like lipase/acylhydrolase	hypothetical protein	ribosomai large subunit pseudouridine synthase A (EC:4.2.1.70)	alpha-1,2-fucosyltransferase	hypothetical protein	polynucleotide phosphorylase (EC:2.7.7.8)	TetR family transcriptional regulator	hypothetical protein	hypothetical protein	glycosyl transferase family protein	FkbM family methyltransferase	thiol-specific antioxidant protein	aldehyde dehydrogenase	hypothetical protein	gamma-butyrobetaine hydroxylase	LysR family transcriptional regulator	benzoate transporter	hypothetical protein	ribosomal RNA small subunit methyltransferase B	hypothetical protein	3.5,4.10)
1.22	1.81	1.43	1.15	3.83	х	-1.00	1.40	х	3.95	4.96	1.35	6.07	2.31	6.35	4.67	5.46	1.23	1.20	2.19	Х	-1.57	-7.48	-1.20	1.25	1.41	1.42	-1.43	1.07	2.17	1.20	-1.31	3.40	1.04	1.28	1.56	-2.28	-1.58	
0.67	0.01	0.03	0.84	0.07	x	0.95	0.89	×	0.01	0.00	0.09	0.10	0.11	0.00	0.23	0.00	х	0.65	0.01	х	0.20	0.06	0.80	0.36	0.36	0.17	0.22	0.91	0.01	0.18	0.11	0.14	0.98	0.93	0.58	0.23	0.01	
1.38	1.27	1.86	2.43	2.75	1.27	1.22	1.35	1.17	1.44	1.47	1.40	1.65	1.15	1.57	1.28	1.89	1.41	-1.08	1.06	-1.06	1.11	-2.43	1.55	-1.03	-1.11	-1.15	1.20	1.33	1.21	1.31	1.08	1.06	-1.13	1.45	1.82	-1.27	-1.40	
0.01	0.49	0.03	0.01	0.02	0.36	0.36	0.32	0.20	0.03	0.17	0.00	0.01	0.74	0.17	0.72	0.13	0.02	0.88	0.75	0.79	0.08	0.04	0.31	0.93	0.82	0.71	0.12	0.15	0.47	0.09	0.25	0.38	0.87	0.01	0.05	0.62	0.01	
1.30	1.54	1.65	1.79	3.29	N/A	0.11	1.38	N/A	2.70	3.22	1.38	3.86	1.73	3.96	2.98	3.68	1.32	0.06	1.63	N/A	-0.23	-4.96	0.18	0.11	0.15	0.14	-0.12	1.20	1.69	1.26	-0.12	2.23	-0.04	1.37	1.69	-1.78	-1.49	
0.08	0.27	0.22	0.64	0.54	N/A	1.11	0.02	N/A	1.26	1.75	0.02	2.21	0.58	2.39	1.70	1.79	0.09	1.14	0.57	N/A	1.34	2.53	1.38	1.14	1.26	1.29	1.32	0.13	0.48	0.06	1.20	1.17	1.09	0.09	0.13	0.51	0.09	
-2.25	2.00	-4.24	-7.68	2.38	-0.96	1.14	2.35	3.96	-2.11	-1.36	-2.35	-1.06	-1.45	-1.40	-1.10	-1.12	2.47	2.56	-1.56	2.46	1.60	0.87	3.88	-1.08	-1.03	-1.33	1.42	0.94	2.94	1.74	8.50	-1.12	1.07	1.19	1.22	-2.24	-1.30	
0.02	0.00	0.00	0.01	0.18	0.10	0.13	0.56	0.01	0.02	0.89	0.00	0.97	0.32	0.58	0.99	0.72	0.00	0.32	0.02	0.00	0.07	0.20	0.25	0.80	0.75	0.73	0.04	0.56	0.01	0.00	0.01	0.18	1.00	0.97	0.34	0.03	0.15	
1.82	1.11	3.14	4.54	1.37	-1.29	-1.08	-1.14	-1.50	-1.65	-1.36	-1.48	-2.07	-1.43	-1.06	-1.38	-1.02	-1.10	1.38	1.64	-1.49	- 1.07	-1.10	-1.51	-1.33	-1.20	-1.12	-1.54	1.03	1.56	-1.29	-1.23	-1.05	-1.02	1.09	1.44	-1.10	-1.06	
0.01	0.55	0.01	0.00	0.01	0.19	0.30	0.63	0.08	0.01	X	0.08	0.04	0.31	0.81	0.47	0.85	0.53	0.58	0.09	0.15	0.40	0.20	0.15	0.36	0.49	0.24	0.57	0.93	0.02	0.18	0.34	0.78	0.96	0.18	0.15	0.86	0.44	
-0.22	1.56	-0.55	-1.57	1.88	-1.13	0.03	0.61	1.23	-1.88	-1.36	-1.92	-1.57	-1.44	-1.23	-1.24	-1.07	0.69	1.97	0.04	0.49	0.27	-0.11	1.19	-1.21	-1.12	-1.23	-0.06	0.98	2.25	0.23	3.64	-1.09	0.03	1.14	1.33	-1.67	-1.18	
2.04	0.44	3.69	6.11	0.51	0.16	1.11	1.75	2.73	0.23	0.00	0.44	0.51	0.01	0.17	0.14	0.05	1.79	0.59	1.60	1.98	1.34	0.99	2.70	0.13	0.09	0.11	1.48	0.05	0.69	1.52	4.87	0.04	1.05	0.05	0.11	0.57	0.12	
-1.38	1.34	-0.95	-1.24	1.92	-1.05	1.12	1.02	-1.02	-1.28	-1.17	-1.55	1.10	-1.62	-1.33	-1.43	1.09	1.21	1.40	1.35	-1.40	1.12	-1.25	1.20	1.14	1.21	-1.26	1.41	1.25	1.28	1.07	1.18	-1.31	1.29	1.63	1.23	-1.10	-1.20	
3 0.00	0.11	0.66	0.43	0.09	0.40	0.11	0.98	0.97	0.10	0.37	0.08	0.25	0.14	0.20	0.42	0.42	0.17	0.22	0.02	0.14	0.64	0.12	0.61	0.53	0.50	0.11	0.20	0.32	0.18	0.59	0.23	0.18	0.19	0.04	0.42	0.57	0.03	
-1.67	-0.99	x	x	1.61	-1.21	1.12	-1.07	-1.70	-1.57	-1.15	-1.37	-1.29	-1.54	-1.32	-1.05	1.12	-1.53	2.30	2.06	-1.12	-1.45	1.33	-1.22	-1.54	-1.24	-1.15	-1.43	1.59	1.70	1.27	-1.50	-1.88	1.78	-0.96	1.26	1.47	-2.07	
0.28	0.76	x	x	0.10	0.82	0.41	0.92	0.24	0.04	1.00	0.26	0.32	0.15	0.36	0.96	0.23	0.15	0.05	0.08	0.79		0.37	0.43	0.35	0.64	0.96	0.06	0.09	0.09	0.05	0.24	<sup>8</sup> 0.04	0.53	0.43	0.63	0.43	0.00	
-1.53	0.18	N/A	N/A	1.77	-1.13	1.12	-0.03	4 -1.36		-1.16	-1.46	-0.10	-1.58	-1.33	-1.24	3 1.11	-0.16	5 1.85	3 1.71	-1.26		0.04			4 -0.02	-1.21	-0.01	1.42	1.49	5 1.17	4 -0.16	4 -1.60	3 1.54	_	3 1.25	3 0.19	-1.64	
3 0.15			. N/A	0.16	0.08	0.00	3 1.05	0.34	3 0.14	0.01	0.09	1.20	0.04	3 0.01	4 0.19	0.02	5 1.37	0.45	0.36	0.14	1.29	1.29	1.21	1.34	1.23	0.06	1 1.42	0.17	0.21	0.10	5 1.34	0.29	0.25	1.29	0.02	1.29	0.44	

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SPO3450 guar	SPO3449 fatty	SPO3448 mutY A/G	SPO3447 hype	SPO3446 Dsb.	SPO3445 <i>lpxK</i> tetra	SPO3444 3-de	SPO3443 inosi	SPO3442 hype	SPO3441 hype	SPO3440 20-b	SPO3439 enoy	SPO3438 hype	SPO3437 mec	SPO3436 hype	SPO3435 glutz	SPO3434 hype	SPO3433 lipoj	SPO3432 <i>leuS</i> leuc	SPO3431 hype	SPO3430 porin	SPO3429 alani	SPO3428 hype	ribBA	SPO3426 DN/	SPO3425 exoc	SPO3424 hype	SPO3423 thior	SPO3422 ATF	SPO3421 hype	SPO3420 hype	SPO3418 ami Ubil SPO3419 bios	SPO3417 class	SPO3416 DN/	SPO3415 oute	
guanylate cyclase	fatty acid desaturase	A/G-specific adenine glycosylase (EC:3.2.2)	hypothetical protein	DsbA family thiol:disulfide interchange protein	tetraacyldisaccharide 4'-kinase (EC:2.7.1.130)	3-deoxy-D-manno-octulosonic-acid transferase	inositol monophosphatase	hypothetical protein	hypothetical protein	20-beta-hydroxysteroid dehydrogenase	enoyl-CoA hydratase	hypothetical protein	mechanosensitive ion channel protein MscS	hypothetical protein	glutathione S-transferase	hypothetical protein	lipoprotein	leucyl-tRNA synthetase (EC:6.1.1.4)	hypothetical protein	in .	alanine racemase	hypothetical protein	3,4-dihydroxy-2-butanone 4-phosphate synthase (EC:3.5.4.25)	DNA-binding response regulator	exodeoxyribonuclease III	hypothetical protein	thioredoxin	ATP-dependent protease La	hypothetical protein	hypothetical protein	amidase UbiH/UbiF/VisC/COQ6 family ubiquinone biosynthesis hydroxylase	class I and II aminotransferase	DNA translocase FtsK	outer membrane lipoprotein carrier protein LolA	

-1.10	3.65	1.18	-1.06	1.64	-1.80	-1.40	1.19	1.19	0.96	1.34	1.25	-1.06	-1.07	х	2.09	-1.24	-1.54	1.09	1.55	1.70	1.17	-1.77	-2.84	-1.25	1.63	-2.25	1.52	1.05	-1.07	1.25	1.67	1.69	-1.79	0.94	-1.22	-1.27	-1.05
0.85	0.05	0.31	0.30	0.01	0.10	0.53	0.92	0.14	0.02	0.79	0.68	0.90	0.89	×	0.42	0.09	0.33	0.29	0.02	0.66	0.79	0.44	0.00	0.08	0.01	0.13	0.06	0.82	0.87	0.10	0.12	0.05	0.03	0.69	0.00	0.02	0.96
-1.47	1.41	-1.25	1.31	-1.14	1.33	1.08	-1.04	1.25	-1.05	1.08	1.11	-1.19	1.15	1.36	1.32	1.30	1.02	1.14	-1.05	1.33	-1.11	-1.06	-1.04	-1.31	1.15	-1.14	-1.29	-1.10	1.15	-1.08	1.72	1.29	-1.18	-1.27	-1.90	-1.23	1.01
0.28	0.35	0.01	0.17	0.19	0.10	0.70	0.87	0.27	0.27	0.84	0.52	0.86	0.49	0.04	0.19	0.16	0.98	0.33	0.68	0.51	0.79	0.85	0.78	0.04	0.51	0.71	0.23	0.64	0.39	0.65	0.05	0.05	0.67	0.33	0.07	0.38	0.99
-1.29	2.53	-0.04	0.13	0.25	-0.24	-0.16	0.08	1.22	-0.05	1.21	1.18	-1.13	0.04	N/A	1.71	0.03	-0.26	1.12	0.25	1.52	0.03	-1.42	-1.94	-1.28	1.39	-1.70	0.12	-0.03	0.04	0.09	1.70	1.49	-1.49	-0.16	-1.56	-1.25	-0.02
0.18	1.12	1.22	1.19	1.39	1.57	1.24	1.12	0.03	1.01	0.13	0.07	0.06	1.11	N/A	0.38	1.27	1.28	0.02	1.30	0.18	1.14	0.36	0.90	0.03	0.24	0.56	1.41	1.08	1.11	1.17	0.03	0.20	0.31	1.11	0.34	0.02	1.03
1.62	1.17	1.37	1.00	1.45	8.27	1.34	1.70	2.86	1.98	1.52	2.05	-1.65	2.89	2.44	1.31	1.01	1.06	3.35	2.69	2.19	1.28	-1.22	-1.50	-1.75	4.22	-1.65	1.92	1.12	1.10	1.32	1.75	1.33	-1.63	0.84	1.51	-1.78	1.52
0.17	0.81	0.06	0.95	0.13	0.00	0.58	0.65	0.02	0.06	0.04	0.12	0.59	0.02	0.00	0.79	0.63	0.96	0.01	0.00	0.38	0.39	0.72	0.06	0.01	0.00	0.10	0.00	0.82	0.44	0.02	0.02	0.32	0.29	0.41	0.01	0.00	0.06
-1.16	1.26	1.02	-1.09	-1.08	1.02	-1.18	1.25	-1.18	1.45	1.18	-1.02	1.06	-1.15	-1.19	1.44	1.17	1.20	-1.47	-1.20	-1.76	-1.11	-1.28	-1.07	1.19	0.96	1.10	-1.16	-1.12	-1.05	1.24	1.06	1.24	-1.43	-1.19	1.10	-1.10	-1.25
0.25	0.56	0.88	0.07	0.50	0.85	0.31	0.18	0.15	0.12	0.17	0.78	0.94	0.50	0.34	0.36	0.35	0.70	0.03	0.27	0.09	0.64	0.27	0.66	0.38	0.19	0.85	0.20	0.15	0.69	0.30	0.89	0.13	0.41	0.41	0.50	0.29	0.05
0.23	1.22	1.20	-0.05	0.19	4.65	0.08	1.48	0.84	1.72	1.35	0.52	-0.30	0.87	0.63	1.38	1.09	1.13	0.94	0.75	0.22	0.09	-1.25	-1.29	-0.28	2.59	-0.28	0.38	0.00	0.03	1.28	1.41	1.29	-1.53	-0.18	1.31	-1.44	0.14
1.39	0.05	0.18	1.05	1.27	3.63	1.26	0.22	2.02	0.27	0.17	1.54	1.36	2.02	1.82	0.06	0.08	0.07	2.41	1.95	1.98	1.20	0.03	0.21	1.47	1.63	1.38	1.54	1.12	1.08	0.04	0.35	0.05	0.10	1.02	0.20	0.34	1.39
1.62	-1.12	-1.59	-1.10	1.32	1.13	1.32	1.08	1.30	1.24	1.03	1.18	1.12	-1.43	1.08	1.29	1.36	1.04	-1.44	1.01	1.28	-1.15	-1.00	1.19	-1.29	-1.14	1.11	-1.07	-1.07	1.20	-1.07	-1.05	1.16	1.08	1.15	1.40	1.10	-1.19
0.02	0.59	0.02	0.05	0.07	0.44	0.41	0.72	0.23	0.19	0.89	0.22	0.85	0.12	0.29	0.48	0.10	0.96	0.02	0.72	0.34	0.65	0.99	0.15	0.02	0.56	0.72	0.04	0.51	0.06	0.71	0.73	0.18	0.89	0.43	0.01	0.70	0.02
1.91	1.06	-1.85	-1.37	1.68	-1.62	-1.17	-1.15	-1.02	1.74	-1.17	-1.22	1.39	-2.12	-1.94	1.40	-1.00	-1.26	-2.09	0.96	1.22	1.18	1.08	1.06	-1.18	-1.34	1.99	0.99	-1.16	1.24	-1.13	1.48	1.21	-1.13	-1.26	1.84	1.14	-1.39
0.01	0.91	0.06	0.14	0.01	0.00	0.54	0.31	0.82	0.08	0.06	0.16	0.61	0.03	0.01	0.20	0.93	0.26	0.02	0.23	0.93	0.87	0.60	0.50	0.07	0.11	0.14	0.40	0.12	0.05	0.96	0.54	0.19	0.76	0.37	0.09	0.47	0.58
1.77	-0.03	-1.72	-1.24	1.50	-0.25	0.08	-0.03	0.14	1.49	-0.07	-0.02	1.26	-1.78	-0.43	1.35	0.18	-0.11	-1.77	0.99	1.25	0.02	0.04	1.13	-1.24	-1.24	1.55	-0.04	-1.12	1.22	-1.10	0.22	1.19	-0.02	-0.06	1.62	1.12	-1.29
0.15	1.09	0.13	0.14	0.18	1.38	1.25	1.12	1.16	0.25	1.10	1.20	0.14	0.35	1.51	0.05	1.18	1.15	0.33	0.02	0.03	1.17	1.04	0.06	0.06	0.10	0.44	1.03	0.04	0.02	0.03	1.27	0.03	1.11	1.21	0.22	0.02	0.10

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peptidoglycan binding domain-containing protein	hypothetical protein	hypothetical protein	hypothetical protein	heat shock protein 20	hypothetical protein	serine protease	hypothetical protein	glycolate oxidase, iron-sulfur subunit	glycolate oxidase subunit GlcE	glycolate oxidase subunit GlcD	hypothetical protein	hypothetical protein	polyamine ABC transporter permease	polyamine ABC transporter permease	poryannic race transporter substance ontening	polyamine ABC transporter ATP-binding protein	aminotransferase	GntR family transcriptional regulator	putrescine ABC transporter substrate-binding protein	putrescine ABC transporter ATP-binding protein	putrescine ABC transporter permease	putrescine ABC transporter permease	hypothetical protein	hypothetical protein	flagellar basal body rod modification protein	flagellar hook-length control protein	flagellar protein FlgJ	hypothetical protein	flagellin protein	flagellar biosynthesis regulatory protein FlaF	flagellar biosynthesis repressor FlbT	flagellar protein	guanylate cyclase	hypothetical protein	hypothetical protein	ribonuclease HII (EC:3.1.26.4)	modification methylase
-1.34	-1.09	1.03	1.06	-2.10	-1.08	1.11	1.20	1.89	1.27	-1.03	-1.54	1.83	1.04	1.00	-2.11	-1.41	2.18	-1.41	-2.08	1.36	1.55	1.56	-1.29	-1.22	1.46	1.45	1.79	1.41	2.16	3.71	3.02	1.40	1.40	-1.29	0.99	-1.16	1.19
0.03	0.49	0.98	0.88	0.44	0.33	0.83	x	0.03	0.46	0.93	0.21	0.27	0.69	0.61	0.00	0.15	0.19	0.33	0.17	0.57	0.64	0.48	0.20	0.37	0.44	0.04	0.67	0.20	0.21	0.05	0.05	0.32	0.77	0.11	0.05	0.17	0.94
-1.23	-1.22	-1.18	-1.07	-2.90	1.30	1.37	1.06	1.22	1.34	1.07	1.23	1.31	-1.22	-1.59	-2.44	-1.97	1.36	-1.02	-1.77	1.02	1.30	1.58	-1.14	-1.05	-1.08	1.23	1.27	1.09	2.62	1.86	1.56	1.09	-1.36	1.02	-1.07	-1.44	-1.27
0.23	0.31	0.79	0.80	0.28	0.23	0.04	0.73	0.48	0.17	0.87	0.07	0.26	0.34	0.06	0.05	0.25	0.47	0.16	0.03	0.98	0.40	0.37	0.76	0.87	0.79	0.57	0.52	0.71	0.01	0.02	0.07	0.90	0.17	0.93	0.60	0.06	0.36
-1.29	-1.16	-0.08	-0.01	-2.50	0.11	1.24	1.13	1.56	1.31	0.02	-0.16	1.57	-0.09	-0.30	-2.28	-1.69	1.77	-1.22	-1.93	1.19	1.43	1.57	-1.22	-1.14	0.19	1.34	1.53	1.25	2.39	2.79	2.29	1.25	0.02	-0.14	-0.04	-1.30	-0.04
0.06	0.06	1.11	1.07	0.40	1.19	0.13	0.07	0.34	0.04	1.05	1.39	0.26	1.13	1.30	0.17	0.28	0.41	0.20	0.16	0.17	0.13	0.01	0.08	0.09	1.27	0.11	0.26	0.16	0.23	0.93	0.73	0.15	1.38	1.16	1.03	0.14	1.23
0.95	-1.10	-1.36	-1.50	0.84	-1.72	-1.90	1.70	1.34	1.02	-1.30	1.45	-2.06	1.60	1.14	-1.49	1.27	1.22	-1.34	-3.21	-1.30	-1.20	1.44	-1.63	1.57	-1.99	-3.65	-10.80	-7.15	-2.39	-1.42	-1.70	-1.55	-1.24	-1.20	-1.58	1.25	-1.83
0.32	0.38	0.22	0.13	0.64	0.19	0.06	0.05	0.07	0.24	0.82	0.10	0.13	0.36	0.87	0.02	0.35	0.58	0.16	0.00	0.25	0.53	0.45	0.25	0.05	0.15	0.01	0.00	0.00	0.28	0.86	0.42	0.49	0.28	0.67	0.03	0.02	0.07
-1.10	-1.24	-1.37	1.20	-1.11	1.18	-1.03	-1.08	5.28	6.05	4.70	1.07	1.34	-1.15	1.00	-1.28	-1.48	-1.49	-1.27	1.59	1.23	1.45	1.27	-1.16	-1.02	1.09	2.19	2.47	2.85	4.71	4.64	2.66	-1.14	-1.06	-1.00	1.06	1.13	-1.33
0.55	х	0.14	0.03	0.79	0.15	0.87	0.64	0.00	0.01	0.10	0.52	0.10	0.23	0.70	0.03	0.04	0.34	0.03	0.08	0.56	0.10	0.62	0.24	0.89	0.87	0.05	0.08	0.04	0.00	0.01	0.01	0.85	0.42	0.93		0.28	0.19
-0.07	-1.17	-1.37	-0.15	-0.13	-0.27	-1.47	0.31	3.31	3.54	1.70	1.26	-0.36	0.23	1.07	-1.39	-0.11	-0.14	-1.31	-0.81	-0.04	0.13	1.36	-1.40	0.28	-0.45	-0.73	-4.17	-2.15	1.16	1.61	0.48	-1.35	-1.15	-1.10	-0.26	1.19	-1.58
1.03	0.07	0.01	1.35	0.98	1.45	0.44	1.39	1.97	2.52	3.00	0.19	1.70	1.38	0.07	0.11	1.38	1.36	0.04	2.40	1.27	1.33	0.09	0.23	1.30	1.54	2.92	6.64	5.00	3.55	3.03	2.18	0.21	0.09	0.10	1.32	0.06	0.25
1.28	-1.01	1.10	1.26	1.57	-1.35	-1.18	1.21	-0.98	-0.99	-0.98	1.19	1.66	1.49	1.05	1.40	-1.24	1.01	-1.10	1.00	1.02	1.05	2.05	-1.24	-1.24	-1.03	-1.66	-1.24	-1.47	-1.12	-1.25	-1.13	-1.08	1.49	-1.03	1.05	-1.87	-1.04
0.28	0.87	0.07	0.08	0.52	0.30	0.17	0.08	0.79	0.86	0.93	0.29	0.01	0.29	1.00	0.04	0.02	0.96	0.32	0.67	0.99	0.67	0.20	0.13	0.29	0.97	0.26	0.78	0.12	0.70	0.19	0.74	0.94	0.23	0.46	0.72	0.01	0.77
1.01	-1.38	2.02	1.21	2.44	-1.17	1.16	1.15	-1.31	-1.37	-0.98	-1.25	-1.01	1.87	1.45	1.49	-1.20	-1.06	-1.09	2.06	1.39	1.08	1.61	1.63	-1.51	1.23	-1.44	1.52	1.16	-1.11	Х	-0.95	1.29	1.59	-1.08	-1.15	-2.66	-1.20
0.87	×	0.10	0.10	0.45	0.82	0.68	0.51	0.62	0.26	0.79	0.49	0.49	0.04	0.23	0.21	0.08	0.79	0.41	0.02	0.34	0.52	0.28	0.14	0.17	0.52	0.26	0.50	0.38	0.91	×	0.63	0.84	0.04	0.92	0.66	0.02	0.08
1.15	-1.20	1.56	1.24	2.01	-1.26	-0.01	1.18	-1.14	-1.18	-0.98	-0.03	0.33	1.68	1.25	1.45	-1.22	-0.03	-1.10	1.53	1.21	1.07	1.83	0.20	-1.38	0.10	-1.55	0.14	-0.16	-1.12	N/A	-1.04	0.11	1.54	-1.06	-0.05	-2.27	-1.12
0.14	0.19	0.46	0.03	0.44	0.09	1.17	0.03	0.17	0.19	0.00	1.22	1.34	0.19	0.20	0.05	0.02	1.04	0.01	0.53	0.18	0.02	0.22	1.44	0.14	1.13	0.11	1.38	1.32	0.01	N/A	0.09	1.19	0.05	0.03	1.10	0.40	0.08

SPO3477 SPO3478 SPO3479 SPO3480

glcD glcE glcF

SPO3475 SPO3474

SPO3473

SPO3476

SPO3485 SPO3486 SPO3487

SPO3484 SPO3483 SPO3482 SPO3481

SPO3488

SPO3469 SPO3470 SPO3471 SPO3472

potF

SPO3467 SPO3468

potI potH potG

SPO3465 SPO3464

SPO3466

SPO3461 SPO3462 SPO3463

flgD

SPO3460 SPO3459

flgJ

SPO3458 SPO3457 SPO3456

flbT flaF

SPO3453 SPO3454 SPO3455

SPO3451 SPO3452

rnhB

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SPO3525	SPO3524	SPO3523	SPO3522	SPO3521	SPO3520	SPO3519	SPO3518	SPO3517	SPO3516	SPO3515	SPO3514	SPO3513	SPO3512	SPO3511	SPO3510	SPO3509	SPO3508	SPO3507	SPO3506	SPO3505	SPO3504	SPO3503	SPO3502	SPO3501	SPO3500	SPO3499	SPO3498	SPO3496	SPO3495	SPO3494	SPO3493	SPO3492	SPO3491	SPO3490	SPO3489
ccoO-1	ccoQ-1	ccoP-1				ccoS		secE	nusG		rplK	rplA			rplJ	rplL	rpoB	rpoC						rpsL	rpsG	fusA	tuf-2		mscL						phe T
cbb3-type cytochrome c oxidase subunit $\Pi$	cytocntome c oxtdase, cob3-type subunit i v (EC:1.9.3.1)	(EC:19.3.1) (EC:19.3.1)	family	fixH protein	copper-translocating P-type ATPase (EC:3.6.3.4)	cbb3-type cytochrome oxidase maturation protein	hypothetical protein	preprotein translocase subunit SecE	rranscription termination/antitermination factor NusG	hypothetical protein	50S ribosomal protein L11	50S ribosomal protein L1	hypothetical protein	hypothetical protein	50S ribosomal protein L10	50S ribosomal protein L7/L12	EC:2.7.7.6)	DNA-directed RNA polymerase subunit beta (EC:2.7.7.6)	DNA-binding protein	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	30S ribosomal protein S12	30S ribosomal protein S7	elongation factor G	elongation factor Tu (EC:3.6.5.3)	mechanosensitive ion channel protein MscS	ratge conductance mechanosensitive channer	glutathione S-transferase	transporter	hypothetical protein	UbiE/COQ5 family methlytransferase	LysR family transcriptional regulator	phenylalanyl-tRNA synthetase subunit beta (EC:6.1.1.20)
2.14	2.11	2.24	1.56	3.52	2.31	1.03	-1.06	-1.37	-1.30	Х	1.15	-1.58	-1.23	-1.34	-2.35	-1.83	1.47	-1.17	-1.45	1.07	-0.98	1.04	1.02	1.70	1.51	2.36	х	1.53	-1.45	1.06	1.09	-1.07	-1.44	х	1.74
0.07	0.00	0.01	0.50	0.02	0.01	0.22	0.25	0.01	0.61	Х	0.87	0.05	0.03	0.04	0.01	0.02	0.38	0.60	0.04	0.57	0.62	0.66	0.93	0.03	0.19	0.08	x	0.06	0.02	0.20	0.66	0.33	0.03	Х	0.01
1.30	1.34	2.10	1.61	1.58	1.77	1.28	1.45	-1.03	1.26	1.28	-1.08	1.90	-1.22	1.35	-1.47	2.19	-1.25	1.43	-1.06	-1.16	1.20	1.03	-1.09	-1.36	-1.67	-1.40	x	-1.94	-2.13	-2.38	1.12	-1.11	-1.33	-1.05	1.67
0.31	0.07	0.02	0.03	0.06	0.01	0.43	0.18	0.73	0.33	0.75	0.12	0.05	0.40	0.09	0.10	0.06	0.48	0.10	0.65	0.02	0.42	0.84	0.59	0.17	0.09	0.21	×	0.01	0.05	0.02	0.46	0.53	0.10	0.93	0.07
1.72	1.73	2.17	1.59	2.55	2.04	1.16	0.20	-1.20	-0.02	N/A	0.03	0.16	-1.23	0.01	-1.91	0.18	0.11	0.13	-1.26	-0.04	0.11	1.04	-0.04	0.17	-0.08	0.48	N/A	-0.21	-1.79	-0.66	1.11	-1.09	-1.39	N/A	1.71

1.30	1.34	2.10	1.61	1.58	1.77	1.28	1.45	-1.03	1.26	1.28	-1.08	1.90	-1.22	1.35	-1.47	2.19	-1.25	1.43	-1.06	-1.16	1.20	1.03	-1.09	-1.36	-1.67	-1.40	Х	-1.94	-2.13	-2.38	1.12	-1.11	-1.33	-1.05	1.67
0.31	0.07	0.02	0.03	0.06	0.01	0.43	0.18	0.73	0.33	0.75	0.12	0.05	0.40	0.09	0.10	0.06	0.48	0.10	0.65	0.02	0.42	0.84	0.59	0.17	0.09	0.21	x	0.01	0.05	0.02	0.46	0.53	0.10	0.93	0.07
1.72	1.73	2.17	1.59	2.55	2.04	1.16	0.20	-1.20	-0.02	N/A	0.03	0.16	-1.23	0.01	-1.91	0.18	0.11	0.13	-1.26	-0.04	0.11	1.04	-0.04	0.17	-0.08	0.48	N/A	-0.21	-1.79	-0.66	1.11	-1.09	-1.39	N/A	1.71
0.42	0.38	0.07	0.03	0.97	0.27	0.13	1.26	0.17	1.28	N/A	1.12	1.74	0.01	1.35	0.44	2.01	1.36	1.30	0.20	1.12	1.09	0.01	1.06	1.53	1.59	1.88	N/A	1.74	0.34	1.72	0.02	0.02	0.05	N/A	0.04
6.96	7.30	8.51	3.51	7.86	3.36	0.97	-1.23	1.54	3.97	-1.31	10.10	3.05	1.30	1.09	6.97	5.97	4.58	2.86	-1.59	1.38	2.27	1.09	1.06	7.08	6.86	9.93	Х	2.10	0.87	-1.24	-0.98	2.03	-1.32	-1.24	2.83
0.00	0.00	0.00	0.02	0.00	0.00	0.36	0.14	0.03	0.02	х	0.00	0.05	0.10	0.25	0.00	0.00	0.04	0.09	0.06	0.21	0.02	0.19	0.95	0.00	0.00	0.01	х	0.03	0.04	0.01	0.20	0.03	0.10	0.70	0.00
1.17	1.01	-1.08	1.03	-1.20	-1.13	1.31	1.52	-1.40	-1.78	-1.15	-3.66	-4.39	1.34	1.14	-4.81	-6.38	-1.15	-1.48	1.68	-1.19	-1.30	1.37	1.27	-1.08	-1.99	-1.34	Х	-1.40	-1.19	1.09	-1.02	1.48	1.12	-1.11	-1.14
0.17	0.63	0.22	0.84	0.19	0.45	0.23	0.10	0.02	0.02	0.93	0.02	0.01	0.08	0.08	0.00	0.00	0.15	0.03	0.02	0.43	0.10	0.11	0.35	0.21	0.15	0.06	x	0.01	0.07	0.63	0.95	0.03	0.34	0.49	0.38
4.07	4.16	3.72	2.27	3.33	1.12	1.14	0.15	0.07	1.10	-1.23	3.22	-0.67	1.32	1.12	1.08	-0.21	1.72	0.69	0.04	0.10	0.49	1.23	1.17	3.00	2.44	4.30	N/A	0.35	-0.16	-0.08	-1.00	1.76	-0.10	-1.18	0.85
2.90	3.15	4.80	1.24	4.53	2.25	0.17	1.38	1.47	2.88	0.08	6.88	3.72	0.02	0.02	5.89	6.18	2.87	2.17	1.64	1.29	1.79	0.14	0.11	4.08	4.43	5.64	N/A	1.75	1.03	1.17	0.02	0.28	1.22	0.06	1.99
-1.18	1.02	2.09	1.34	1.71	1.19	1.74	1.57	-1.03	-1.29	-1.20	-1.28	1.05	1.21	1.43	-1.61	1.46	-1.51	1.47	1.11	-1.00	-1.01	-1.18	-1.01	-1.60	-2.59	-1.59	Х	-1.04	1.12	1.35	1.15	1.40	-1.15	1.27	1.18
0.09	0.29	0.00	0.18	0.08	0.18	0.06	0.02	0.71	0.10	x	0.01	0.97	0.22	0.04	0.04	0.02	0.01	0.16	0.51	0.99	0.79	0.17	0.94	0.05	0.00	0.11	×	0.55	0.11	0.36	0.35	0.03	0.17	0.11	0.16
-1.14	1.04	1.37	1.85	1.71	-1.11	2.05	1.24	1.04	-1.81	1.90	-2.29	-2.45	1.51	-1.15	-4.23	-3.47	-1.68	1.25	1.22	1.17	-1.08	-1.19	-1.05	-2.57	-3.02	-1.93	Х	1.10	1.56	1.86	1.30	1.36	-1.09	1.13	-1.36
0.35	0.58	0.03	0.04	0.05	0.12	0.07	0.30	0.69	0.07	x	0.03	0.03	0.08	0.29	0.00	0.00	0.01	0.28	0.08	0.13	0.83	0.34	0.66	0.01	0.01	0.08	×	0.83	0.12	0.04	0.04	0.16	0.81	0.19	0.07
-1.16	1.03	1.73	1.60	1.71	0.04	1.90	1.41	0.01	-1.55	0.35	-1.79	-0.70	1.36	0.14	-2.92	-1.01	-1.60	1.36	1.17	0.09	-1.05	-1.19	-1.03	-2.09	-2.81	-1.76	N/A	0.03	1.34	1.61	1.23	1.38	-1.12	1.20	-0.09
0.02	0.01	0.36	0.26	0.00	1.15	0.16	0.17	1.04	0.26	1.55	0.51	1.75	0.15	1.29	1.31	2.47	0.09	0.11	0.05	1.09	0.04	0.01	0.02	0.49	0.22	0.17	N/A	1.07	0.22	0.26	0.08	0.02	0.03	0.07	1.27

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SPO3562	SPO3561	SPO3560	SPO3559	SPO3558	SPO3557	SPO3556	SPO3555	SPO3554	SPO3553	SPO3552	SPO3551	SPO3550	SPO3549	SPO3548	SPO3547	SPO3546	SPO3545	SPO3544	SPO3543	SPO3542	SPO3541	SPO3540	SPO3539	SPO3538	SPO3537	SPO3536	SPO3535	SPO3534	SPO3533	SPO3532	SPO3531	SPO3530	SPO3529	SPO3528	SPO3527	SPO3526	
tauR	XSC	pta				dapA						dnaX												cycM	yejA	yejB	yejE	yejF	fusA	hemN	fnrL		glyA-3			ccoN-1	
taurine transcriptional regulator	sulfoacetaldehyde acetyltransferase (EC:2.3.3.15)	phosphate acetyltransferase (EC:2.3.1.8)	molybdopterin-binding oxidoreductase	(Fe-S)-binding protein	dimethyl sulfoxide reductase subunit C	dihydrodipicolinate synthase (EC:4.2.1.52)	Slt family transglycosylase	hypothetical protein	hypothetical protein	oxidoreductase, FAD-binding	acetyltransferase	DNA polymerase III subunits gamma and tau (EC:2.7.7.7)	hypothetical protein	hypothetical protein	hypothetical protein	zinc metallopeptidase	TetR family transcriptional regulator	hypothetical protein	hypothetical protein	2`,3`-cyclic-nucleotide 2`-phosphodiesterase	NUDIX family hydrolase	hypothetical protein	prephenate dehydratase (EC:4.2.1.51)	cytochrome c552	binding protein	oligopeptide/dipeptide ABC transporter permease	oligopeptide/dipeptide ABC transporter permease	oligopeptide/dipeptide ABC transporter ATP- binding protein	elongation factor G	coproporphyrinogen III oxidase (EC:1)	transcriptional activator protein FnrL	LysR family transcriptional regulator	serine hydroxymethyltransferase (EC:2.1.2.1)	hypothetical protein	universal stress protein family protein	cbb3-type cytochrome c oxidase subunit I (EC:1.9.3.1)	(EC:1.9.3.1)
-1.97	1.37	-1.16	-1.44	-1.72	-1.30	-1.07	-1.20	1.30	-1.15	1.26	-1.02	-1.41	1.27	-1.58	2.49	-1.02	-1.26	-1.57	-1.40	1.31	1.93	-1.03	-1.10	2.95	1.27	-1.16	-0.94	-1.41	-1.34	1.12	2.86	-1.14	1.54	0.87	1.86	2.27	
0.00	0.88	0.66	0.08	0.01	0.66	0.83	0.16	0.09	0.40	0.04	0.97	0.01	0.06	0.75	0.12	0.95	0.79	0.28	0.33	0.20	0.00	x	0.50	0.05	0.61	0.66	0.44	0.04	0.12	0.32	0.00	0.66	0.02	0.04	0.01	0.19	
-1.34	-1.00	-1.30	1.92	1.45	1.81	1.64	-1.57	1.13	1.45	1.06	1.16	-1.02	1.23	1.13	2.21	1.07	-1.21	-1.30	-1.28	1.85	1.07	1.55	1.25	1.60	-1.08	1.06	-1.07	-1.01	1.08	1.33	3.01	-1.08	-1.31	1.09	1.42	1.62	
0.54	1.00	0.14	0.00	0.18	0.19	0.10	0.01	0.56	0.03	0.76	0.43	0.87	0.07	0.85	0.03	0.70	0.78	0.39	0.16	0.02	0.66	0.10	0.63	0.06	0.78	0.88	0.57	0.88	0.44	0.02	0.00	0.41	0.29	0.74	0.02	0.30	
-1.66	0.19	-1.23	0.24	-0.14	0.26	0.29	-1.39	1.22	0.15	1.16	0.07	-1.22	1.25	-0.23	2.35	0.03	-1.24	-1.44	-1.34	1.58	1.50	0.26	0.08	2.28	0.10	-0.05	-1.00	-1.21	-0.13	1.23	2.94	-1.11	0.12	0.98	1.64	1.95	
0.32	1.19	0.07	1.68	1.59	1.56	1.36	0.19	0.09	1.30	0.10	1.09	0.20	0.02	1.36	0.14	1.05	0.03	0.14	0.06	0.27	0.43	1.29	1.18	0.67	1.18	1.11	0.07	0.20	1.21	0.11	0.07	0.03	1.43	0.11	0.22	0.33	
-2.13	-1.11	-1.54	-6.55	-7.75	-3.09	2.68	1.38	1.59	5.06	2.12	1.21	1.31	-1.04	-1.39	2.66	0.91	-1.53	-2.73	-1.29	1.46	2.58	4.85	4.13	6.92	2.58	2.66	2.82	-1.17	-2.93	6.46	2.30	-1.39	-3.40	-2.37	3.34	9.29	
0.01	0.98	0.30	0.01	0.00	0.07	0.06	0.14	0.02	0.00	0.00	0.11	0.12	0.30	0.79	0.02	0.69	0.72	0.03	0.96	0.02	0.02	0.08	0.00	0.02	0.11	0.19	0.00	0.14	0.01	0.01	0.00	0.51	0.01	0.01	0.01	0.01	
-1.40	-1.46	x	-1.32	-1.19	-1.35	-1.19	1.17	-1.33	-1.45	-1.50	-1.50	1.07	1.10	1.58	3.20	1.45	-1.20	-1.39	-1.53	2.66	-1.49	-1.64	-1.68	1.14	1.15	-1.03	-1.23	-1.31	2.16	1.01	1.61	-1.20	-1.83	1.41	-1.29	-1.10	
0.46	0.36	×	0.12	0.75	0.30	0.26	0.48	0.34	0.03	0.05	0.24	0.78	0.45	0.52	0.02	0.01	0.62	0.39	0.19	0.06	0.32	0.15	0.01	0.73	0.48	0.86	0.19	0.12	0.00	0.94	0.04	0.54	0.08	0.12	0.06	0.49	
-1.77	-1.29	N/A	-3.94	-4.47	-2.22	0.75	1.28	0.13	1.81	0.31	-0.15	1.19	0.03	0.10	2.93	1.18	-1.37	-2.06	-1.41	2.06	0.55	1.61	1.23	4.03	1.87	0.82	0.80	-1.24	-0.39	3.74	1.96	-1.30	-2.62	-0.48	1.03	4.10	
0.37	0.17	N/A	2.62	3.28	0.87	1.94	0.11	1.46	3.26	1.81	1.36	0.12	1.07	1.49	0.27	0.27	0.17	0.67	0.12	0.60	2.04	3.25	2.91	2.89	0.72	1.85	2.03	0.07	2.55	2.73	0.34	0.10	0.78	1.89	2.32	5.20	
1.04	1.30	-1.33	1.24	1.55	1.44	1.01	1.71	-1.21	-1.22	-1.15	-1.39	1.33	-1.26	1.76	1.35	-1.24	-1.05	-1.15	1.17	-1.05	-1.09	-1.02	-1.33	-1.04	1.42	1.18	1.23	1.09	2.41	1.44	1.56	1.21	1.14	1.19	1.92	1.15	
0.42	0.50	0.12	0.08	0.02	0.29	0.94	0.05	0.50	0.08	0.45	0.20	0.28	0.06	0.31	0.50	0.40	0.86	0.76	0.22	0.72	0.26	0.98	0.13	0.68	0.14	0.46	0.26	0.24	0.02	0.01	0.01	0.66	0.32	0.56	0.00	0.52	
-0.98	1.61	x	1.24	1.16	-1.03	-1.29	1.50	1.07	-1.67	-1.80	-1.36	1.18	-1.26	1.51	1.21	-1.06	1.44	1.25	1.73	-1.34	-1.40	-2.72	-2.44	-1.15	1.47	-1.09	-1.11	-1.10	2.50	-1.10	1.23	1.67	1.12	1.11	1.64	0.98	
0.76	0.33	x	Х	0.62	Х	0.10	0.02	0.87	0.04	0.04	0.68	0.44	0.45	0.43	0.53	0.57	0.57	Х	0.01	0.13	0.23	0.03	0.04	0.23	0.30	0.79	0.70	0.48	0.00	0.34	0.05	0.09	0.21	0.73	0.12	0.50	
0.03	1.46	N/A	1.24	1.36	0.21	-0.14	1.61	-0.07	-1.45	-1.48	-1.38	1.26	-1.26	1.64	1.28	-1.15	0.20	0.05	1.45	-1.20	-1.25	-1.87	-1.89	-1.10	1.45	0.04	0.06	-0.01	2.46	0.17	1.40	1.44	1.13	1.15	1.78	1.07	
1.01	0.16	N/A	0.00	0.20	1.24	1.15	0.11	1.14	0.23	0.33	0.01	0.08	0.00	0.13	0.07	0.09	1.25	1.20	0.28	0.15	0.15	0.85	0.56	0.05	0.03	1.14	1.17	1.10	0.04	1.27	0.17	0.23	0.01	0.04	0.14	0.08	

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pyridoxamine 5-phosphate oxidase amine oxidase <i>rplT</i> 50S ribosomal protein L20 50S ribosomal protein L35	ŏ ŏ	SPO3598 SPO3599
pyridoxamine 5'-phosphate amine oxidase 50S ribosomal protein L20	õ	SPO359
5'-phosphate		
5'-phosphate	Γ	SPO3597
	6	SPO3596
AraC family transcriptional regulator	Ū	SPO3595
phenylalanyl-tRNA synthetase subunit alpha pheS (EC:6.1.1.20)	4	SPO3594
en lfatace	ا در	SP03593
LysR family transcriptional regulator	0,	SPO3592
hypothetical protein	- 0	SPO3590
hypothetical protein	9	SPO3589
hypothetical protein	œ	SPO3588
hypothetical protein	7	SPO3587
<i>fadH</i> 2,4-dienoyl-CoA reductase (EC:1.3.1.34)		SPO3586
NUDIX family hydrolase NudH subfamily hydrolase	Ŭn .	SPO3585
pecS transcriptional regulator PecS drug/metabolite transporter family membrane		SPO3583
PfkB family kinase		SPO3582
nth endonuclease III (EC:4.2.99.18)	<u>`</u>	SPO3581
isochorismatase	0	SPO3580
MarR family transcriptional regulator	9	SPO3579
ADA regulatory protein	80	SPO3578
hypothetical protein	7	SPO3577
OmpA domain-containing protein	6	SPO3576
MerR family transcriptional regulator	U.	SPO3575
short chain dehydrogenase	4	SPO3574
NADPH-dependent FMN reductase domain- containing protein	ω	SPO3573
isochorismatase	2	SPO3572
AraC family transcriptional regulator	1	SPO3571
hypothetical protein	0	SPO3570
recR recombination protein RecR	9	SPO3569
hypothetical protein	õõ	SPO3568
ribonuclease T2 family protein	7	SPO3567
hypothetical protein	6	SPO3566
hypothetical protein	Ċ1	SPO3565
permease	4	SPO3564
ArsR family transcriptional regulator	ω	SPO3563

1.46	-1.13	-1.12	-1.55	1.29	Х	1.47	1.63	1.08	1.35	1.80	2.89	1.20	-1.69	1.28	1.30	Х	-1.43	-1.03	-1.01	-1.26	1.32	-1.20	2.15	0.89	-1.88	1.19	-1.43	-2.21	х	1.05	1.37	1.02	-1.37	-1.16	-1.79	0.95	1.13
0.42	0.01	0.02	0.53	0.16	x	0.53	0.31	0.88	0.17	0.23	0.00	х	0.00	0.83	0.69	x	0.18	0.70	0.86	0.07	0.04	0.15	0.02	0.34	0.18	0.51	0.75	0.39	x	0.09	0.20	0.54	0.51	0.10	0.63	0.37	0.09
2.00	-1.28	1.11	1.35	1.03	-1.33	1.31	-1.18	1.18	1.37	1.01	1.06	-1.05	1.11	-1.01	1.18	1.04	-1.06	1.10	-1.05	1.07	1.02	-1.02	-1.02	-2.02	-1.49	-1.44	-1.92	-1.34	1.06	1.22	1.09	1.11	1.39	-1.82	-1.15	1.02	-1.04
0.04	0.12	0.68	0.40	0.96	0.10	0.67	0.57	0.45	0.03	0.98	0.48	0.81	0.46	0.95	0.32	0.60	0.54	0.49	0.63	0.68	0.79	0.89	0.95	0.06	0.31	0.20	0.12	0.62	0.35	0.40	0.64	0.73	0.27	0.11	0.63	0.91	0.80
1.73	-1.21	-0.01	-0.10	1.16	N/A	1.39	0.23	1.13	1.36	1.41	1.98	0.08	-0.29	0.14	1.24	N/A	-1.25	0.04	-1.03	-0.10	1.17	-1.11	0.57	-0.57	-1.69	-0.13	-1.68	-1.78	N/A	1.14	1.23	1.07	0.01	-1.49	-1.47	0.98	0.04
0.27	0.08	1.12	1.45	0.13	N/A	0.08	1.41	0.05	0.01	0.40	0.92	1.13	1.40	1.15	0.06	N/A	0.18	1.07	0.02	1.17	0.15	0.09	1.59	1.45	0.19	1.32	0.25	0.44	N/A	0.09	0.14	0.05	1.38	0.33	0.32	0.04	1.09
1.48	3.35	3.53	-1.99	1.52	2.14	4.49	1.51	-2.44	1.14	-4.29	-1.19	-2.08	-1.53	-1.67	1.86	1.99	-1.59	1.03	2.60	3.37	1.45	-1.32	-1.27	2.22	-1.53	-1.88	1.67	-3.48	2.06	0.79	1.82	-1.27	1.65	3.98	0.87	0.93	-1.06
0.20	0.00	0.00	0.33	0.09	0.01	0.09	0.49	0.09	0.19	0.03	0.37	0.07	0.02	0.10	0.15	0.02	0.18	0.93	0.02	0.00	0.00	0.10	0.63	0.10	0.55	0.00	0.00	0.21	0.01	0.00	0.01	0.15	0.22	0.00	0.83	0.27	0.61
5.01	-1.33	-1.38	1.21	-1.10	-1.00	-1.40	1.43	-1.09	-1.21	1.42	-1.52	-1.45	-1.09	1.16	-1.08	-1.56	-1.01	-1.74	0.99	-1.07	-1.16	1.24	1.05	1.23	-1.36	1.31	-1.32	-1.50	-1.50	-1.52	-1.67	1.00	1.18	-1.09	-1.06	-1.65	-1.38
0.01	0.07	0.20	0.18	0.80	0.89	0.42	0.38	0.42	0.29	0.31	0.11	x	0.82	0.76	0.51	0.05	0.82	0.02	0.59	0.62	0.25	0.60	0.55	0.42	0.12	0.29	0.02	0.34	0.09	0.02	0.01	0.40	0.55	0.21	0.73	0.12	0.08
3.25	1.01	1.08	-0.39	0.21	0.57	1.55	1.47	-1.77	-0.04	-1.44	-1.36	-1.77	-1.31	-0.26	0.39	0.22	-1.30	-0.36	1.79	1.15	0.15	-0.04	-0.11	1.73	-1.45	-0.29	0.18	-2.49	0.28	-0.37	0.08	-0.14	1.42	1.45	-0.10	-0.36	-1.22
1.77	2.34	2.46	1.60	1.31	1.57	2.95	0.04	0.68	1.18	2.86	0.16	0.31	0.22	1.42	1.47	1.78	0.29	1.39	0.81	2.22	1.31	1.28	1.16	0.50	0.09	1.60	1.50	0.99	1.78	1.15	1.75	1.14	0.23	2.54	0.96	1.29	0.16
-1.85	-1.98	-1.50	1.63	1.10	1.04	-1.06	1.10	-1.95	1.41	-1.26	1.15	-0.94	1.58	1.48	-1.18	1.13	-1.09	1.16	-1.48	1.33	1.30	1.20	-1.08	1.28	1.39	-1.00	1.06	-1.09	-1.07	-1.13	-1.04	1.19	1.39	-1.55	1.04	1.20	-1.24
0.03	0.04	0.05	0.03	0.82	0.52	0.80	0.83	0.03	0.02	0.30	0.30	0.53	0.01	0.37	0.04	0.27	0.21	0.45	0.06	0.09	0.11	0.23	0.78	0.41	0.09	0.81	0.88	0.89	0.79	0.66	0.26	0.06	0.21	0.00	1.00	0.05	0.11
-3.14	-1.97	-1.91	-1.02	-1.13	-1.21	-1.57	1.40	-1.51	-1.28	1.04	1.35	Х	1.58	3.12	-1.55	-0.96	-1.05	-1.26	-1.43	1.13	-1.08	1.27	-0.92	1.16	1.22	-1.25	1.11	1.38	-1.32	-1.43	-1.20	1.14	1.08	-1.39	0.95	1.59	-1.16
0.01	0.08	0.04	0.66	0.88	0.57	0.35	0.44	0.05	0.33	0.98	0.08	x	0.01	0.13	0.01	0.34	0.70	0.11	0.08	0.28	0.67	0.57	0.16	0.84	0.55	0.20	0.96	0.52	×	0.04	0.13	0.56	0.73	0.15	0.71	0.02	0.48
-2.50	-1.98	-1.71	0.31	-0.01	-0.09	-1.32	1.25	-1.73	0.06	-0.11	1.25	N/A	1.58	2.30	-1.37	0.08	-1.07	-0.05	-1.46	1.23	0.11	1.24	-1.00	1.22	1.31	-1.12	1.09	0.15	-1.20	-1.28	-1.12	1.17	1.24	-1.47	1.00	1.40	-1.20
0.64	0.01	0.20	1.33	1.12	1.13	0.26	0.15	0.22	1.35	1.15	0.10	N/A	0.00	0.82	0.19	1.05	0.02	1.21	0.03	0.10	1.19	0.04	0.08	0.06	0.09	0.13	0.03	1.24	0.13	0.15	0.08	0.03	0.16	0.08	0.05	0.20	0.04

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SPO3639	SPO3638	SPO3637	SPO3636	SPO3635	SPO3634	SPO3633	SPO3632	SPO3631	SPO3630	SPO3629	SPO3628	SPO3627	SPO3626	SPO3625	SPO3623	SPO3622	SPO3621	SPO3620	SPO3619	SPO3618	SPO3617	-	SPO3615	SPO3614	SPO3613	SPO3612	SPO3611	SPO3610	SPO3609	SPO3608	SPO3607	SPO3606	SPO3605	SPO3604	SPO3603	SPO3602	3PU30UI
		uvrC		pgsA	moaD				ubiA				gshA	cspA								accA							linC-2	mclA							
K+-dependent Na+/Ca+ exchanger-like protein	short chain dehydrogenase	excinuclease ABC subunit C	beta-lactamase	CDP-diacylglycerolglycerol-3-phosphate 3- phosphatidyltransferase (EC:2.7.8.5)	molybdopterin converting factor subunit 1	molybdopterin converting factor subunit 2	hypothetical protein	OmpA domain-containing protein	4-hydroxybenzoate polyprenyltransferase (EC:2.5.1)	16S ribosomal RNA methyltransferase RsmE	acetyltransferase	hypothetical protein	glutamatecysteine ligase (EC:6.3.2.2)	cold shock protein CspA	hypothetical protein	hypothetical protein	GntR family transcriptional regulator	sulfonate ABC transporter substrate-binding protein	sulfonate ABC transporter ATP-binding protein	sulfonate ABC transporter permease	peptidoglycan-binding protein	acetyl-CoA carboxylase carboxyltransferase subunit alpha (EC:6.4.1.2)	AraC family transcriptional regulator	hypothetical protein	amino acid transporter LysE	AsnC family transcriptional regulator	hypothetical protein	hypothetical protein	short chain dehydrogenase (EC:1.1.1)	malyl-CoA lyase (EC:4.1.3.24)	hypothetical protein	mandelate racemase	hypothetical protein	D-amino acid aminotransferase	thioesterase	hypothetical protein	in-torinyigiutamate annuonyuroiase

1.16	1.54	1.08	1.24	1.04	1.09	-1.11	1.54	1.19	2.69	1.05	1.04	1.22	1.71	-1.74	1.27	-1.17	1.08	1.37	1.30	1.77	0.95	1.48	1.29	-1.00	2.73	1.61	-2.14	1.06	х	-1.91	1.33	1.86	3.43	4.07	1.21	1.38	-1.40
0.74	0.09	0.29	0.81	0.97	0.99	0.63	0.02	0.78	0.03	0.79	0.81	0.96	0.43	0.00	0.07	0.30	0.52	0.24	0.20	0.65	0.13	0.34	0.80	0.63	x	0.05	0.08	1.00	×	0.00	0.86	0.08	0.15	0.04	0.54	0.04	0.06
-1.13	-1.24	-1.37	-1.50	1.37	1.88	1.75	-1.07	-1.41	1.56	-1.17	-1.08	-1.13	1.15	1.33	1.42	1.10	1.21	-1.11	-1.07	-1.15	-1.56	1.21	16.00	4.70	2.34	-1.10	-1.16	1.21	1.04	-1.18	-1.09	1.98	1.58	1.20	-1.21	-1.13	-1.13
0.77	0.13	0.06	0.08	0.27	0.06	0.08	0.48	0.14	0.04	0.47	0.34	0.62	0.75	0.09	0.10	0.19	0.46	0.51	0.75	0.71	0.27	0.18	0.00	0.02	0.07	0.71	0.75	0.75	0.77	0.10	0.97	0.02	0.01	0.51	0.68	0.48	0.55
0.02	0.15	-0.15	-0.13	1.21	1.49	0.32	0.24	-0.11	2.13	-0.06	-0.02	0.05	1.43	-0.21	1.35	-0.03	1.15	0.13	0.12	0.31	-0.31	1.35	8.65	1.85	2.54	0.26	-1.65	1.14	N/A	-1.55	0.12	1.92	2.51	2.64	0.00	0.13	-1.27
1.15	1.39	1.23	1.37	0.17	0.40	1.43	1.31	1.30	0.57	1.11	1.06	1.18	0.28	1.54	0.08	1.14	0.06	1.24	1.19	1.46	1.25	0.14	7.36	2.85	0.20	1.36	0.49	0.08	N/A	0.37	1.21	0.06	0.93	1.44	1.21	1.26	0.14
2.36	2.47	1.07	-1.82	1.71	1.33	-1.29	2.84	2.18	2.86	1.89	2.01	1.98	3.21	1.45	6.31	3.12	1.08	-1.25	0.95	1.57	1.09	3.26	-1.45	-1.19	6.17	-1.08	-2.94	1.49	1.64	-1.34	-1.73	3.19	7.22	6.01	1.01	0.89	1.68
0.11	0.11	0.21	0.07	0.01	0.89	0.60	0.02	0.13	0.01	0.03	0.02	0.73	0.08	0.13	0.00	0.01	0.40	0.25	0.56	0.65	0.39	0.02	0.23	0.91	0.03	0.31	0.06	0.63	0.01	0.02	0.82	0.01	0.01	0.05	0.04	0.17	0.03
-1.62	-1.09	-1.33	1.96	-1.04	1.03	-1.05	-1.07	1.14	-1.20	-1.18	-1.81	-1.05	1.11	-1.07	-1.60	-1.15	1.20	1.60	1.50	-1.03	-1.11	-1.21	-1.03	1.19	-1.61	-1.78	-1.08	-1.11	-1.05	-1.11	-1.06	-2.11	-1.87	-1.62	-1.16	1.00	-1.19
0.26	0.55	0.06	0.03	0.13	0.98	0.75	0.44	0.73	0.15	0.25	0.02	0.73	0.89	0.21	0.01	0.08	0.18	0.18	0.03	0.90	0.10	0.16	0.61	0.55	0.18	0.07	0.78	0.86	0.91	0.15	0.97	0.04	0.02	0.06	0.70	0.35	0.31
0.37	0.69	-0.13	0.07	0.34	1.18	-1.17	0.89	1.66	0.83	0.36	0.10	0.47	2.16	0.19	2.36	0.99	1.14	0.18	1.23	0.27	-0.01	1.03	-1.24	0.00	2.28	-1.43	-2.01	0.19	0.30	-1.23	-1.40	0.54	2.68	2.20	-0.08	0.94	0.25
1.99	1.78	1.20	1.89	1.38	0.15	0.12	1.96	0.52	2.03	1.54	1.91	1.52	1.05	1.26	3.96	2.14	0.06	1.43	0.27	1.30	1.10	2.24	0.21	1.19	3.89	0.35	0.93	1.30	1.35	0.12	0.34	2.65	4.55	3.82	1.09	0.05	1.44
1.19	1.16	-1.03	1.41	1.24	1.61	-1.04	1.36	-1.07	1.22	1.05	1.26	1.35	-1.05	1.30	-1.25	-1.37	-1.27	1.83	1.64	1.34	1.35	-1.09	1.05	-1.58	-1.04	1.17	1.25	1.26	1.28	-1.05	1.03	1.78	-1.40	1.18	-1.07	-1.32	-1.59
0.74	0.42	0.44	0.04	0.32	0.11	0.91	0.06	0.57	0.34	0.49	0.08	0.32	0.75	0.01	0.22	0.02	0.03	0.18	0.02	0.42	0.01	0.14	0.64	0.06	0.99	0.07	0.50	0.73	0.12	0.63	0.99	0.11	0.08	0.24	0.57	0.05	0.01
1.18	1.14	1.18	1.41	-1.08	1.43	-1.24	1.16	1.22	-1.23	-1.14	-1.04	1.61	-1.28	-1.12	-2.20	-1.94	-1.30	1.93	1.46	1.06	1.37	-1.17	25.30	3.04	-1.64	-1.09	1.54	-0.95	1.07	1.13	1.31	1.45	-1.22	1.23	1.40	-1.28	-2.13
0.81	0.58	0.34	0.02	0.18	0.28	0.44	0.17	0.64	0.04	0.20	0.66	0.23	0.42	0.17	0.02	0.01	0.06	0.03	0.09	0.80	0.26	0.14	0.00	0.02	х	0.95	0.24	0.89	×	0.67	х	0.02	0.11	0.66	0.52	0.07	0.00
1.19	1.15	0.08	1.41	0.08	1.52	-1.14	1.26	0.08	-0.01	-0.04	0.11	1.48	-1.17	0.09	-1.73	-1.66	-1.29	1.88	1.55	1.20	1.36	-1.13	13.18	0.73	-1.34	0.04	1.40	0.16	1.18	0.04	1.17	1.62	-1.31	1.21	0.17	-1.30	-1.86
0.01	0.01	1.11	0.00	1.16	0.09	0.10	0.10	1.15	1.23	1.10	1.15	0.13	0.12	1.21	0.48	0.29	0.02	0.05	0.09	0.14	0.01	0.04	12.13	2.31	0.30	1.13	0.14	1.10	0.11	1.09	0.14	0.17	0.09	0.03	1.24	0.02	0.27

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### Chapter 10: Appendix

argE-2	catB	hem C hem F	hem E
LysR family transcriptional regulator hypothetical protein muticopper oxidase domain-containing protein hypothetical protein ISSpo7, transposase hypothetical protein acetylornithine deacetylase (EC:3.5.1.16) LysR family transcriptional regulator Rieske (2Fe-2S) domain-containing protein hypothetical protein	HAD superfamily hydrolase D-alanyl-D-alanine carboxypeptidase LamB/YcsF family protein urea amidolyase allophanate hydrolase hypothetical protein TR AP dicarboxylate transporter subunit DctM hypothetical protein solute-binding family 7 protein oxidoreductase, FAD-binding muconate cycloisomerase I (EC:5.5.1.1)	porphobilinogen deaminase (EC.2.5.1.61) guanylate cyclase fatty acid desaturase hypothetical protein coproporphyrinogen III oxidase (EC:1.3.3.3) 7-alpha-hydroxysteroid dehydrogenase methyltransferase ATP-dependent Clp protease, adaptor protein ClpS	S49 family peptidase ABC transporter permease ABC transporter ATP-binding protein hypothetical protein transcriptional regulator enoyl-CoA hydratase (EC:4.2.1.17) CAIB/BAIF family protein uroporphyrinogen decarbox vlase (EC:4.1.1.37)
-1.44 X 1.44 1.30 1.67 X 1.67 X 1.60 1.14 1.01 1.09	-1.68 0.96 -1.14 1.29 -1.04 -1.24 -1.37 -1.31 -2.26 1.05 1.50	1.41 -1.48 0.98 1.58 1.42 1.51 -1.34 0.98	-1.46 1.18 1.22 -1.03 X -0.99 1.30 2.92 2.14
0.47 X 0.72 0.39 0.39 X X 0.03 0.29 0.97 0.93	0.45 0.78 0.07 X 1.00 0.60 0.60 0.02 0.55	0.71 0.02 0.01 0.17 0.17 0.79 0.79 0.21	$0.26 \\ 0.86 \\ 0.83 \\ 0.98 \\ X \\ 0.93 \\ 0.23 \\ 0.24 \\ 0.48 \\ 0.4$
-1.16 -1.41 -1.34 1.10 1.20 X 1.21 1.33 1.02 1.07	-1.22 -1.82 -1.10 -1.19 -1.17 -1.16 -2.12 -2.24 -2.94 -2.00 1.05	1.62 -1.15 -1.18 1.29 1.51 1.51 -1.03 1.13 1.13	-1.42 1.04 1.08 1.40 -1.40 -1.22 -1.10 1.99 3.30
0.79 0.57 0.87 0.87 0.43 0.43 0.43 0.58 0.64	0.26 0.07 0.78 0.31 0.27 0.27 0.04 0.05 0.04 0.05 0.04 0.24	0.10 0.17 0.57 0.51 0.51 0.83 0.83	0.39 0.94 0.48 0.22 0.14 0.73 0.73
-1.30 N/A 0.05 1.20 1.44 N/A N/A N/A 1.47 1.08 1.04	-1.45 -0.43 -1.08 0.05 -1.11 -1.44 -1.75 -2.28 -2.28 -2.60 -0.48 1.28	1.52 -1.32 -0.10 1.44 1.47 0.24 -0.11	-1,44 1.11 1.15 0.19 N/A -1.11 0.10 0.10 2.46 2.72
0.14 N/A 1.39 0.10 0.23 N/A 0.14 0.06 0.03	0.23 1.39 0.06 1.24 0.06 0.19 0.38 0.97 0.34 1.53 0.23	0.11 0.17 1.08 0.15 0.05 1.27 1.27 1.24	0.02 0.07 0.07 1.22 N/A 0.11 1.20 0.47 0.58
-1.27 -1.16 -1.73 -1.45 3.28 3.28 X 4.71 -1.02 -1.66 -2.08	-1.79 0.79 1.11 3.40 2.23 1.72 1.99 1.04 2.49	2.69 -2.12 -17.70 -1.40 1.82 1.94 -1.12 1.38	1.55 1.77 1.68 1.58 -0.87 -1.25 -4.28 -1.63 3.14
0.88 X 0.02 0.40 0.12 0.13 0.13 0.13 0.26 0.26	0.52 0.02 0.16 0.01 0.00 0.01 0.01 0.01 0.41 0.41 0.23 0.23	0.26 0.02 0.00 0.04 0.03 0.63 0.71	0.12 0.07 0.07 0.16 0.16 0.15 0.15 0.10
-1.40 -1.03 1.87 2.12 1.26 X -1.36 1.09 1.20 1.04	1.30 0.97 1.26 1.97 2.10 4.35 3.67 3.67 3.47 4.17 4.17	 1.10 1.26 1.46 1.20 1.07 1.07 1.07 1.07	-1.40 1.24 -1.02 -1.19 -1.56 1.06 1.70 1.70 1.36
0.51 1.00 0.20 0.22 0.22 X 0.22 0.59	0.45 0.12 0.03 0.04 0.00 0.00 0.00 0.01 0.00 0.01 0.00	0.84 0.16 0.62 0.44 0.94 0.51	0.22 0.45 0.59 0.56 0.56 0.59 0.59 0.04
-1.34 -1.10 0.07 0.34 2.27 N/A 1.68 0.04 -0.12 -1.57	-0.25 0.88 1.19 2.69 2.32 3.34 2.41 2.41 2.83 2.26 3.33 -1.31	-1.12 -0.43 -0.43 -8.12 -0.10 1.45 1.51 -1.12 -1.12	0.08 1.51 0.33 0.20 -1.22 -0.10 -1.29 0.09
0.06 0.06 1.80 1.79 1.01 1.01 1.04 1.06 1.32 1.35	1.55 0.09 0.72 0.72 1.01 0.69 0.84 1.22 0.84	0.80 1.69 9.58 1.30 0.44 0.01	1.48 0.27 1.35 1.39 0.34 1.16 2.99 2.99
-1.15 -1.71 -1.17 1.14 0.97 X -1.75 1.31 -1.11 1.11	-1.08 1.09 -1.06 -1.48 -1.29 -1.17 1.03 -1.17 1.03 -1.22 1.41 -1.39	-1.46 -1.11 -0.98 1.08 1.24 0.99 1.10 -1.23	1.21 1.04 -1.04 1.74 -1.40 1.37 1.06 -1.06
0.80 0.17 0.38 0.68 0.65 0.65 X 0.20 0.11 0.71 0.71 0.68	0.79 0.75 0.63 0.02 0.08 0.40 0.40 0.92 0.14 0.14 0.37 0.40	0.21 0.22 0.55 0.77 0.87 0.87 0.87 0.21	0.38 0.94 0.62 0.28 0.28 0.28 0.70 0.72
-1.23 1.19 1.41 1.75 1.07 X -2.95 X -1.31 -1.01 1.02	1.30 1.03 -1.02 -1.86 -2.12 -1.13 1.06 1.15 2.12 -1.21 -1.21	-1.06 1.56 3.50 1.26 -1.28 1.02 1.02 1.09	1.60 -1.07 -1.21 -0.99 X 1.28 2.55 2.30 -1.46
0.87 0.48 0.37 0.67 0.67 X 0.07 X 0.07 X 0.05 0.92	0.25 0.37 0.80 0.09 0.09 0.97 0.97 0.56 0.27 0.27 0.21	0.47 0.47 0.00 0.62 0.62 0.95 0.95 0.48	0.16 0.73 0.19 0.94 X 0.05 0.03 0.13
-1.19 -0.26 0.12 1.45 1.02 N/A -2.35 N/A -1.21 0.05	0.11 1.06 -1.04 -1.67 -1.71 -1.71 -1.15 -0.04 1.77 -1.30 -1.30	-1.26 0.23 1.26 1.17 -0.02 1.01 1.10 1.10	1.41 -0.02 -1.13 0.38 N/A 1.33 1.81 1.81 1.81
0.04 1.45 1.29 0.31 0.60 N/A 0.60 N/A 0.10 1.06	1.19 0.03 0.12 0.19 0.42 0.02 1.19 0.35 0.09	0.20 1.34 2.24 0.09 1.26 0.01 0.01	0.20 1.06 1.36 1.36 0.05 0.75 1.68

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#### SPO3714 SPO3711 SPO3704 SPO3715 SPO3713 SPO3712 SPO3681 SPO3716 SPO3710 SPO3709 SPO3703 SPO3702 SPO3701 SPO3700 SPO3699 SPO3698 SPO3697 SPO3696 SPO3695 SPO3694 SPO3693 SPO3692 SPO3691 SPO3690 SPO3689 SPO3688 SPO3687 SPO3686 SPO3685 SPO3684 SPO3682 SPO3708 SPO3707 SPO3706 SPO3705 SPO3680 asd badA-1 gtdA-1 fcsboxA boxBaspartate-semialdehyde dehydrogenase (EC:1.2.1.11) endoribonuclease L-PSP benzoyl-CoA oxygenase, A subunit anaerobic benzoate catabolism transcriptional hydrolase benzoate-coenzyme A ligase (EC:6.2.1.25) gentisate 1,2-dioxygenase (EC:1.13.11.4) hypothetical protein carbonic anhydrase hypothetical protein hypothetical protein hypothetical protein hypothetical protein branched-chain amino acid ABC transporter branched-chain amino acid ABC transporter binding protein branched-chain amino acid ABC transporter ATPbinding protein branched-chain amino acid ABC transporter ATPsubstrate-binding protein branched-chain amino acid ABC transporter hypothetical protein benzoyl-CoA oxygenase subunit B benzoyl-CoA-dihydrodiol lyase (EC:4.2.1.17) hypothetical protein feruloyl-CoA synthase hypothetical protein TRAP dicarboxylate transporter subunit DctP salicylate hydroxylase (EC:1.14.13.1) fumarylacetoacetate hydrolase MarR family transcriptional regulator extradiol ring-cleavage dioxygenase decarboxylase, pyridoxal-dependent hypothetical protein LacI family transcriptional regulator hypothetical protein Rieske (2Fe-2S) domain-containing protein permease permease regulator TRAP dicarboxylate transporter subunit DctM HpcH/HpaI aldolase 2.21 0.89 0.99 1.03 1.08 1.19 -1.86 1.12 1.10 Ė 1.66 1.58 1.141.59 1.1 1.06 1.68 1.34 1.36 1.09 1.14 1.06 1.13 1.24 1.28 × 1.27 1.09 1.48 .42 0.72 0.39 0.19 0.98 0.91 0.82 0.72 0.960.52 0.05 0.16 0.90 0.20 0.010.13 0.990.47 0.09 0.06 0.95 0.05 0.86 0.98 0.19 0.00 0.02 0.63 0.34 0.74 0.00 0.88 0.03 0.00 0.00 0.02 $\times$ -1.34 -1.57 -1.32 -1.39 1.19 -1.07 1.35 1.35 1.90 1.04 1.13 0.19 1.061.20 1.02 1.7 1.07 1.12 1.10 1.31 .09 4 4 4 0.31 0.18 0.02 0.89 0.69 0.65 0.57 0.90 0.01 0.340.45 0.080.220.05 0.99 0.58 0.41 0.14 0.44 0.37 0.42 0.57 0.35 0.00 0.03 0.43 0.02 0.78 0.78 0.60 0.44 0.27 0.80 0.55 0.79 0.02 0.03 N/A -0.40 -0.25 0.10 1.30 1.78 -0.01 -1.20 -1.59 -0.19 -0.67 <u>-</u> -0.05 1.39 1.341.75 1.08 1.091.20 -1.67 1.35 1.23 0.43 0.14 0.21 0.04 0.27 0.24 0.14 0.02 0.16 0.15 0.10 0.33 1.43 1.42 1.32 N/A 0.05 0.03 0.01 0.27 1.77 1.15 1.23 1.09 1.331.41 1.47 1.361.25 1.29 1.29 1.03 1.341.11 1.23 1.17 -11.302.05 6.31 0.98 2.52 -1.20 -3.29 1.26 -2.83 4.21 4.326.9 × 1.6 0.72 0.06 0.04 0.19 0.10 0.84 0.04 0.230.00 0.03 0.04 0.01 0.07 0.010.00 0.06 0.09 0.04 0.160.93 0.35 0.04 0.00 0.07 0.52 0.01 0.36 0.09 0.13 0.49 0.02 0.01 0.03 0.01 0.03 × -1.01 2.03 2.95 5.37 1.91 -1.16 3.54 4.26 2.67 1.93 1.55 1.16 1.07 1.00 1.67 1.37 1.43 1.62 1.96 1.55 1.62 1.81 1.57 1.47 1.05 1.27 1.28 1.43 1.42 1.30 1.30 0.71 0.38 0.98 0.46 0.14 0.49 0.08 0.13 0.140.01 0.610.220.03 0.26 0.31 0.13 0.19 0.07 0.26 0.26 0.320.05 0.15 0.01 0.87 0.51 0.410.05 0.10 0.47 0.73 0.00 0.02 0.03 0.06 0.02 N/A 0.68 -0.03 0.330.41 0.49 -1.30-1.18 -0.72 0.10-0.49 -0.06 0.100.22 0.080.11 -0.77 -0.84 -0.52 2.51 1.17 1.45 1.52 ... 82 0.53 N/A 0.05 3.81 0.19 1.71 2.55 2.37 3.78 8.34 0.44 2.92 2.17 2.38 2.10 2.92 3.14 2.27 2.11 5.36 1.16 2.32 0.46 0.18 2.00 1.59 1.84 1.31 1.64 1.37 1.71 1.70 1.33 1.14 1.19 1.33 2.18 -0.99 1.04 1.19 1.38 1.19 1.15 1.03 1.03 1.19 1.53 1.38 1.56 1.03 1.18 1.20 1.24 1.04 1.24 1.22 1.05 0.30 0.68 0.33 0.54 0.15 0.340.340.97 0.97 0.11 0.00 0.48 0.66 0.07 0.90 0.66 0.36 0.58 0.33 0.99 0.64 0.17 0.95 0.940.47 0.21 0.14 0.01 0.970.08 0.60 0.85 0.16 0.61 0.58 0.89 -1.83 2.14-0.97 2.01 2.54 -1.41 1.27 -0.95 1.141.12 2.34 1.06 1.37 1.77 1.70 1.45 1.14 1.35 1.07 1.28 1.43 1.88 1.40 1.43 1.22 1.19 1.23 1.30 1.27 1.02 0.19 X 0.08 0.01 0.06 0.620.24 0.36 0.24 0.04 0.250.02 0.13 0.03 0.95 0.56 0.98 0.84 0.26 0.35 0.16 0.24 0.22 0.02 0.140.29 0.36 0.16 0.50 0.02 0.54 0.32 0.05 0.48 0.23 0.02 0.140.06 0.05 0.07 0.10 2.05 2.161.13 0.21 0.63 1.05 1.04 1.62 1.42 1.09 1.12 1.16 -1.14 1.34 1.23 1.28 1.50 1.20 1.22 1.22 1.23 .60 .28 6 0.05 0.02 1.72 0.09 0.27 0.09 0.20 0.20 0.01 0.04 0.49 0.05 0.07 1.21 0.02 0.02 0.08 0.100.10 1.23 0.28 0.02 0.29 1.19 0.09 0.16 0.05 0.04 0.13 1.17 1.09 1.05 0.41 1.18 1.15

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-1.19	Х	х	Х	14.60	10.20	-3.14	1.10	-1.15	1.22	1.04	1.76	2.18	2.27	2.00	1.44	1.81	1.22	1.88	Х	-1.76	-1.20	1.20	2.01	1.32	1.35	1.20	1.13	1.18	-1.30	Х	-2.15	-0.97	Х	3.12	2.17	1.10	1.19
0.03	x	x	x	0.00	0.00	0.04	0.49	0.71	0.12	0.53	0.11	0.01	0.02	0.00	0.32	0.00	0.20	0.25	×	0.03	0.13	0.83	0.04	0.73	0.67	0.87	0.95	0.74	0.25	×	0.11	0.43	×	0.00	0.01	0.18	0.45
-1.08	1.70	1.26	1.07	4.58	3.57	-1.37	1.07	-1.84	-1.54	0.97	1.50	1.10	1.28	1.37	1.37	-1.01	1.03	1.12	х	1.04	1.25	-1.45	1.06	1.00	0.99	-1.09	-1.55	-1.58	1.03	-1.13	-1.12	1.01	1.60	1.45	-1.04	1.20 1.19	1 20
0.51	0.04	0.10	0.59	0.00	0.02	0.56	0.71	0.00	0.19	0.79	0.20	0.48	0.35	0.02	0.50	0.90	0.89	0.78	Х	0.59	0.36	0.06	0.87	0.92	0.81	0.68	0.09	0.02	0.96	0.54	0.89	0.66	0.00	0.03	0.80	0.10 0.03	0.10
-1.14	N/A	N/A	N/A	9.59	6.89	-2.26	1.09	-1.50	-0.16	1.00	1.63	1.64	1.78	1.69	1.41	0.40	1.13	1.50	N/A	-0.36	0.03	-0.13	1.54	1.16	1.17	0.05	-0.21	-0.20	-0.14	N/A	-1.64	0.02	N/A	2.29	0.57	1.20	1 20
0.05	N/A	N/A	N/A	5.01	3.32	0.89	0.02	0.35	1.38	0.04	0.13	0.54	0.50	0.31	0.03	1.41	0.10	0.38	N/A	1.40	1.23	1.33	0.47	0.16	0.18	1.15	1.34	1.38	1.17	N/A	0.52	0.99	N/A	0.84	1.61	0.04	0.01
-2.62	2.25	х	-0.97	-1.03	-1.56	-7.31	-1.22	1.87	3.45	2.24	2.40	-2.68	0.91	-1.31	-2.03	2.24	1.34	1.43	Х	-1.42	-1.13	1.71	3.75	2.41	2.62	2.53	2.41	2.56	1.16	-1.62	-1.84	-1.09	1.57	3.51	2.23	1.51	1 45
0.00	0.01	×	0.27	0.08	0.09	0.02	0.20	0.08	0.01	0.41	0.02	0.00	0.14	0.01	0.07	0.00	0.11	0.40	×	0.44	0.37	0.38	0.01	0.22	0.32	0.08	0.01	0.23	0.71	0.22	0.48	0.36	×	0.00	0.06	0.05	0.09
1.75	-5.44	-8.54	-7.34	1.20	1.01	-1.15	1.28	1.21	-1.19	-1.26	-0.99	-1.05	1.65	1.38	-1.04	-1.14	-1.09	1.06	х	-1.28	1.26	-1.23	1.31	1.01	1.17	1.17	1.12	-1.14	-1.04	-0.97	1.10	-1.03	-1.28	-1.11	1.00	1.39 1.29	1.35
0.05	0.01	0.00	0.00	0.15	0.98	0.64	0.19	0.35	0.11	0.42	0.97	0.92	0.03	0.06	0.70	0.17	0.27	0.99	×	0.36	0.19	0.22	0.14	0.80	0.56	0.35	0.78	0.27	0.74	0.78	0.84	0.81	0.24	0.17	0.50	0.02	0.09
-0.44	-1.60	N/A	-4.16	0.09	-0.28	-4.23	0.03	1.54	1.13	0.49	0.70	-1.87	1.28	0.03	-1.54	0.55	0.13	1.25	N/A	-1.35	0.07	0.24	2.53	1.71	1.90	1.85	1.77	0.71	0.06	-1.30	-0.37	-1.06	0.15	1.20	1.61	1.40	1.40
2.19	3.85	N/A	3.18	1.12	1.29	3.08	1.25	0.33	2.32	1.75	1.70	0.81	0.37	1.35	0.50	1.69	1.22	0.18	N/A	0.07	1.20	1.47	1.22	0.70	0.73	0.68	0.65	1.85	1.10	0.32	1.47	0.03	1.43	2.31	0.62	0.11	0.05
1.22	-3.39	-6.48	-5.55	-1.41	1.33	-1.10	1.04	-1.41	-1.23	1.19	-1.09	-0.97	1.19	1.27	-1.34	1.05	-1.08	-1.06	-1.15	1.19	2.06	1.38	1.04	1.17	1.77	1.78	-1.31	1.18	1.16	-1.16	-1.14	-1.59	-1.00	1.17	-1.17	1. <del>7</del> 7 1.38	1.47
0.25	0.01	0.00	0.01	0.10	0.35	0.77	0.78	0.17	0.02	0.80	0.57	0.65	0.43	0.08	0.06	0.83	0.57	0.17	х	0.28	0.00	0.13	0.77	0.39	0.12	0.06	0.18	0.44	0.58	0.56	0.64	0.03	0.52	0.20	0.38	0.20 0.02	0.28
1.20	-5.05	-11.30	-6.32	-1.38	1.44	2.08	-1.07	1.00	-1.26	0.98	-1.25	-1.17	1.08	1.05	1.10	-1.27	-1.30	1.00	Х	1.14	1.32	1.33	1.08	1.44	1.59	2.38	0.98	1.53	1.01	-0.95	1.12	-1.43	-1.18	-1.19	0.95	1.30 1.46	1.30
0.24	0.01	0.00	0.00	0.13	0.38	0.18	0.96	0.26	0.02	0.66	0.18	0.91	0.95	0.64	0.47	0.22	0.21	0.82	x	0.12	0.12	0.20	0.72	0.19	0.08	0.03	0.54	0.12	0.93	0.69	0.63	0.15	0.86	0.04	0.13	0.30	0.10
1.21	-4.22	-8.89	-5.94	-1.40	1.39	0.49	-0.02	-0.21	-1.25	1.08	-1.17	-1.07	1.14	1.16	-0.12	-0.11	-1.19	-0.03	N/A	1.17	1.69	1.36	1.06	1.31	1.68	2.08	-0.16	1.36	1.09	-1.05	-0.01	-1.51	-1.09	-0.01	-0.11	1 1.42	1.39
0.01	0.83	2.41	0.39	0.02	0.05	1.59	1.06	1.20	0.02	0.11	0.08	0.10	0.05	0.11	1.22	1.16	0.11	1.03	N/A	0.03	0.37	0.02	0.02	0.14	0.09	0.30	1.15	0.18	0.08	0.11	1.13	0.08	0.09	1.18	1.06	0.04	0.09

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# Chapter 10: Appendix

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acoB	acoC		acoR																	gltD		gltB		mtgA					map	sfsA					
subunit beta (EC:1.1.1)	subunit (EC:2.3.1.12) acetoin dehydrogenase complex. El component	alkylhydroperoxidase	acetoin catabolism regulatory protein	sugar ABC transporter substrate-binding protein	sugar ABC transporter permease	sugar ABC transporter permease	sugar ABC transporter ATP-binding protein	sugar ABC transporter ATP-binding protein	hypothetical protein	hypothetical protein	hypothetical protein	LysR family transcriptional regulator	oligopeptide/dipeptide ABC transporter ATP- binding protein	oligopeptide/dipeptide ABC transporter A1P- binding protein	oligopeptide/dipeptide ABC transporter permease	oligopeptide/dipeptide ABC transporter permease	ongopepude and transporter substrate- binding protein	NADH ubiquinone oxidoreductase	UDP pyrophosphate phosphatase	oxidoreductase (EC:1.4.1.13)	hypothetical protein	glutamate synthase, large subunit (EC:1.4.1.13)	hypothetical protein	monotunetionai piosyninetic peptidoglycan transglycosylase (EC:2.4.2)	hypothetical protein	glutathione S-transferase	(Fe-S)-binding protein	HAD family hydrolase	methionine aminopeptidase (EC:3.4.11.18)	sugar fermentation stimulation protein A	noryodenum coracior prosynthesis domain- containing protein	acetyltransferase	alkylhydroperoxidase	OmpA domain-containing protein	hypothetical protein
3.20	2.13	2.92	1.12	1.77	1.34	1.73	2.23	1.72	1.31	-1.78	-1.04	-1.42	-1.27	-1.29	-1.36	-1.44	-3.83	3.89	1.83	2.81	2.21	2.22	-2.34	-1.43	1.78	-1.11	-1.32	3.14	1.09	-2.74	1.20	2.61	3.43	2.14	1.90
0.05	0.45	0.07	0.77	0.29	0.12	x	0.19	0.11	0.90	0.37	0.37	0.85	0.02	0.24	0.73	0.70	0.21	0.00	0.58	0.06	0.58	0.02	0.04	0.06	0.46	0.33	0.12	0.03	1.00	×	0.68	0.02	0.01	0.02	0.26
-1.67	-1.30	-1.15	-2.03	-1.05	-1.25	-1.06	-1.08	-1.06	1.08	1.06	1.13	-1.21	-1.44	-1.47	-1.41	-1.44	-2.23	1.72	1.36	2.39	1.92	3.30	-2.43	-1.34	-1.14	1.14	-1.02	-1.13	1.08	-1.21	1.22	1.36	1.04	1.43	1.50
0.01	0.76	0.71	0.50	0.94	0.30	0.92	0.80	0.74	0.79	0.88	0.32	0.69	0.00	0.23	0.39	0.32	0.15	0.02	0.73	0.06	0.30	0.00	0.00	0.30	0.16	0.44	0.82	0.05	0.42	0.08	0.44	0.13	0.95	0.16	0.21
0.77	0.42	0.89	-0.46	0.36	0.05	0.34	0.58	0.33	1.20	-0.36	0.04	-1.32	-1.36	-1.38	-1.39	-1.44	-3.03	2.81	1.60	2.60	2.07	2.76	-2.39	-1.39	0.32	0.01	-1.17	1.01	1.09	-1.98	1.21	1.99	2.24	1.79	1.70
2.44	1.72	2.04	1.58	1.41	1.30	1.40	1.66	1.39	0.12	1.42	1.09	0.11	0.09	0.09	0.02	0.00	0.80	1.09	0.24	0.21	0.15	0.54	0.05	0.04	1.46	1.13	0.15	2.14	0.01	0.77	0.01	0.63	1.20	0.35	0.20
-8.30	-4.28	-2.27	-4.82	-5.62	-2.48	-1.79	-1.14	-1.11	-1.15	2.15	1.80	-1.06	1.16	1.58	1.59	1.12	0.88	3.23	4.35	5.01	4.95	7.10	-2.16	1.12	1.26	2.33	1.82	2.92	2.36	-1.87	2.01	0.98	1.36	1.46	2.08
0.00	0.30	0.03	0.10	0.04	0.04	0.55	0.83	0.50	0.96	0.45	0.02	0.93	0.28	0.08	0.38	0.84	0.46	0.00	0.21	0.02	0.29	0.00	0.02	0.45	0.97	0.01	0.03	0.03	0.02	0.14	0.03	0.30	0.15	0.05	0.02
1.77	1.72	x	1.70	1.53	1.25	-0.97	1.26	-0.98	-1.43	-1.21	-1.03	-1.43	1.05	-1.02	-1.08	-1.04	1.08	- 1.09	-1.20	3.27	1.92	3.14	-1.84	-1.12	1.04	-1.14	-1.50	-1.76	1.05	-1.35	-1.35	2.18	1.84	1.14	1.15
0.04	0.13	×	0.26	0.16	0.17	0.82	0.12	0.79	0.35	0.55	0.61	0.48	0.91	0.72	0.81	0.74	0.83	0.50	0.79	0.00	0.30	0.01	0.04	0.39	0.98	0.15	0.03	0.06	0.95	0.05	0.12	0.00	0.05	0.62	0.07
-3.27	-1.28	N/A	-1.56	-2.05	-0.62	-1.38	0.06	-1.05	-1.29	0.47	0.39	-1.25	1.11	0.28	0.26	0.04	0.98	1.07	1.58	4.14	3.44	5.12	-2.00	0.00	1.15	0.60	0.16	0.58	1.71	-1.61	0.33	1.58	1.60	1.30	1.62
5.04	3.00	N/A	3.26	3.58	1.87	0.41				1.68	1.42	0.18	0.05	1.30	1.34	1.08	0.10	2.16	2.78	0.87	1.52	1.98	0.16	1.12	0.11	1.74	1.66	2.34	0.65	0.26	1.68	0.60	0.24	0.16	0.47
-1.22	-0.97	-1.12	1.15	-1.14	-1.18	-1.16	-1.24	-1.09	-1.15	-1.09	1.41	1.18	-1.25	-1.43	-1.18	-1.01	1.08	1.13	-1.02	-1.06	-1.63	-1.36	-1.06	1.08	1.10	1.45	1.30	-1.10	-1.04	-1.39	1.16	-1.11	1.14	1.47	1.10
0.28	0.88		0.71		0.58	0.72					0.02	0.74	0.32	0.04	_					0.68		0.12	0.45	0.78		0.01		-	0.11	0.17				0.24	
-1.19	1.25	×	1.37	×	-1.25	-1.69	-1.74		1.18	-1.32	-1.04	-1.06	1.02	-1.27	-1.26	1.02						-2.49	-1.98	-1.24				-2.08	1.01		-1.25	1.41	1.46	1.50	
X	0.67	×	X	x	5 0.39		X		0.68		0.52	5 0.96	0.87	0.17		0.97		0.29				0.00	3 0.24	0.42			0.40		0.63	_	0.02		0.04		0.50
-1.21	7 0.14	N/A		N/A	9 -1.22						2 0.19		7 -0.12	-1.35	-1.22		0 1.21	9 0.01					4 -1.52	-			0 0.10		3 -0.02		2 -0.05	6 0.15	4 1.30		0 1.11
0.02	1.11		0.11					4 0.05			1.23	1.12	1.14	0.08			0.13			3 0.27		3 0.57					1.20	0.49	1.03		1.21	1.26			0.01

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SPO3786 SPO3787 SPO3788 SPO3789

SPO3790 SPO3791 SP03778 SP03779 SP03780 SP03781 SP03782 SP03783 SP03783 SP03785 SPO3775 SPO3776

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SPO3777

SPO3767 SPO3768 SPO3769 SPO3770 SPO3771 SPO3772

SPO3765 SPO3766

SPO3759 SPO3760 SPO3761 SPO3762 SPO3763 SPO3764 SPO3755 SPO3756 SPO3757 SPO3758

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Kirkwood
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# Chapter 10: Appendix

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dapB				pepT		sthA		rpsJ	rplC	rplD	rplW					folC	accD							gpmI				smpB	dmdD	dmdC			rpoN		acoX	acoA
dihydrodipicolinate reductase (EC:1.3.1.26)	DEAD/DEAH box helicase	ornithine cyclodeaminase/mu-crystallin family	alpha/beta hydrolase	peptidase T (EC:3.4.11.4)	S-formylglutathione hydrolase	зонноге ругилие пистеоние паньпулгоденазе (EC:1.6.1.1)	Yail/YqxD family protein	30S ribosomal protein S10	50S ribosomal protein L3	50S ribosomal protein L4	50S ribosomal protein L23	HAD-superfamily hydrolase	protein	major facilitator superfamily transporter	AFG1 family ATPase	synthase/dihydrofolate synthase	acetyl-CoA carboxylase, carboxyl transferase subunit beta (EC.64.1.2)	CAAX amino terminal protease	ABC transporter permease	hypothetical protein	hypothetical protein	carboxyl-terminal protease	hypothetical protein	phosphoglyceromutase (EC:5.4.2.1)	ArsR family transcriptional regulator	hypothetical protein	hypothetical protein	SsrA-binding protein	methylthioacryloyl-CoA hydratase (EC:4.2.1.17)	3-methylmercaptopropionyl-CoA dehydrogenase	AraC family transcriptional regulator	hypothetical protein	RNA polymerase sigma-54 factor	hypothetical protein	acetoin catabolism protein X	acetoin dehydrogenase complex, E1 component subunit alpha (EC:1.1.1)
-1.06	-0:20 X	86 0-	1.80	2.41	1.62	1.40	1.07	-1.39	-1.61	-1.35	1.07	-1.01	1.00	1.42	1.45	1.47	1.15	1.29	1.12	-1.41	0.98	-1.22	-1.28	-1.24	1.05	0.97	-1.89	-1.00	6.53	4.36	x	-1.33	1.36	-1.62	1.75	2.92
0.98	X .	0 70	0.32	0.01	0.00	0.53	0.98	0.01	0.01	0.01	0.82	0.44	0.75	0.67	0.12	0.03	0.86	0.02	0.72	0.01	0.11	0.54	0.37	0.83	1.00	0.80	0.28	0.76	0.00	0.09	×	0.62	0.18	0.14	0.02	0.02
1.40	1.02 1.62	1 02	-1.27	1.05	-1.08	1.17	1.07	-1.57	-1.44	-1.12	1.61	-1.11	-1.20	1.55	1.27	-1.33	-1.77	1.01	1.00	-1.13	1.14	-1.07	1.04	-1.18	-1.04	1.01	-1.03	-1.33	3.96	4.75	-1.02	-1.04	-1.02	-1.32	-1.78	-1.69
0.46	0.38	0 95	0.02	0.70	0.71	0.53	0.90	0.01	0.08	0.13	0.13	0.28	0.38	0.32	0.18	0.02	0.10	0.84	0.99	0.39	0.30	0.76	0.72	0.59	0.96	0.99	0.86	0.17	0.01	0.01	0.99	0.94	0.97	0.09	×	0.00
0.17	0.02 N/A	0.02	0.27	1.73	0.27	1.29	1.07	-1.48	-1.53	-1.24	1.34	-1.06	-0.10	1.49	1.36	0.07	-0.31	1.15	1.06	-1.27	1.06	-1.15	-0.12	-1.21	0.01	0.99	-1.46	-1.16	5.25	4.56	N/A	-1.19	0.17	-1.47	-0.02	0.62
1.23	N/A	1 00	1.54	0.68	1.35	0.12	0.00	0.09	0.09	0.12	0.27	0.05	1.10	0.07	0.09	1.40	1.46	0.14	0.06	0.14	0.08	0.08	1.16	0.03	1.05	0.02	0.43	0.17	1.29	0.20	N/A	0.15	1.19	0.15	1.77	2.31
2.63	7.51	1 37	2.22	2.57	2.64	2.40	1.27	9.22	9.32	9.01	14.10	1.97	1.40	3.97	2.14	3.83	2.29	2.00	1.57	-1.48	1.02	1.18	2.37	1.99	1.25	-1.33	-2.98	2.23	1.81	1.23	-1.18	-1.79	-1.43	-4.14	-4.43	-14.90
0.01	0.00	0 07	0.01	0.01	0.01	0.14	0.92	0.00	0.00	0.00	0.01	0.05	0.51	0.16	0.04	0.01	0.06	0.01	0.08	0.04	0.98	1.00	0.11	0.47	0.92	0.43	0.02	0.00	0.07	0.82	0.85	0.54	0.63	0.00	0.06	0.00
-1.57	-1.19	1 24	1.36	1.18	-1.15	-1.10	-1.04	-1.71	-1.33	-2.02	-2.10	-1.10	1.16	-1.01	-1.67	-1.23	1.18	-1.11	1.07	-1.09	-1.32	-1.08	-1.67	-1.51	1.10	-1.05	1.49	-1.20	1.74	1.44	1.05	1.19	1.15	-1.26	×	2.12
0.02	0.17	0 14	0.04	0.41	0.09	0.60	0.95	0.00	0.02	0.01	0.02	0.52	0.06	0.91	0.05	0.13	0.39	0.59	0.31	0.17	0.07	0.62	0.00	0.20	0.89	0.98	0.10	0.28	0.03	0.15	0.64	0.76	0.65	0.57	×	0.02
0.53	3.16	1 31	1.79	1.88	0.75	0.65	0.12	3.76	4.00	3.50	6.00	0.44	1.28	1.48	0.24	1.30	1.74	0.45	1.32	-1.29	-0.15	0.05	0.35	0.24	1.18	-1.19	-0.75	0.52	1.78	1.34	-0.06	-0.30	-0.14	-2.70	N/A	-6.39
2.10	4.35	0 07	0.43	0.70	1.90	1.75	1.16	5.47	5.33	5.52	8.10	1.54	0.12	2.49	1.91	2.53	0.56	1.56	0.25	0.19	1.17	1.13	2.02	1.75	0.08	0.14	2.24	1.72	0.04	0.11	1.12	1.49	1.29	1.44	N/A	8.51
1.30	-1.93	-1 02	-1.22	1.25	-1.41	1.02	1.08	-1.44	-1.43	-1.04	-1.11	-1.10	-1.13	-1.01	-1.12	-1.50	-1.27	1.18	-1.07	-1.32	1.26	1.27	-1.17	-1.61	1.40	1.35	1.41	-1.19	-1.08	1.22	1.08	1.34	-1.00	-1.43	-1.83	-1.78
0.14	0.02	0.62	0.06	0.11	0.07	0.93	0.91	0.01	0.10	0.57	0.09	0.53	0.55	0.93	0.08	0.07	0.06	0.21	0.65	0.05	0.03	0.33	0.25	0.13	0.63	0.79	0.01	0.29	0.40	0.18	0.67	0.53	0.99	0.49	×	0.05
-1.24	-2.30	1 23	-1.14	1.32	-1.58	-1.24	1.22	-2.58	-2.39	-2.29	-3.29	-1.18	-1.22	-1.11	-1.91	-1.99	1.03	-0.94	1.11	-1.40	-1.45	1.54	-1.07	-1.28	1.92	1.53	1.40	-1.30	1.32	1.35	-1.27	1.19	-1.28	2.26	×	×
0.53	0.04	0 79	0.23	0.09	0.07	0.33	0.84	0.01	0.00	0.01	0.00	0.16	0.24	0.84	0.02	0.00	0.03	0.45	0.32	0.02	0.08	0.07	0.54	0.31	0.40	0.87	0.14	0.00	0.39	0.13	0.42	0.60	0.70	0.03	×	x
0.03	-2.12	0 1 1	-1.18	1.29	-1.50	-0.11	1.15	-2.01	-1.91	-1.67	-2.20	-1.14	-1.18	-1.06	-1.52	-1.75	-0.12	0.12	0.02	-1.36	-0.10	1.41	-1.12	-1.45	1.66	1.44	1.41	-1.25	0.12	1.29	-0.10	1.27	-1.14	0.42	N/A	N/A
1.27	0.19	1 13	0.04	0.04	0.09	1.13	0.07	0.57	0.48	0.63	1.09	0.04	0.05	0.05	0.39	0.25	1.15	1.06	1.09	0.04	1.36	0.14	0.05	0.17	0.26	0.09	0.01	0.06	1.20	0.07	1.18	0.08	0.14	1.85	N/A	N/A

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SPO3832 SPO3833 SPO3834 SPO3828 SPO3829 SPO3830 SPO3831 SP03821 SP03822 SP03823 SP03824 SP03824 SP03825 SP03826 SP03827 SPO3818 SPO3819 SPO3820 SPO3817

SPO3809 SPO3810 SPO3811 SPO3812 SPO3813 SPO3813 SPO3814 SPO3815 SPO3806 SPO3807

SPO3808

SPO3795 SPO3802 SPO3803 SPO3804 SPO3805

SPO3792 SPO3793 SPO3794

SPO3872	SPO3871	SPO3870	SPO3869	SPO3868	SPO3867	SPO3866	SPO3865	SPO3864	SPO3863	SPO3862	SPO3861	SPO3860	SPO3859	SPO3858	SPO3857	SPO3856	SPO3855	SPO3854	SPO3853	SPO3852	SPO3851	SPO3850	SPO3849	SPO3848	SPO3847	SPO3846	SPO3845	SPO3844	SPO3843	SPO3842	SPO3841	SPO3840	SPO3839	SPO3838	SPO3837	SPO3836	SPO3835
					regB	senC	regA				ahcY						gpsA											polA				rpsO			truB		rbfA
hypothetical protein	nucleotidyltransferase	hypothetical protein	hypothetical protein	hypothetical protein	sensor histidine kinase RegB (EC:2.7.3)	regulatory protein SenC	photosynthetic apparatus regulatory protein RegA	hypothetical protein	HD domain-containing protein	lipoprotein	S-adenosyl-L-homocysteine hydrolase (EC:3.3.1.1)	hypothetical protein	NAD(P)H-dependent glycerol-3-phosphate dehydrogenase (EC:1.1.1.94)	DNA-binding/iron metalloprotein/AP endonuclease	uroporphyrinogen-III synthase	hypothetical protein	HemY domain-containing protein	giuannone-rependent tormadenyde dehydrogenase (EC:1.1.284)	hypothetical protein	sterol desaturase	ABC transporter ATP-binding protein	hypothetical protein	hypothetical protein	DNA polymerase I (EC:2.7.7.7)	malonate transporter	hypothetical protein	TetR family transcriptional regulator	30S ribosomal protein S15	type I secretion target repeat-containing protein	hypothetical protein	tRNA pseudouridine synthase B (EC:4.2.1.70)	hypothetical protein	ribosome-binding factor A				
1.86	1.92	1.13	1.12	1.38	-1.03	3.95	-1.19	-1.66	-1.35	-3.76	1.17	1.11	0.98	1.42	1.33	1.59	1.23	-1.40	1.04	2.17	2.35	3.57	-0.94	-1.22	-1.13	1.98	1.64	1.07	1.28	-1.17	-1.42	-1.17	1.02	-1.12	-1.38	-1.11	-1.23
0.00	0.05	0.41	0.71	0.53	0.83	0.00	0.04	0.00	0.34	0.29	0.91	0.66	0.39	0.02	0.14	0.80	0.72	0.02	0.79	0.13	0.05	0.17	0.35	0.15	0.12	0.03	0.05	0.92	0.25	0.05	0.00	0.70	0.85	0.88	0.62	0.35	0.06
1.10	1.05	-1.01	-1.11	2.43	-1.16	1.88	-1.03	1.02	-1.09	-1.38	1.15	1.18	-1.38	0.99	1.38	1.10	-1.14	1.06	-1.18	2.14	1.93	1.18	1.17	-1.05	1.13	1.16	1.57	1.33	1.29	1.66	-1.20	1.18	-1.06	-1.15	1.27	-1.15	-1.17
0.30	0.86	0.95	0.43	0.01	0.66	0.04	0.57	0.99	0.72	0.81	0.80	0.07	0.25	0.80	0.43	0.47	0.29	0.90	0.04	0.07	0.00	0.83	0.10	0.61	0.64	0.11	0.02	0.44	0.11	0.09	0.40	0.09	0.71	0.21	0.06	0.24	0.56
1.48	1.49	0.06	0.01	1.91	-1.10	2.92	-1.11	-0.32	-1.22	-2.57	1.16	1.15	-0.20	1.21	1.36	1.35	0.05	-0.17	-0.07	2.16	2.14	2.38	0.12	-1.14	0.00	1.57	1.61	1.20	1.29	0.25	-1.31	0.01	-0.02	-1.14	-0.05	-1.13	-1.20
0.38	0.44	1.07	1.12	0.53	0.06	1.04	0.08	1.34	0.13	1.19	0.01	0.03	1.18	0.22	0.02	0.24	1.19	1.23	1.11	0.01	0.21	1.20	1.06	0.09	1.13	0.41	0.03	0.13	0.01	1.42	0.11	1.18	1.04	0.01	1.33	0.02	0.03
2.66	2.86	1.83	1.65	-4.87	1.39	5.19	1.08	0.89	2.01	-5.05	4.34	-3.17	-1.39	0.84	1.29	1.56	2.02	1.56	1.28	2.23	1.96	1.03	-1.12	-1.97	1.34	2.02	3.01	1.52	1.79	-1.45	-1.65	11.40	-2.56	2.81	1.87	1.79	1.66
0.00	0.02	0.05	0.10	0.02	0.06	0.00	0.24	0.27	0.52	0.16	0.03	0.00	0.08	0.07	0.16	0.75	0.01	0.00	0.08	0.05	0.07	0.92	0.45	0.02	0.01	0.03	0.01	0.40	0.06	0.07	0.01	0.03	0.02	0.15	0.18	0.02	0.01
-1.01	1.18	-1.06	1.34	2.73	-1.41	0.99	1.18	1.01	-1.12	1.08	-1.08	1.92	1.47	1.18	-1.18	1.13	0.97	-1.60	1.09	1.32	1.68	1.38	1.45	1.44	-1.42	0.98	-1.30	-1.19	1.48	1.28	1.31	-2.44	-1.13	-1.28	-1.80	-1.19	-1.14
0.52	0.24	0.50	0.28	0.01	0.04	0.82	0.23	0.83	0.27	0.94	0.71	0.01	0.07	0.24	0.26	0.87	0.46	0.07	0.56	0.16	0.03	0.65	0.19	0.06	0.05	0.32	0.03	0.36	0.03	0.05	0.23	0.01	0.36	0.15	0.15	0.25	0.04
0.83	2.02	0.39	1.50	-1.07	-0.01	3.09	1.13	0.95	0.45	-1.99	1.63	-0.63	0.04	1.01	0.06	1.35	1.49	-0.02	1.19	1.78	1.82	1.21	0.17	-0.27	-0.04	1.50	0.86	0.17	1.64	-0.09	-0.17	4.48	-1.85	0.77	0.04	0.30	0.26
1.84	0.84	1.45	0.16	3.80	1.40	2.10	0.05	0.06	1.57	3.07	2.71	2.55	1.43	0.17	1.24	0.22	0.53	1.58	0.10	0.46	0.14	0.17	1.29	1.71	1.38	0.52	2.16	1.36	0.16	1.37	1.48	6.92	0.72	2.05	1.84	1.49	1.40
1.02	1.16	-0.99	-1.18	-0.98	-1.11	-1.22	1.09	1.53	-1.10	-1.13	-1.18	1.18	1.05	1.59	1.46	1.78	-1.34	1.16	1.05	1.63	1.11	1.20	-1.02	-1.06	-1.26	1.03	-1.10	1.07	1.23	1.26	1.60	-1.20	-1.02	-1.21	-1.07	1.41	1.42
0.68	0.39	0.86	0.46	0.93	0.52	0.05	0.80	0.01	0.53	0.88	0.68	0.20	0.94	0.06	0.15	0.29	0.20	0.52	0.43	0.01	0.40	0.79	0.91	0.45	0.04	0.88	0.03	0.82	0.11	0.02	0.05	0.34	0.85	0.21	0.74		0.01
-0.99	-1.20	-1.15	-1.11	1.46	-1.46	-1.41	1.02	1.36	-1.17	1.45	-1.31	1.42	-1.11	1.99	-1.17	1.55	-1.37	-1.31	-1.16	1.76	1.66	1.30	x	1.23	-1.58	-1.17	-1.68	-1.34	-1.39	1.27	1.91	-1.73	1.44	-1.56	-1.28	-1.06	1.20
0.68	0.60	0.51	0.46	0.07	0.12	0.04	0.36	0.16	0.40	0.76	0.54	0.30	0.16	0.01	0.40	0.56	0.11	0.48	0.48	0.07	0.04	0.74	Х	0.13	0.03	0.19	0.03	-		0.08	0.05	0.01	0.07	0.07	0.48	0.58	0.51
0.02	-0.02	-1.07	-1.15	0.24	-1.29	-1.32	1.06	1.45	-1.14	0.16	-1.25	1.30	-0.03	1.79	0.15	1.67	-1.36	-0.08	-0.05	1.70	1.39	1.25	N/A	0.09	-1.42	-0.07	-1.39	-0.14	-0.08	1.27	1.76	-1.47	0.21	-1.39	-1.18	0.18	1.31
1.00	1.18	0.08	0.03	1.22		0.10	0.04	0.09	0.03	1.29	0.07	0.12	1.08	0.20	1.32	0.12	0.02	1.24	1.11	0.07	0.28	0.05	N/A	1.15	0.16	1.10	0.29	1.21	1.31	0.01	0.16	0.27	1.23	0.18	0.11		0.11

SPOA0006	SPOA0005	SPOA0004	SPOA0003	SPOA0002	SPOA0001	SPO3903	SPO3902	SPO3901	SPO3900	SPO3899	SPO3898	SPO3897	SPO3896	SPO3895	SPO3894	SPO3893	SPO3892	SPO3891	SPO3890	SPO3889	SPO3888	SPO3887	SPO3886	SPO3885	SPO3884	SPO3883	SPO3882	SPO3881	SPO3880	SPO3879	SPO3878	SPO3877	SPO3876	SPO3875	SPO3874	SPO3873
				miaB										trmE	rho		maf-1	aroE	coaE	dmaQ	secB	fxsA					hslU		hslV		gap-4				trx	uvrD
PhoH family protein	lipoprotein	OmpA family protein	hypothetical protein	(unieulyianyi)aucilosine uxiva methylthiotransferase	Aimstructure of the second sec	6-pyruvoyl tetrahydrobiopterin synthase	transglycosylase domain-containing protein	carbon monoxide dehydrogenase G protein	thermonuclease	hypothetical protein	DNA-binding protein	resolvase site-specific recombinase	TetR family transcriptional regulator	tRNA modification GTPase TrmE	transcription termination factor Rho	hypothetical protein	hypothetical protein	shikimate 5-dehydrogenase (EC:1.1.1.25)	dephospho-CoA kinase (EC:2.7.1.24)	DNA polymerase III subunit epsilon (EC:2.7.7.7)	preprotein translocase subunit SecB	FxsA protein	Tim44 family protein	MltA/3D domain-containing protein	smr domain-containing protein	lipoprotein	A 17-dependent protease A 17-binding subunit HsIU	twin-arginine translocation pathway signal sequence domain-containing protein	A 17-dependent protease peptidase subunit (EC:3.4.25.1)	A TD dependent motoric postidance subunit	уусстаниспунс-э-риозриате испунгоденизе, турс 1 (ЕС:1.2.1)	transcriptional regulator/arsenate reductase	hypothetical protein	hypothetical protein	thioredoxin	ATP-dependent DNA helicase UvrD
2.95	2.34	2.26	2.07	1.05	-1.20	-1.71	-1.48	1.63	1.65	1.48	10.80	-1.08	1.16	-1.21	-1.24	1.22	-1.42	1.27	1.37	-1.22	1.57	1.06	1.19	-1.32	-1.19	1.10	1.08	1.63	1.36	1.08	1.44	x	1.16	1.03	1.79	-1.60
0.14	0.01	0.01	0.05	Х	0.10	×	0.14	0.89	0.07	0.13	0.01	0.87	0.89	0.81	0.56	0.35	0.16	0.74	0.01	0.35	0.01	0.91	0.94	0.77	0.47	0.51	0.54	0.01	0.05	0.90	×	×	0.61	0.93	0.08	0.16
1.49	2.04	1.41	1.30	1.28	1.04	1.26	-1.25	1.31	1.40	-1.20	-1.19	-1.38	-1.49	1.31	1.21	1.08	1.20	1.30	1.17	1.06	-1.53	-1.68	-1.05	-1.19	-1.16	-1.12	-1.51	-1.27	1.19	-1.22	1.28	х	-1.02	-1.07	1.27	-1.32
0.15	0.04	0.08	0.06	0.04	0.88	0.10	0.47	0.42	0.22	0.41	0.03	0.31	0.17	0.13	0.48	0.76	0.38	0.05	0.33	0.64	0.10	0.03	0.66	0.64	0.03	0.65	0.15	0.40	0.63	0.64	0.24	×	0.92	0.66	0.35	0.49
2.22	2.19	1.84	1.69	1.17	-0.08	-0.23	-1.37	1.47	1.53	0.14	4.81	-1.23	-0.17	0.05	-0.02	1.15	-0.11	1.29	1.27	-0.08	0.02	-0.31	0.07	-1.26	-1.18	-0.01	-0.22	0.18	1.28	-0.07	1.36	N/A	0.07	-0.02	1.53	-1.46
0.73	0.15	0.43	0.38	0.12	1.12	1.49	0.12	0.16	0.13	1.34	6.00	0.15	1.33	1.26	1.23	0.07	1.31	0.02	0.10	1.14	1.55	1.37	1.12	0.07	0.02	1.11	1.30	1.45	0.09	1.15	0.08	N/A	1.09	1.05	0.26	0.14
0.93	-2.28	-2.12	-1.83	4.06	1.27	-1.82	-1.82	2.28	-1.42	1.51	1.00	-0.97	0.95	-1.51	6.05	2.73	-0.99	2.06	1.83	1.00	2.94	1.50	2.08	1.89	1.54	3.25	5.54	-1.09	3.12	0.84	1.62	х	1.33	1.30	2.07	-1.96
0.29	0.00	0.01	0.03	0.01	0.05	0.39	0.03	0.77	0.12	0.05	0.98	0.22	0.65	0.12	0.02	0.00	0.52	0.01	0.00	0.91	0.00	0.26	0.05	0.67	0.03	0.02	0.01	0.75	0.01	0.29	0.01	х	0.21	0.08	0.01	0.04
1.60	1.53	1.76	1.53	-1.09	1.38	-5.41	1.61	-1.39	1.07	1.14	-1.13	-1.47	1.11	-1.10	-2.11	-1.23	-1.30	-1.30	1.06	-1.23	-1.20	-1.32	-1.10	-1.10	-1.09	-1.06	-1.26	1.20	-1.17	-1.04	1.21	×	1.09	1.12	-1.09	1.16
0.17	0.02	0.06	0.06	0.38	0.02	0.00	0.10	0.63	0.83	0.25	0.09	0.36	0.32	0.22	0.00	0.10	0.25	0.17	0.71	0.16	0.05	0.43	0.24	0.75	0.49	0.58	0.01	0.08	0.41	0.92	0.56	×	0.56	0.58	0.38	0.55
1.27	-0.38	-0.18	-0.15	1.49	1.33	-3.62	-0.11	0.45	-0.18	1.33	-0.06	-1.22	1.03	-1.31	1.97	0.75	-1.15	0.38	1.45	-0.12	0.87	0.09	0.49	0.40	0.23	1.10	2.14	0.05	0.98	-0.10	1.42	N/A	1.21	1.21	0.49	-0.40
0.33	1.91	1.94	1.68	2.58	0.05	1.80	1.72	1.84	1.25	0.19	1.07	0.25	0.08	0.20	4.08	1.98	0.16	1.68	0.39	1.11	2.07	1.41	1.59	1.50	1.32	2.16	3.40	1.15	2.15	0.94	0.21	N/A	0.12	0.09	1.58	1.56
2.08	1.11	1.48	1.31	-1.40	-1.31	-4.41	-1.06	1.10	1.14	-1.55	-1.06	-1.32	-1.22	-1.19	1.22	1.35	-1.04	1.28	-1.35	1.06	0.97	1.53	1.65	1.22	-1.02	1.28	1.14	-1.55	1.07	-1.33	-1.01	1.16	-0.99	-1.19	1.40	-1.15
0.06	0.89	0.08	0.25	0.06	0.12	0.01	0.76	0.77	0.58	0.10	0.73	0.62	0.28	0.33	0.05	0.06	0.64	0.21	0.05	0.79	0.61	0.16	0.06	0.48	0.87	0.44	0.27	0.02	0.88	0.48	0.88	0.40	0.92			0.28
2.38	-1.06	1.85	1.22	-1.73	-1.25	-3.61	1.10	-1.09	-1.23	-1.70	-1.22	1.27	-1.23	-1.66	-1.42	1.59	-1.06	-1.25	-1.81	-1.08	1.06	1.84	2.28	1.16	1.14	1.15	1.15	-1.91	-1.56	1.37	-1.23	Х	-0.98	-1.60	1.35	1.87
0.06	0.55	0.03	0.19	0.02	0.13	0.01	0.44	0.97	0.52	0.10	0.13	0.53	0.25	0.02	0.16	0.34	0.85	0.13		0.86	0.56	0.00	0.01	0.80	0.38	0.37	0.63	0.01	0.32	0.43	0.53	x				0.38
5 2.23	0.03	3 1.67	1.27	-1.57	-1.28	-4.01						-0.03	-1.23		-0.10	1 1.47	-1.05	3 0.02	4 -1.58	-0.01	1.01	) 1.69	1.97	) 1.19	3 0.06	1.22	3 1.15	-1.73	-0.25	3 0.02		N/A				3 0.36
	1.09	0.18	0.05	0.17	0.03	0.40	1.08	1.10	1.19	0.08	4 0.08		0.01	0.24	1.32		0.01		0.23	1.07	0.05	0.16	0.32	0.03	1.08	0.07	0.01	0.18	1.32	1.35	0.11	N/A	3 0.01			1.51

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SPOA0042	SPOA0041	SPOA0040	SPOA0039	SPOA0038	SPOA0037	SPOA0036	SPOA0035	SPOA0034	SPOA0033	SPOA0032	SPOA0031	SPOA0030	SPOA0029	SPOA0028	SPOA0027	SPOA0026	SPOA0025	SPOA0024	SPOA0023	SPOA0022	SPOA0021	SPOA0020	SPOA0019	SPOA0018	SPOA0017	SPOA0015	SPOA0014	SPOA0013	SPOA0012	SPOA0011	SPOA0010	SPOA0009	SPOA0008	SPOA0007
						apbE			nqrF	nqrE	nqrD	nqrC	nqrB	nqrA	hpcC	hpcB									cmk	aroA	trmB			metK	cutE			
hypothetical protein	crcB family protein	crcB family protein	hypothetical protein	LuxR family transcriptional regulator	hypothetical protein	thiamin biosynthesis lipoprotein ApbE	hypothetical protein	CBS domain-containing protein	Na(+)-translocaung NADH-quinone reductase subunit F (EC:1.6.5)	Na(+)-translocating NADH-quinone reductase subunit E (EC:1.6.5)	subunit D (EC:1.6.5)	subunit C (EC:1.6.5) Na(+)-translocating NADH-quinone reductase	subunit B (EC:1.6.5) Na(+)-translocating NADH-quinone reductase	subunit A (EC:1.6.5) Na(+)-translocating NADH-quinone reductase	dehydrogenase (EC:1.2.1.60) Na(+)-translocating NADH-quinone reductase	(EC:1.13.11.15)	fumarylacetoacetate hydrolase family protein 3,4-dihydroxyphenylacetate 2,3-dioxygenase	2-oxo-hepta-3-ene-1,7-dioic acid hydratase	TetR family transcriptional regulator	cytochrome c5	oxidoreductase, FAD-binding	alanine racemase	hypothetical protein	hypothetical protein	cytidylate kinase (EC:2.7.4.14)	3-phosphoshikimate 1-carboxyvinyltransferase (EC:2.5.1.19)	(EC:2.1.1.33)	hypothetical protein	DNA topology modulation kinase FlaR	S-adenosylmethionine synthetase (EC:2.5.1.6)	apolipoprotein N-acyltransferase (EC:2.3.1)	hemolysin	hypothetical protein	hypothetical protein
1.21	х	х	0.98	1.46	1.54	1.18	-1.05	1.50	1.31	1.27	1.51	-1.21	1.27	1.37	1.57	2.23	2.65	2.85	-1.48	-1.25	-1.12	-1.83	1.26	-1.28	1.52	1.69	-1.88	-1.34	-1.81	1.25	-1.16	-2.04	1.58	1.46
0.95	×	×	0.09	0.71	0.61	0.13	0.98	0.00	0.53	0.76	0.63	0.41	0.34	0.62	0.04	0.33	0.00	0.24	0.14	0.25	0.58	0.14	0.38	0.68	0.07	0.02	0.33	0.16	0.04	0.26	0.72	0.00	0.71	0.32
1.49	1.18	1.28	1.09	1.14	1.63	1.37	1.31	1.38	1.57	1.44	1.20	1.46	1.24	1.82	1.10	2.22	1.67	1.80	-1.15	-1.16	-1.09	-1.34	-1.02	-1.01	1.63	1.27	1.01	1.21	1.20	-1.22	-1.10	1.43	1.16	1.62
0.64	0.16	0.15	0.45	0.82	0.04	0.14	0.32	0.29	0.29	0.33	0.65	0.02	0.32	0.29	0.63	0.01	0.05	0.12	0.26	0.22	0.74	0.21	0.92	0.98	0.17	0.10	0.88	0.25	0.46	0.35	0.46	0.16	0.19	0.21
1.35	N/A	N/A	1.03	1.30	1.59	1.28	0.13	1.44	1.44	1.36	1.36	0.13	1.26	1.60	1.34	2.23	2.16	2.33	-1.32	-1.21	-1.11	-1.59	0.12	-1.15	1.58	1.48	-0.44	-0.07	-0.31	0.02	-1.13	-0.31	1.37	1.54
0.14	N/A	N/A	0.06	0.16	0.04	0.10	1.18	0.06	0.13	0.09	0.16	1.34	0.02	0.22	0.24	0.00	0.49	0.53	0.17	0.05	0.02	0.25	1.14	0.14	0.05	0.21	1.45	1.28	1.51	1.24	0.03	1.74	0.21	0.08
-1.49	2.73	1.80	0.93	-2.51	2.54	5.37	1.80	-1.83	4.11	2.82	5.86	1.29	4.67	6.09	-1.78	1.16	1.29	1.50	-2.47	-2.73	-1.42	-3.80	-1.41	-1.59	3.87	4.03	1.95	-1.23	1.08	5.46	2.01	1.15	-1.08	0.91
0.86	0.01	0.00	0.09	0.30	0.04	0.00	0.02	0.00	0.10	0.18	0.09	0.32	0.03	0.11	0.04	0.90	0.22	0.10	0.07	0.03	0.43	0.00	0.30	0.61	0.01	0.00	0.10	0.25	0.66	0.00	0.08	0.09	0.74	0.62
1.32	-1.16	-0.98	1.72	1.31	-1.26	-1.33	-1.43	1.93	-1.34	-1.20	-1.41	-1.53	-1.50	-1.61	-1.20	1.42	1.47	-1.08	1.90	1.97	1.73	1.34	1.83	1.92	-1.45	-1.28	-1.17	1.33	-1.12	1.21	-1.34	-1.14	1.49	1.40
0.78	0.25	0.80	0.04	0.71	0.25	0.03	0.06	0.01	0.49	0.24	0.04	0.05	0.06	0.35	0.14	0.03	0.01	0.51	0.09	0.02	0.15	0.16	0.03	0.19	0.03	0.07	0.28	0.26	0.69	0.21	0.07	0.15	0.03	0.12
-0.09	0.79	0.41	1.33	-0.60	0.64	2.02	0.19	0.05	1.39	0.81	2.23	-0.12	1.59	2.24	-1.49	1.29	1.38	0.21	-0.29	-0.38	0.16	-1.23	0.21	0.17	1.21	1.38	0.39	0.05	-0.02	3.34	0.34	0.01	0.21	1.16
1.41	1.95	1.39	0.39	1.91	1.90	3.35	1.62	1.88	2.73	2.01	3.64	1.41	3.09	3.85	0.29	0.13	0.09	1.29	2.19	2.35	1.58	2.57	1.62	1.76	2.66	2.66	1.56	1.28	1.10	2.13	1.68	1.15	1.29	0.24
-1.07	-1.73	-1.26	1.52	-1.04	2.13	1.04	1.20	1.25	0.99	1.06	-1.12	-1.13	-1.15	1.10	-1.43	1.10	1.00	-1.02	-1.43	-1.25	-1.15	-1.92	-1.13	-1.08	1.15	-1.21	-1.25	1.60	-1.17	-1.20	1.11	2.03	-1.30	1.22
0.93	0.02	0.50	0.05	0.94	0.06	0.93	0.09	0.44	0.95	0.95	0.29	0.29	0.11	0.88	0.07	0.66	0.86	0.76	0.18	0.03	0.59	0.15	0.31	0.80	0.13	0.08	0.30	0.03	0.37	0.17	0.41	0.01	0.07	0.08
-1.32	-2.51	-1.40	1.91	1.64	2.38	-1.16	-1.56	1.56	-1.27	0.94	-1.37	-1.41	-1.22	-1.33	-1.35	1.17	-1.10	-1.25	-1.91	-1.35	-1.61	1.38	-1.54	1.30	-1.47	-1.41	1.06	1.19	-1.20	1.10	-1.13	1.58	-1.43	1.26
0.75	0.02	x	0.05	0.41	0.08	0.27	0.03	0.04	0.63	0.38	0.09	0.12	0.06	0.54	0.14	0.54	0.33	0.31	0.02	0.02	0.04	0.22	0.07	0.56	0.00	0.05	0.96	0.20	0.91	0.88	0.36	0.05	0.01	0.69
-1.20	-2.12	-1.33	1.72	0.30	2.26	-0.06	-0.18	1.41	-0.14	1.00	-1.25	-1.27	-1.19	-0.12	-1.39	1.14	-0.05	-1.14	-1.67	-1.30	-1.38	-0.27	-1.34	0.11	-0.16	-1.31	-0.10	1.40	-1.19	-0.05	-0.01	1.81	-1.37	1.24
0.13	0.39	0.07	0.20	1.34	0.13	1.10	1.38	0.16	1.13	0.06	0.13	0.14	0.04	1.22	0.04	0.03	1.05	0.12	0.24	0.05	0.23	1.65	0.20	1.19	1.31	0.10	1.16	0.21	0.02	1.15	1.12	0.22	0.06	0.02

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SPOA0078	SPOA0077	SPOA0076	SPOA0075	SPOA0074	SPOA0073	SPOA0072	SPOA0071	SPOA0070	SPOA0069	SPOA0068	SPOA0067	SPOA0066	SPOA0065	SPOA0064	SPOA0063	SPOA0062	SPOA0061	SPOA0060	SPOA0059	SPOA0058	SPOA0057	SPOA0056	SPOA0055	SPOA0054	SPOA0053	SPOA0052	SPOA0051	SPOA0050	SPOA0049	SPOA0048	SPOA0047	SPOA0046	SPOA0045	SPOA0044	SPOA0043
maf-2	infA				arsC-2												katG		$g_{CV}P$	gcvH-2	gcvT			nosL	nosY	nosF	nosD	nosZ	nosR		pcaQ	pobA	pcaC	pcaH	pcaG
maf protein	translation initiation factor IF-1	carbon-nitrogen family hydrolase	hypothetical protein	hypothetical protein	arsenate reductase (EC:1.20.4.1)	hypothetical protein	polar amino acid ABC transporter permease	polar amino acid ABC transporter permease	polar amino acid ABC transporter periplasmic substrate-binding protein	protein	TetR family transcriptional regulator nolar amino acid ARC transporter ATP-binding	aspartate aminotransferase	hypothetical protein	NG,NG-dimethylarginine dimethylaminohydrolase	FAD-dependent oxidoreductase	aspartate racemase (EC:5.1.1.13)	catalase/peroxidase HPI (EC:1.11.1.6)	AraC family transcriptional regulator	glycine dehydrogenase (EC:1.4.4.2)	glycine cleavage system protein H	glycine cleavage system T protein (EC:2.1.2.10)	DNA-binding protein	cytochrome c family protein	nitrous oxide reductase accessory protein	nitrous oxide maturation protein NosY	ABC transporter, ATP-binding protein NosF	nitrous oxide maturation protein NosD	nitrous-oxide reductase (EC:1.7.99.6)	nitrous-oxide reductase transcriptional activator NosR	hypothetical protein	pca operon transcriptional activator PcaQ	тиуциолу оснасовые 5-тионоголу усназе (EC:1.14.13.2)	EC:4.1.1.44)	protocatecritate 3,4-citoXygenase subuitit beta (EC:1.13.11.3) 4-carboxymiconolactone decarboxylase	protocatechuate 3,4-dioxygenase alpha subunit (EC:1.13.11.3) reconstructure 3.4 dioxygenases subunit beta
-1.21	1.13	2.59	3.07	2.40	3.65	1.12	-1.42	х	-5.54	-2.79	-2.63	-2.00	-1.95	-1.16	1.10	1.18	-1.67	1.60	1.19	-1.43	-1.61	-1.15	1.26	×	1.84	-1.08	x	-1.33	1.00	-2.40	х	-1.48	-1.24	-1.25	1.34
0.81	0.56	0.02	0.01	0.00	0.40	0.65	0.08	x	0.08	0.00	0.02	0.02	0.08	0.37	0.72	0.45	0.01	0.75	0.62	0.01	0.23	0.03	0.28	×	0.12	0.72	×	×	х	0.09	x	0.71	0.52	0.22	0.71
-1.36	-1.40	1.18	1.08	1.21	1.05	-1.35	-1.05	-1.32	-1.89	-1.46	-1.20	1.18	-1.30	-1.24	-1.15	-1.12	0.99	-1.78	-2.26	-5.32	-5.58	-2.89	-1.21	1.09	-1.11	-1.34	х	-1.30	-0.99	-1.38	-1.01	-1.25	-1.14	-1.12	-1.18
0.01	0.03	0.07	0.85	0.37	0.88	0.09	0.96	0.27	0.07	0.08	0.57	0.17	0.10	0.57	0.38	0.18	0.65	0.07	0.01	0.00	0.01	0.00	0.10	0.39	0.59	0.48	×	0.74	0.83	0.44	0.98	0.50	0.68	0.26	0.23
-1.29	-0.14	1.89	2.08	1.81	2.35	-0.12	-1.24	N/A	-3.72	-2.13	-1.92	-0.41	-1.63	-1.20	-0.02	0.03	-0.34	-0.09	-0.54	-3.38	-3.60	-2.02	0.03	N/A	0.37	-1.21	N/A	-1.32	0.00	-1.89	N/A	-1.37	-1.19	-1.19	0.08
0.08	1.27	0.71	0.99	0.60	1.30	1.24	0.19	N/A	1.83	0.67	0.72	1.59	0.33	0.04	1.13	1.15	1.33	1.69	1.73	1.95	1.99	0.87	1.24	N/A	1.48	0.13	N/A	0.02	1.00	0.51	N/A	0.12	0.05	0.06	1.26
1.20	2.74	2.01	1.29	1.00	2.10	2.86	1.23	-1.65	-6.93	-4.53	-3.00	-1.65	-2.06	-1.45	-2.04	1.35	1.38	2.49	1.49	1.27	1.44	-4.06	-1.47	-0.95	2.15	-2.05	х	-1.15	1.45	3.62	1.54	-2.84	-1.83	-2.83	-2.24
0.17	0.00	0.01	0.28	0.27	0.08	0.02	0.30	х	0.01	0.01	0.03	0.04	0.12	0.41	0.05	0.37	0.45	0.55	0.07	0.12	0.54	0.00	0.13	0.24	0.05	0.03	x	0.99	0.81	0.01	0.02	0.04	0.52	0.01	0.04
1.01	0.96	-1.05	1.11	1.06	1.04	1.47	x	-1.88	-1.43	-1.33	-1.08	-1.38	-1.42	-1.20	1.42	-1.06	3.81	1.34	1.23	0.98	1.32	-1.40	1.23	x	-1.15	-1.44	x	-1.48	Х	1.11	-1.62	-1.44	1.47	1.27	1.60
0.71	0.28	0.84	0.77	0.96	0.93	0.04	x	X	0.10	0.20	0.91	0.13	0.02	0.50	0.02	0.83	0.00	0.05	0.31	0.33	0.28	0.12	0.20	x	0.61	0.16	x	0.84	х	0.82	0.01	0.08	0.03	0.13	0.02
1.11	1.85	0.48	1.20	1.03	1.57	2.17	N/A	-1.77	-4.18	-2.93	-2.04	-1.52	-1.74	-1.33	-0.31	0.15	2.60	1.92	1.36	1.12	1.38	-2.73	-0.12	N/A	0.50	-1.75	N/A	-1.32	N/A	2.37	-0.04	-2.14	-0.18	-0.78	-0.32
0.10	0.89	1.53	0.09	0.03	0.53	0.70	N/A	0.12	3 2.75	3 1.60	0.96	0.14	0.32	0.13	1.73	1.21	1.22	0.58	0.13	0.15	0.06	1.33	1.35	N/A	1.65	0.31	N/A	0.17	N/A	1.26	1.58	0.70	3 1.65	3 2.05	1.92
0 -1.3	9 -1.15	-1.06	9 -1.07	3 -1.10	3 -1.46			-1.03		0 -1.49	-1.49	4 1.22	2 -1.69	-1.32	3 -1.18	-1.00	2 1.24	-1.26	3 1.16	-2.63		-1.15		-1.03	-1.04	-1.03	×	-1.04	-1.15	6 -0.99	8 1.17	0 -1.19	-1.01	-1.13	-1.45
0.06	5 0.12	0.86	0.50	0 0.15	.6 0.08	2 0.02	<b>8</b> 0.54	0.83	0.77	9 0.04	9 0.16	2 0.19	9 0.03	0.38	8 0.17	0 0.96	4 0.09	0.17	6 0.11	3 0.06	8 0.09	5 0.34	.3 0.09	0.90	0.98	0.98	x	4 0.99	5 X	9 0.92	7 0.19	9 0.37	0.88	3 0.07	.5 0.05
6 -1.4	2 -1.40	-1.21	0 -1.06		8 -1.80	2 -1.83	4 X	3 -1.34	-1.08	4 -1.46	6 1.40	9 -1.28	3 -2.69	8 8	7 -1.14	6 1.19	9 2.92	7 -1.39	1 1.13	6 -2.77	9 -1.49	4 1.06	9 -1.21	0 X		8 1.88	x	9 1.20	Х	2 1.42	9 1.29	7 -1.43	8 1.02	-1.35	-1.07
43 0.12	40 0.03	21 0.32	0.72	0.62	30 0.01	33 <u>0.00</u>	×	34 X	0.55	46 0.14	0 0.55	0.23	59 <u>0.01</u>	×	14 0.85	9 0.40	<sup>2</sup> 0.00	39 0.46	.3 0.64	77 0.01	49 0.11	0.52	0.61		I.44 X	38 0.10	×	0 0.88	×	12 0.35	X 6	43 0.23	0.76	0.06	0.42
12 -1.	03 -1.28	32 -1.14	-1.07	62 -0.06	01 -1.63	00 -2.08	ς Ν/Α	-1.19	-1.07	14 -1.48	-0.05	-0.03	01 -2.19	ζ N/A	85 -1.16	40 0.10	2.08	46 -1.33	64 1.15	01 -2.70	-1.59	52 -0.04	-1.22	۲ N/A		10 0.43	۲ ۸/۸	88 0.08	ς N/Α	35 0.22	ζ 1.23	-1.31	76 0.01	06 -1.24	42 -1.26
40 0.(	28 0.13	14 0.08	07 0.01	06 1.04	63 0.17	08 0.24	'A N/A	19 0.16	07 0.01	48 0.02	05 1.45	03 1.25	19 0.50	'A N/A	16 0.02	10 1.10	0.84	33 0.06	0.02	0.07	59 0.10	04 1.11	0.01	A N/A	1.24 0.20	13 1.46	A N/A	)8 1.12	'A N/A	1.20	0.06	31 0.12	01 1.02	0.11	26 0.19
33	3	8(	)1	¥	17	ÿ4	A	16	Ĭ	72	5	35	50	A	2	10	¥	9	2	7(	10	Ξ	ĭ	A	0	16	A	12	A	30	9	12	)2	Ξ	9

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SPOA0116	SPOA0115	SPOA0114	SPOA0113	SPOA0112	SPOA0111	SPOA0110	SPOA0109	SPOA0108	SPOA0107	SPOA0106	SPOA0105	SPOA0104	SPOA0103	SPOA0102	SPOA0101	SPOA0100	SPOA0099	SPOA0098	SPOA0097	SPOA0096	SPOA0095	SPOA0094	SPOA0093	SPOA0091	SPOA0090	SPOA0089	SPOA0088	SPOA0087	SPOA0086	SPOA0085	SPOA0084	SPOA0083	SPOA0082	SPOA0081	SPOA0079
	gtdA-2			feaB																															
fumarylacetoacetate hydrolase family protein	gentisate 1,2-dioxygenase (EC:1.13.11.4)	nitrilase family protein	hypothetical protein	phenylacetaldehyde dehydrogenase (EC: 1.2.1.39)	indolepyruvate oxidoreductase, IorA subunit	Indolepyruvate oxidoreductase subunit B (EC:1.2.7.8)	hypothetical protein	hypothetical protein	LysR family transcriptional regulator	hypothetical protein	transcriptional regulator	aldehyde dehydrogenase	hypothetical protein	LuxR family transcriptional regulator	periplasmic branched-chain amino acid-binding protein	binding protein	branched-chain amino acid ABC transporter, ATP- binding protein branched-chain amino acid ABC transporter ATP-	permease protein	permease protein branched chain amino acid ABC transnorter	histone deacetylase/AcuC/AphA family protein branched-chain amino acid ABC transporter.	hypothetical protein	hypothetical protein	FAD binding domain-containing protein	IS3 family transposase orfA	ISS po7, transposase	hypothetical protein	hypothetical protein	phage integrase family site specific recombinase	hypothetical protein	DNA-binding protein	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	ribonuclease
-2.10	-1.67	-1.22	-1.51	-1.00	-1.56	-1.39	2.94	-1.53	-1.33	-1.77	-1.30	-1.87	0.98	-1.19	-1.38	Х	х	1.24	1.46	1.16	1.47	1.50	1.98	-1.06	x	x	1.68	-1.77	×	-2.16	-2.70	1.03	1.10	-1.25	-0.96
0.01	0.07	0.57	0.52	0.99	0.21	0.56	0.01	×	0.73	0.68	0.09	0.03	0.08	0.62	×	×	×	0.88	0.92	0.10	0.60	0.08	0.22	0.93	×	x	0.48	0.00	×	×	0.20	0.90	0.58	0.48	0.75
-1.61	-1.63	-1.34	-1.13	-1.26	-1.22	-1.17	1.44	-1.68	-1.31	-1.02	1.04	-1.08	-1.33	-1.26	-1.18	х	-1.40	-0.97	1.07	1.04	-1.15	1.06	-1.31	-1.22	х	x	1.14	-1.20	-1.34	1.44	-1.13	-1.30	-1.23	1.11	-1.28
0.02	0.08	0.32	0.86	0.51	0.61	0.74	0.04	0.25	0.66	0.95	0.46	0.73	0.07	0.57	0.82	×	0.08	0.38	0.94	0.45	0.82	0.36	0.39	0.79	×	х	0.73	0.09	0.04	0.08	0.21	0.41	0.07	0.75	0.34
-1.86	-1.65	-1.28	-1.32	-1.13	-1.39	-1.28	2.19	-1.61	-1.32	-1.40	-0.13	-1.48	-0.18	-1.23	-1.28	N/A	N/A	0.13	1.27	1.10	0.16	1.28	0.34	-1.14	N/A	N/A	1.41	-1.49	N/A	-0.36	-1.92	-0.14	-0.06	-0.07	-1.12
0.25	0.02	0.06	0.19	0.13	0.17	0.11	0.75	0.08	0.01	0.38	1.17	0.40	1.15	0.04	0.10	N/A	N/A	1.11	0.19	0.06	1.31	0.22	1.65	0.08	N/A	N/A	0.27	0.29	N/A	1.80	0.79	1.17	1.17	1.18	0.16
-3.02	-2.78	-1.93	-2.26	-1.55	-2.14	-2.02	2.49	-1.48	-0.93	-1.17	-1.02	-2.31	-2.84	-2.05	-2.40	Х	-2.64	1.02	1.66	-1.25	1.66	1.32	1.52	-2.22	х	-1.39	1.14	-1.82	-1.15	-1.19	-3.33	-1.40	1.54	-1.29	1.98
0.02	0.01	0.28	0.27	0.82	0.13	0.36	0.06	0.90	0.71	0.91	0.13	0.02	0.00	0.21	0.21	x	0.11	0.96	0.88	0.71	0.51	0.11	0.22	0.67	×	0.96	0.93	0.00	0.68	0.82	0.04	0.38	0.06	0.53	0.39
-1.04	-1.03	-1.46	-1.17	-1.24	-1.25	-1.25	1.08	-1.20	-1.48	1.10	-1.36	-1.05	1.10	1.03	-1.24	Х	-1.46	Х	-1.02	1.23	-1.51	-1.02	-1.08	1.23	Х	Х	1.68	1.02	-1.01	-1.10	-1.09	1.35	1.59	1.74	-1.09
0.72	0.83	0.31	0.73	0.35	0.13	0.66	0.35	0.82	0.67	0.91	0.05	0.89	0.39	0.91	x	×	0.29	×	0.99	0.08	0.45	0.67	0.42	0.76	×	х	0.26	0.87	0.71	0.75	0.78	0.34	0.06	0.22	0.61
-2.03	-1.91	-1.70	-1.72	-1.40	-1.70	-1.64	1.79	-1.34	-1.21	-0.03	-1.19	-1.68	-0.87	-0.51	-1.82	N/A	-2.05	N/A	0.32	-0.01	0.08	0.15	0.22	-0.50	N/A	N/A	1.41	-0.40	-1.08	-1.15	-2.21	-0.02	1.57	0.23	0.45
0.99	0.88	0.24	0.55	0.16	0.45	0.39	0.71	0.14	0.27	1.14	0.17	0.63	1.97	1.54	0.58	N/A	0.59	N/A	1.34	1.24	1.59	1.17	1.30	1.73	N/A	N/A	0.27	1.42	0.07	0.04	1.12	1.38	0.03	1.52	1.54
-1.65	-1.55	-1.43	-1.09	-1.48	-1.55	-1.42	-1.41	-1.34	1.37	-1.05	-1.17	1.32	-1.22	1.02	-1.04	-1.13	1.08	1.13	1.22	-1.25	1.26	1.49	1.02	-1.19	х	x	1.30	-1.14	-1.08	-2.15	-1.12	1.06	-1.07	-1.03	-1.27
0.08	0.04	0.18	0.81	0.14	0.20	0.45	0.03	0.72	0.68	0.93	0.17	0.49	0.43	0.94	0.98	х	0.73	0.45	0.80	0.06	0.62	0.11	0.93	0.80	x	x	0.36	0.30	0.60	0.02	0.82	0.89	0.61	0.96	0.18
-1.55	-1.06	-1.24	1.22	-1.17	-0.97	-1.28	-1.49	-0.93	-1.25	1.25	-1.52	2.41	-1.22	1.14	×	Х	2.50	Х	2.05	-1.55	1.28	1.28	1.27	1.21	x	x	-1.16	-0.99	х	-1.92	1.54	1.11	1.20	1.15	-1.55
0.11	0.97	0.67	×	0.88	0.88	0.60	0.08	0.55	0.91	0.77	0.07	0.06	0.60	0.65	×	Х	0.04	х	х	Х	0.56	0.07	0.27	0.72	х	Х	0.52	0.75	x	0.11	0.05	0.76	0.10	0.69	0.17
-1.60	-1.31	-1.34	0.06	-1.33	-1.26	-1.35	-1.45	-1.14	0.06	0.10	-1.35	1.87	-1.22	1.08	N/A	N/A	1.79	N/A	1.64	-1.40	1.27	1.39	1.15	0.01	N/A	N/A	0.07	-1.07	N/A	-2.04	0.21	1.09	0.06	0.06	-1.41
0.05	0.24	0.10	1.16	0.15	0.29	0.07	0.04	0.21	1.31	1.15	0.18	0.54	0.00	0.06	N/A	N/A	0.71	N/A	0.42	0.15	0.01	0.11	0.12	1.20	N/A	N/A	1.23	0.07	N/A	0.12	1.33	0.03	1.14	1.09	0.14

# Chapter 10: Appendix

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_	hypothetical protein	)154	SPOA0154
	hypothetical protein	)153	SPOA0153
otein	di-haem cytochrome c peroxidase family protein	)152	SPOA0152
	cytochrome c family protein	)151	SPOA0151
	AraC family transcriptional regulator	)150	SPOA0150
	GntR family transcriptional regulator	)149	SPOA0149
	major facilitator transporter	)148	SPOA0148
nily	ornithine cyclodeaminase/mu-crystallin family protein	0147	SPOA0147
	threonine aldolase (EC:4.1.2.5)	)146	SPOA0146
	pyridoxal-phosphate dependent enzyme	)145	SPOA0145
	serineglyoxylate transaminase	)144	SPOA0144
	IclR family transcriptional regulator	)143	SPOA0143
	hypothetical protein	)142	SPOA0142
	hypothetical protein	0141	SPOA0141
6)	glucosamnetructose-6-phosphate glmS aminotransferase (isomerizing) (EC:2.6.1.16)		SPOA0140
	acetyltransferase	)139	SPOA0139
	hypothetical protein	)138	SPOA0138
	murA carboxyvinyltransferase (EC:2.5.1.7)		SPOA0137
	hypothetical protein	)136	SPOA0136
	ISSpo7, transposase	)134	SPOA0134
	oxidoreductase	)133	SPOA0133
	aromatic 1,2-dioxygenase, alpha subunit	)132	SPOA0132
	aromatic 1,2-dioxygenase, beta subunit	)131	SPOA0131
	AMP-binding protein	)130	SPOA0130
	hypothetical protein	)129	SPOA0129
	snort criain denydrogenase/reductase family oxidoreductase	)128	SPOA0128
		)127	SPOA0127
	short chain denydrogenase/reductase family oxidoreductase	)126	SPOA0126
	hypothetical protein	)125	SPOA0125
	hypothetical protein	)124	SPOA0124
	hypothetical protein	)123	SPOA0123
	ISSpo9, transposase	)122	SPOA0122
	sulfatase family protein	)121	SPOA0121
	hypothetical protein	)120	SPOA0120
	hypothetical protein	)1 19	SPOA0119
	hypothetical protein	)118	SPOA0118
L	Asp/Glu/hydantoin racemase family protein	0117	SPOA0117
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-1.41	x	×	x	-1.92	x	-1.49	-1.11	1.07	-1.01	-2.55	-1.01	-2.23	2.59	1.54	1.52	1.21	1.69	х	х	-1.06	х	-1.24	х	1.82	1.07	-1.15	1.11	1.24	-1.78	-1.90	-1.18	-3.45	-1.96	-1.54	-1.39
X 0.78	х	х	Х	0.37	×	0.01	0.78	0.92	0.88	0.08	0.97	0.00	0.06	0.87	0.05	0.43	0.04	х	×	0.92	×	0.18	x	0.28	0.88	0.94	0.90	0.43	0.27	0.15	0.92	0.05	0.24	0.21	0.25
X -1.16	-1.38	×	-0.98	-1.25	-1.22	-1.14	-1.16	-1.15	-1.19	1.06	-1.43	-1.37	-1.03	-1.04	1.06	-1.19	-1.39	1.33	-1.47	-1.13	-1.10	-1.09	-1.17	-1.26	-1.18	-1.08	-1.25	-1.09	1.05	-1.60	-1.00	-1.56	-1.54	-1.46	-1.29
0.81	Х	Х	0.73	0.18	0.73	0.38	0.80	0.48	0.38	0.78	0.04	0.08	0.94	0.96	0.82	0.25	0.14	0.01	0.15	0.83	0.41	0.80	0.70	0.66	0.38	0.91	0.09	0.82	0.93	0.18	0.98	0.05	0.08	0.66	0.05
N/A -1.29	N/A	N/A	N/A	-1.59	N/A	-1.32	-1.14	-0.04	-1.10	-0.75	-1.22	-1.80	0.78	0.25	1.29	0.01	0.15	N/A	N/A	-1.10	N/A	-1.17	N/A	0.28	-0.05	-1.12	-0.07	0.08	-0.37	-1.75	-1.09	-2.51	-1.75	-1.50	-1.34
N/A 0.13	N/A	N/A	N/A	0.34	N/A	0.18	0.02	1.11	0.09	1.81	0.21	0.43	1.81	1.29	0.23	1.20	1.54	N/A	N/A	0.03	N/A	0.08	N/A	1.54	1.13	0.03	1.18	1.17	1.42	0.15	0.09	0.95	0.21	0.04	0.05
X -1.07	x	-1.18	1.35	-1.79	-2.48	-2.09	-1.81	-3.52	-2.37	-3.36	1.24	-2.18	-1.86	2.22	2.85	1.82	2.21	Х	x	-2.48	-0.97	-1.24	х	1.15	-1.75	-1.65	-1.36	1.03	0.87	-1.71	-1.46	-5.09	-2.82	-3.30	-2.31
0.95	Х	0.28	0.24	0.20	0.09	0.07	0.43	0.02	0.00	0.01	0.17	0.00	0.25	0.73	0.03	0.09	0.03	х	х	0.28	0.43	0.79	X	0.73	0.26	0.84	0.22	0.93	0.44	0.12	0.86	0.03	0.11	0.05	0.04
-1.01	х	х	Х	-1.31	-1.49	-1.11	1.15	1.07	-1.20	-1.87	-1.17	-1.21	-1.96	-1.06	-1.35	1.06	1.28	-1.22	х	-1.17	-1.21	-1.39	Х	-1.19	-1.24	1.16	1.36	1.26	1.28	-1.26	-0.98	-1.48	-1.15	1.06	-1.03
X 0.99	х	х	Х	0.05	0.43	0.23	0.80	0.67	0.59	0.02	0.09	0.37	0.27	0.93	0.04	0.98	0.20	0.05	×	0.76	×	0.21	×	0.86	0.45	0.49	0.18	0.53	0.70	0.16	0.81	0.07	0.70	0.96	0.89
N/A -1.04	N/A	N/A	N/A	-1.55	-1.99	-1.60	-0.33	-1.23	-1.79	-2.62	0.04	-1.70	-1.91	0.58	0.75	1.44	1.75	N/A	N/A	-1.83	-1.09	-1.32	N/A	-0.02	-1.50	-0.25	0.00	1.15	1.07	-1.49	-1.22	-3.29	-1.99	-1.12	-1.67
N/A 0.03	N/A	N/A	N/A	0.24	0.50	0.49	1.48	2.30	0.59	0.74	1.21	0.49	0.05	1.64	2.10	0.38	0.47	N/A	N/A	0.66	0.12	0.08	N/A	1.17	0.26	1.41	1.36	0.12	0.21	0.23	0.24	1.81	0.84	2.18	0.64
X -1.23	х	1.14	-0.98	-1.36	1.08	-0.97	-1.22	-1.28	-1.13	-1.18	1.28	1.42	-1.05	1.30	1.33	-1.08	1.01	-1.42	-1.24	-1.11	-1.44	-1.12	-1.88	-1.31	-1.44	-1.04	-1.13	1.21	1.43	-1.32	-1.02	-1.21	-1.10	-1.26	1.06
X 0.76	х	0.23	0.53	0.15	0.78	0.43	0.69	0.03	0.24	0.06	0.07	0.05	0.90	0.75	0.03	0.96	0.84	0.06	0.12	0.81	0.17	0.66	×	0.77	0.16	0.96	0.38	0.63	0.47	0.82	0.96	0.13	0.44	0.75	0.66
X 1.53	x	×	х	-1.39	1.34	-0.94	1.12	-1.09	-0.90	-1.36	1.13	1.23	1.34	-1.07	-1.17	1.03	1.29	-2.25	х	1.17	-1.19	-0.99	х	1.40	-1.53	1.25	-1.03	1.16	1.55	1.77	-1.04	-1.39	-1.23	-1.03	-0.91
0.56	х	х	Х	0.15	0.56	0.11	0.77	0.95	0.08	0.22	0.73	0.12	0.66	0.95	0.58	0.64	0.21	×	×	0.25	×	0.77	×	0.20	0.24	0.84	0.80	0.63	0.40	0.04	0.77	0.13	0.63	0.97	0.26
N/A 0.15	N/A	N/A	N/A	-1.38	1.21	-0.96	-0.05	-1.19	-1.01	-1.27	1.21	1.33	0.15	0.12	0.08	-0.03	1.15	-1.84	N/A	0.03	-1.32	-1.06	N/A	0.04	-1.49	0.11	-1.08	1.19	1.49	0.23	-1.03	-1.30	-1.17	-1.15	0.08
N/A 1.38	N/A	N/A	N/A	0.01	0.13	0.02	1.17	0.10	0.12	0.09	0.08	0.10	1.20	1.19	1.25	1.06	0.14	0.42	N/A	1.14	0.13	0.06	N/A	1.36	0.05	1.15	0.05	0.03	0.06	1.55	0.01	0.09	0.06	0.12	0.98

# Chapter 10: Appendix

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SPOA0192	SPUA0191	SPOA0190	SPOA0189	SPOA0188	SPOA0187	SPOA0186	SPOA0185	SPOA0184	SPOA0183	SPOA0182	SPOA0181	SPOA0180	SPOA0177	SPOA0176	SPOA0175	SPOA0174	SPOA0173	SPOA0172	SPOA0171	SPOA0170	SPOA0169	SPOA0168	SPOA0167	SPOA0166	SPOA0165	SPOA0164	SPOA0163	SPOA0162	SPOA0161	SPOA0160	SPOA0159	SPOA0158	SPOA0157	SPOA0156	SPOA0155
		ccoN-2	ccoO-2	ccoQ-2	ccoP-2																						mvaA				cuyR	cuyA			
hypothetical protein	пуротненсат ртотеш	(EC:1.9.3.1)	cytochrome c oxidase, cob3-type, subunit I (EC:1.9.3.1)	(EC:1.9.3.1)	cytochrome c oxidase, cbb3-type, subunit III (EC:1.9.3.1) (EC:1.9.3.1)	Rrf2 family protein	cytochrome c oxidase,ba3-type, subunit I	heme-copper respiratory oxidase family	SCO1/SenC family protein	hypothetical protein	FNT family protein	cytochrome c family protein	hydrogen peroxide-inducible genes activator	hypothetical protein	oxidoreductase	MOSC domain-containing protein	hypothetical protein	hypothetical protein	hypothetical protein	MarR family transcriptional regulator	MOSC domain-containing protein	oxidoreductase, FAD-binding	TRAP transporter, 4TM/12TM fusion protein	protein	LysR family transcriptional regulator TR AP transporter solute recentor TAXI family	GntR family transcriptional regulator	hydroxymethylglutaryl-CoA reductase, degradative (EC:1.1.1.88)	subunit	TRAP dicarboxylate transporter, DctQ subunit	TRAP dicarboxylate transporter, DctM subunit	transcriptional regulator CuyR	L-cysteate sulfo-lyase (EC:4.4.1.15)	sulfite exporter, CuyZ	hypothetical protein	type I secretion target repeat-containing protein
-1.86	1.07	X	-1.39	-3.40	Х	-1.38	Х	1.10	-1.97	Х	Х	-1.06	Х	-2.32	1.06	-1.09	-1.23	-1.79	-1.60	-1.99	-1.35	-2.28	Х	-1.85	-2.09	-1.03	Х	-2.49	-2.06	Х	-1.31	1.46	-1.82	-1.26	x
0.06	>	××	0.81	0.19	×	0.24	×	0.77	0.38	×	×	0.97	×	0.04	0.53	0.41	0.59	0.01	0.62	0.04	0.36	0.02	x	0.48	0.33	0.97	×	0.25	х	×	0.50	0.14	0.82	0.74	×
1.32	0.1	1.29	-1.07	-1.41	1.10	-1.47	-1.01	1.17	-1.12	1.12	-1.00	1.38	-1.38	-1.06	-1.03	-1.05	-1.22	-1.38	-1.30	-1.36	-1.20	-1.07	-1.27	-1.79	-0.99	1.13	х	-1.01	-1.27	1.35	-0.99	1.13	-1.07	-1.15	2.39
0.33	0.04	0.32	0.96	0.16	0.81	0.02	0.97	0.47	0.50	0.59	0.93	0.07	Х	0.93	0.94	0.85	0.18	0.23	0.22	0.02	0.31	0.79	0.36	0.62	0.96	0.64	×	0.98	0.81	0.27	0.83	0.27	0.95	0.48	0.00
-0.27	1.30	N/A	-1.23	-2.41	N/A	-1.43	N/A	1.14	-1.55	N/A	N/A	0.16	N/A	-1.69	0.02	-1.07	-1.23	-1.59	-1.45	-1.68	-1.28	-1.68	N/A	-1.82	-1.54	0.05	N/A	-1.75	-1.67	N/A	-1.15	1.30	-1.45	-1.21	N/A
1.59	0.20	N/A	0.16	1.00	N/A	0.05	N/A	0.03	0.43	N/A	N/A	1.22	N/A	0.63	1.05	0.02	0.01	0.20	0.15	0.32	0.08	0.61	N/A	0.03	0.55	1.08	N/A	0.74	0.39	N/A	0.16	0.16	0.38	0.06	N/A
-2.11	1.32	1.31	1.32	-6.39	1.37	1.46	-0.89	-1.01	-2.38	-2.02	2.43	-0.98	-1.34	-3.89	-1.39	-1.70	-1.95	-3.50	-1.04	-1.53	-1.23	-3.13	-1.24	-1.87	-2.68	-1.35	-1.61	-4.53	-1.86	-1.34	-1.37	-1.52	-1.26	1.52	2.79
0.04	0.32	0.04	0.85	0.01	x	0.01	0.40	0.47	0.04	0.08	x	0.40	х	0.01	0.41	0.11	0.02	0.00	0.92	0.10	0.58	0.01	0.64	0.68	0.16	0.93	0.23	0.11	0.73	0.89	0.90	0.15	0.95	0.45	0.10
-1.22	-1.50	-1.23	-1.01	-1.30	-1.18	1.23	-1.50	-1.34	-1.50	-1.21	-1.09	-1.02	-1.19	1.07	-1.11	1.07	-1.14	-1.07	-1.82	-1.12	-1.20	1.52	-1.58	-1.35	-1.38	-1.44	X	-1.21	-1.55	-1.35	-1.29	-1.12	-1.38	1.19	-1.44
0.49			0.99	0.44	0.49	0.07	0.61	0.25	0.41	0.61	0.91	0.96	x	0.75	0.69	0.24	0.50	0.74	0.05	0.29	0.41	0.04	0.23	0.77	0.66	0.17	x	0.55	0.80	0.43	0.52	0.33	0.69		0.10
-1.67			0.16	-3.85	0.10	1.35	-1.19	-1.18		-1.62	0.67	-1.00	-1.27	-1.41		-0.32		-2.29	-1.43	-1.33	-1.22	-0.81	-1.41		-2.03		N/A	-2.87			-1.33			1.36	
0.45			1.17	5 2.55	) 1.28	0.12	0.31	3 0.17	4 0.44	0.41	1.76	0.02	0.08	1 2.48	0.14	2 1.39	5 0.41	€ 1.22	3 0.39	3 0.20	2 0.02	1 2.33	0.17	0.26	3 0.65	0.04	N/A	7 1.66	0.16	5 0.01	3 0.04	2 0.20	0.06		3 2.12
-1.41			7 -1.17	5 -1.62	8 -1.16	2 -1.07	1 -1.60	7 -1.23		1 -1.22	6 -1.32	2 1.05	8 1.42	8 -1.49	4 -1.27	9 -1.36	1 -1.12	2 1.04	9 -1.17	0 -1.34	2 1.03	3 -1.01		6 -1.48	5 1.22	4 -1.03	A X	6 -1.00	6 -1.52	-1.28		0 -1.09			2 1.36
H 0.02			0.87	62 0.05	16 0.77	0.19	50 0.06	0.53	8 0.75	0.23	32 0.51	5 0.53	2 X	0.23	0.36	36 0.05	0.45	4 0.93	0.29	34 0.17	3 0.97	0.88	1 0.74	18 0.63	2 0.75	0.99	x	0 0.94	52 0.72	0.53	7 0.52	0.15			6 0.11
2 -1.09			7 1.15	5 1.30	7 1.63	9 -1.40	6 -1.17	3 -1.21	5 1.23	3 -1.35	1 X	3 -1.70	2.49	3 -1.18	6 -1.64	5 -1.42	5 -1.44				7 -1.18	8 -0.93	4 1.21	3 1.21	5 -0.97	9 8	X	4 1.69	2 -0.96		2 -1.13				1 X
0.93																				61 0.03	18 0.37								96 0.93						
			0.89 -0.	0.51 -0.16	X 0.2	0.11 -1.	X -1.	0.81 -1.	0.23 1.1	X -1.	X N	X -0.	X 1.9	0.81 -1.	0.16 -1.46	0.05 -1.	0.14 -1.	0.09 1.5	0.11 -1.	03 -1.48		0.38 -0.	0.19 1.1	0.88 -0.	0.82 0.13	X V	X N/A	0.04 0.35		X -1.		0.29 -1.			X N/A
-1.25 0			-0.01 1		0.24 1	-1.24 0	-1.39 0	-1.22 0	1.16 0	-1.29 0	N/A N	-0.33 1	1.96 0	-1.34 0		-1.39 0	-1.28 0	1.57 0	-1.30 0		-0.08 1	-0.97 0		-0.14 1								-1.14 0			
0.16	1.10	N/A	1.16	1.46	1.40	0.17	0.22	0.01	0.08	0.07	N/A	1.38	0.53	0.16	0.18	0.03	0.16	0.53	0.13	0.14	1.11	0.04	0.05	1.35	1.09	N/A	N/A	1.35	0.28	0.09	1.10	0.05	1.22	0.04	N/A

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# Chapter 10: Appendix

	nirN	nirJ	nirH	nirG		nirF	nirC	nirE	nirS			norC	norB	norQ	norD												hisD				cycK	dsbE	cycL		hemA-2	cycJ
hypothetical protein LyR family transcriptional regulator glycine betaine/proline ABC transporter, periplasmic substrate-binding protein	nitrite reductase protein N (EC:1.7.2.1)	nitrite reductase heme biosynthesis J protein	nitrite reductase heme biosynthesis H protein	nitrite reductase heme biosynthesis G protein	nitrite reductase heme biosynthesis D/L protein	cytochrome cd1 nitrite reductase (EC:1.7.2.1)	cytochrome c55X	nitrite reductase heme biosynthesis E protein	cytochrome cd1 nitrite reductase (EC:1.7.2.1)	NnrS	magnesium transporter CorA family protein	nitric oxide reductase, small subunit (EC:1.7.99.7)	nitric oxide reductase, large subunit (EC:1.7.99.7)	nitric oxide reductase Q protein	nitric oxide reductase D protein	nitric oxide reductase E protein	nitric oxide reductase F protein	Na/Pi-cotransporter family protein	hypothetical protein	oxidoreductase, FAD-binding	oxidoreductase, FAD-binding	guanylate cyclase	Crp/Fnr family transcriptional regulator	serine protease	hypothetical protein	hypothetical protein	histidinol dehydrogenase (EC:1.1.1.23)	hydroxylamine oxidoreductase (EC:1.7.3.4)	hypothetical protein	GntR family transcriptional regulator	cytochrome c-type biogenesis protein Cyc	thiol:disulfide interchange protein DsbE	cytochrome c-type biogenesis protein CycL	hypothetical protein	5-aminolevulinate synthase (EC:2.3.1.37)	cytochrome c-type biogenesis protein CcmE
1.73 -1.60 0.98	Х	-1.19	-1.24	Х	1.57	-2.23	1.17	Х	1.14	Х	-1.17	Х	1.34	-2.34	Х	х	-1.14	1.09	-1.40	-2.28	-1.52	-1.18	-1.34	0.99	-1.09	-1.40	-1.03	-1.95	-2.15	Х	-1.38	-1.49	Х	1.14	1.47	-1.76
0.25 0.15 0.88	×	0.73	0.34	x	0.02	0.02	0.91	×	0.97	×	0.37	×	×	0.05	×	×	0.88	0.07	0.49	0.04	0.10	0.19	0.22	0.73	0.90	0.23	0.98	х	×	х	×	0.16	×	0.71	0.49	0.33
1.17 -1.25 -1.31	-1.76	-1.46	-1.12	-1.10	-1.23	-1.14	-1.08	-1.23	1.05	-1.00	1.13	1.17	-1.08	-1.40	-1.51	-1.16	-1.20	1.21	-1.01	-1.17	1.02	1.02	-1.21	-1.01	-1.26	1.12	1.01	1.18	-1.13	-1.08	-1.03	-1.53	1.18	-1.26	-1.01	-1.31
0.07 0.19 0.48	×	0.54	0.66	х	0.83	0.66	0.90	0.45	0.78	0.91	0.31	0.21	0.77	0.15	x	0.19	0.83	0.03	0.98	0.29	0.94	0.81	0.19	0.69	0.15	0.76	0.99	0.59	0.87	0.62	0.99	0.16	0.34	0.33	1.00	0.15
1.45 -1.43 -0.16	N/A	-1.33	-1.18	N/A	0.17	-1.69	0.04	N/A	1.10	N/A	-0.02	N/A	0.13	-1.87	N/A	N/A	-1.17	1.15	-1.21	-1.73	-0.25	-0.08	-1.28	-0.01	-1.18	-0.14	-0.01	-0.39	-1.64	N/A	-1.21	-1.51	N/A	-0.06	0.23	-1.54
0.28 0.18 1.15	N/A	0.14	0.06	N/A	1.40	0.55	1.13	N/A	0.04	N/A	1.15	N/A	1.21	0.47	N/A	N/A	0.03	0.06	0.19	0.56	1.27	1.10	0.07	1.00	0.09	1.26	1.02	1.57	0.51	N/A	0.17	0.02	N/A	1.20	1.24	0.22
1.97 -0.99 -3.17	-4.95	-1.61	-1.79	-0.99	-8.53	-2.55	-1.13	-2.09	-1.93	-1.05	-1.80	х	-1.06	-2.93	х	-2.26	-1.08	1.51	-1.64	-2.06	0.93	1.12	1.08	0.87	-1.50	-1.59	3.50	-2.00	-1.82	-0.96	-1.43	-1.36	-1.15	-1.29	-0.93	-1.86
0.02 0.63 0.18	0.01	0.73	0.30	0.07	0.00	0.07	0.92	0.04	0.82	0.22	0.38	x	0.61	0.00	х	0.04	0.99	0.01	0.48	0.02	0.13	0.36	0.67	0.26	0.11	0.30	0.01	0.59	0.79	0.12	0.93	0.26	0.44	0.25	0.52	0.38
-0.95 -1.39 1.68	1.20	-1.02	1.01	x	-0.97	-1.05	-1.12	х	-1.41	-1.47	1.67	1.22	-1.08	-1.51	x	1.38	-1.17	-1.52	-2.27	-1.04	1.28	-1.06	1.15	1.20	1.18	1.13	-1.84	1.06	-1.20	x	1.28	-1.45	-1.10	-1.34	-1.06	-1.05
0.61 0.23 0.31	x	1.00	х	х	0.62	0.88	0.86	x	x	0.15	0.07	х	x	0.12	×	0.31	0.30	0.02	0.08	0.55	0.35	0.56	0.36	0.19	0.29	0.80	0.05	х	0.81	×	x	0.40	х	0.49	0.87	0.92
0.51 -1.19 -0.75	-1.88	-1.32	-0.39	N/A	-4.75	-1.80	-1.13	N/A	-1.67	-1.26	-0.07	N/A	-1.07	-2.22	N/A	-0.44	-1.13	-0.01	-1.96	-1.55	1.10	0.03	1.12	1.03	-0.16	-0.23	0.83	-0.47	-1.51	N/A	-0.08	-1.41	-1.13	-1.32	-0.99	-1.46
1.46 0.20 2.43	3.08	0.30	1.40	N/A	3.78	0.75	0.00	N/A	0.26	0.21	1.74	N/A	0.01	0.71	N/A	1.82	0.04	1.52	0.31	0.51	0.18	1.09	0.03	0.17	1.34	1.36	2.67	1.53	0.31	N/A	1.36	0.04	0.02	0.03	0.07	0.41
-1.02 1.31 -1.04	-1.24	-1.16	-0.98	-1.48	-1.20	-1.35	1.11	1.18	1.23	1.07	1.49	-1.07	1.14	-1.51	1.06	-1.41	-1.16	-1.03	1.40	-1.13	1.69	1.47	1.03	1.77	1.23	1.52	-1.40	1.46	-1.01	-0.97	-1.05	-1.16	1.32	1.17	1.09	-1.21
0.98 0.31 0.94	x	0.85	0.78	0.04	0.24	0.08	0.83	0.15	0.13	0.50	0.04	0.73	0.70	0.09	×	0.20	0.78	0.34	0.16	0.72	0.10	0.02	0.95	0.02	0.33	0.35	0.03	0.06	0.95	0.64	0.99	0.50	0.16	0.28	0.65	0.62
X -0.94 1.26	x	1.32	х	х	-1.47	1.25	1.45	1.42	-1.02	-1.10	1.57	x	x	1.26	x	1.59	1.88	-1.20	-1.08	2.24	2.02	1.03	1.87	1.37	1.43	1.34	-2.18	х	1.56	x	-1.10	2.34	x	3.23	-1.02	1.90
0.60 0.58	x	0.72	х	х	x	0.35	0.38	x			0.06	x	x	0.45	x	0.27	x	0.15	0.62	0.03	0.10	0.81	0.01	0.09	0.05	0.49	0.00	x	x	×	x	0.16	х	0.03	0.82	0.21
N/A 0.19 0.11	N/A	0.08	N/A	N/A	-1.34	-0.05	1.28	1.30	0.11	-0.02	1.53	N/A	N/A	-0.13	N/A	0.09	0.36	-1.12	0.16	0.56	1.86	1.25	1.45	1.57	1.33	1.43	-1.79	N/A	0.28	N/A	-1.08	0.59	N/A	2.20	0.04	0.35
N/A 1.12 1.15	N/A	1.24	N/A	N/A	0.14	1.30	0.17	0.12	1.13	1.09	0.04	N/A	N/A	1.39	N/A	1.50	1.52	0.09	1.24	1.69	0.17	0.22	0.42	0.20	0.10	0.09	0.39	N/A	1.29	N/A	0.03	1.75	N/A	1.03	1.06	1.56

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SPOA0224 SPOA0225 SPOA0226 SPOA0227

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SPOA0214

SPOA0215

SPO A0202 SPO A0203 SPO A0204 SPO A0205 SPO A0206 SPO A0207 SPO A0207 SPO A0208 SPO A0209 SPO A0210 SPO A0211 SPO A0212 SPO A0212

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SPOA0267	SPOA0266	SPOA0265	SPOA0264	SPOA0263	SPOA0262	SPOA0261	SPOA0260	SPOA0259	SPOA0258	SPOA0257	SPOA0256	SPOA0255	SPOA0254	SPOA0253	SPOA0252	SPOA0251	SPOA0250	SPOA0249	SPOA0248	SPOA0247	SPOA0246	SPOA0245	SPOA0244	SPOA0243	SPOA0242	SPOA0241	SPOA0240	SPOA0239	SPOA0238	SPOA0237	SPOA0236	SPOA0235	SPOA0234	SPOA0233	SPOA0232
									rbsA	rbsC-2			rbsC-1								pdxA									dctD-2	dctB	speB-4			opuAA
dihydroxydipicolinate synthase family protein	proline racemase	TRAP transporter DctP family protein	TRAP transporter DctQ family protein	TRAP transporter, DctM subunit	oxidoreductase, FAD-binding	GntR family transcriptional regulator	aldehyde dehydrogenase	LacI family transcription regulator	ribose ABC transporter, ATP-binding protein	ribose ABC transporter, permease protein	protein	sugar ABC transporter, ATP binding protein	ribose ABC transporter, permease protein	protein	LacI family transcription regulator	TRAP dicarboxylate transporter, DctQ subunit	TRAP dicarboxylate transporter, DctM subunit	TRAP dicarboxylate transporter, DctP subunit	hypothetical protein	hypothetical protein	4-nydrox ymreonine-4-pnospnate denydrogenase (EC:1.1.1.262)	galactonate dehydratase	dihydrodipicolinate synthase family protein	aldehyde dehydrogenase	LacI family transcription regulator	gluconolactonase	TRAP transporter, DctM subunit	TRAP dicarboxylate transporter, DctQ subunit	TRAP dicarboxylate transporter, DctP subunit	C4-dicarboxylate transport transcriptional regulatory protein DctD	C4-dicarboxylate transport sensor protein detB (EC:2.7.3)	agmatinase (EC:3.5.3.11)	agmatinase	giycine betaine/proline ABC transporter, permease protein	glycine betaine/proline ABC transporter, ATP- binding protein (EC:3.6.3.2)
1.03	-1.12	1.01	-1.86	х	х	-1.63	-0.95	-1.47	-1.26	-1.17	-1.35	×	х	-3.01	-1.26	-1.21	x	-1.22	-1.25	x	-1.20	-0.97	1.12	-1.01	-1.44	-1.84	-1.43	-1.10	-1.37	1.10	-2.47	-1.05	1.26	х	-1.08
×	0.95	0.97	x	×	×	0.54	1.00	0.19	0.82	0.81	0.61	×	x	×	0.03	0.83	×	0.50	0.28	×	0.65	0.85	0.36	0.96	0.15	0.15	0.06	0.53	0.01	0.43	0.10	0.38	0.82	×	0.97
-1.16	-1.35	-1.59	-1.34	-1.15	-1.19	1.16	-1.35	-1.33	-1.30	-1.45	-1.94	-2.49	-1.47	-1.66	-1.15	-1.21	x	-1.92	-1.31	-1.16	-1.23	-1.46	-1.06	1.00	-1.14	-1.12	-1.47	-1.82	-1.88	1.14	-1.18	1.18	-1.01	-1.07	-1.08
0.67	0.25	0.16	0.83	0.21	х	0.72	0.07	0.10	0.54	0.38	0.36	0.11	0.31	0.66	0.28	0.68	×	0.17	0.09	0.38	0.26	0.04	0.54	0.98	0.80	0.81	0.03	0.02	0.03	0.72	0.64	0.12	0.99	0.84	0.96
-0.06	-1.24	-0.29	-1.60	N/A	N/A	-0.24	-1.15	-1.40	-1.28	-1.31	-1.65	N/A	N/A	-2.34	-1.21	-1.21	N/A	-1.57	-1.28	N/A	-1.22	-1.22	0.03	-0.01	-1.29	-1.48	-1.45	-1.46	-1.63	1.12	-1.83	0.06	0.13	N/A	-1.08
1.10	0.12	1.30	0.26	N/A	N/A	1.40	0.20	0.07	0.02	0.14	0.30	N/A	N/A	0.68	0.06	0.00	N/A	0.35	0.03	N/A	0.02	0.24	1.09	1.01	0.15	0.36	0.02	0.36	0.26	0.02	0.65	1.12	1.14	N/A	0.00
1.28	-2.48	-1.77	-1.09	-1.82	х	-1.10	1.31	-2.56	-3.61	-4.33	-6.26	-2.30	-1.81	-2.11	-2.21	-1.65	x	-2.40	-1.61	-1.44	-2.58	-1.39	-1.17	-1.19	-1.34	-2.11	-1.54	-1.21	-4.86	1.32	-1.95	-1.57	-1.45	-0.87	-1.68
0.58	0.48	0.66	0.98	0.12	x	0.91	0.11	0.02	0.50	0.25	0.08	0.10	0.56	0.68	0.02	0.68	x	0.04	0.22	×	0.06	0.53	0.75	0.84	0.16	0.17	0.04	0.17	0.00	0.20	0.10	0.05	0.78	0.43	0.83
-1.53	-1.36	-1.32	-1.26	-1.39	x	-1.12	-1.35	-1.20	-0.98	-1.33	1.09	-1.08	-1.31	-1.08	-1.02	-1.16	x	1.14	-1.41	-1.18	-1.42	-1.70	-1.06	-1.26	-1.01	-1.31	1.63	1.65	1.69	-1.68	-1.15	-1.11	1.14	-1.02	-1.01
0.14	0.59	0.45	0.91	0.09	x	0.79	0.16	0.44	0.95	0.61	0.84	0.85	0.33	0.94	0.65	0.81	x	0.40	0.13	x	0.09	0.04	0.69	0.60	0.94	0.55	0.04	0.07	0.06	0.08	0.51	0.47	0.82	0.95	1.00
-0.13	-1.92	-1.55	-1.18	-1.61	N/A	-1.11	-0.02	-1.88	-2.30	-2.83	-2.59	-1.69	-1.56	-1.60	-1.62	-1.41	N/A	-0.63	-1.51	-1.31	-2.00	-1.55	-1.12	-1.23	-1.18	-1.71	0.04	0.22	-1.59	-0.18	-1.55	-1.34	-0.16	-0.95	-1.35
1.41	0.56	0.23	0.09	0.22	N/A	0.01	1.33	0.68	1.31	1.50	3.68	0.61	0.25	0.52	0.60	0.25	N/A	1.77	0.10	0.13	0.58	0.16	0.05	0.04	0.17	0.40	1.59	1.43	3.28	1.50	0.40	0.23	1.30	0.07	0.34
-1.36	-1.34	-1.07	1.21	-1.36	х	1.32	1.05	-1.18	-1.37	-1.16	-1.02	-1.19	1.23	-0.97	-1.33	1.08	-1.42	-1.14	-1.23	-1.74	-1.78	1.03	-1.51	-1.16	-1.22	-1.00	1.44	1.40	1.63	1.22	-1.09	-1.03	-1.13	1.07	1.22
0.29	0.39	0.90	0.85	0.08	х	0.54	0.77	0.43	0.51	0.71	0.98	0.63	0.22	0.92	0.00	0.72	0.26	0.71	0.25	0.03	0.04	0.94	0.01	0.73	0.65	0.96	0.02	0.10	0.04	0.04	0.54	0.84	0.81	0.66	0.71
×	1.36	-0.97	1.94	-1.74	×	-1.07	1.31	-1.28	1.39	-1.16	1.08	1.18	-1.19	1.35	-1.19	1.31	x	-1.01	×	×	-1.96	2.19	-1.64	-1.10	-1.30	-1.21	1.17	1.55	2.23	-1.03	1.06	1.54	-1.09	х	1.81
×	X	0.85	x	0.08	x	0.93	0.34	0.30	0.47	0.88	0.80	0.32	x	х	0.53	0.60	x	0.59	x	x	0.29	0.03		_	0.49	0.87	0.31	0.08	0.01	0.87	0.20	0.24	0.99	x	0.79
N/A	0.01	-1.02	1.58	-1.55	N/A	0.13	1.18	-1.23	0.01	-1.16			0.02	0.19	-1.26	1.20	N/A	-1.08	N/A	N/A	-1.87	1.61	-1.58	-1.13	-1.26	-1.10	1.31	1.48	1.93	0.10	-0.02		-1.11	N/A	1.52
N/A	1.35		0.37	0.19	N/A	1.20	0.13	0.05	1.38	0.00	1.05	1.19	1.21	1.16	0.07	0.12	N/A	0.06	N/A	N/A	0.09	0.58	0.06		0.04	0.11	0.14	0.08	0.30	1.13	1.08	1.29	0.02	N/A	0.29

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SPOA0302	SPOA0301	SPOA0300	SPOA0299	SPOA0298	SPOA0297	SPOA0296	SPOA0295	SPOA0294	SPOA0293	SPOA0292	SPOA0291	SPOA0290	SPOA0289	SPOA0288	SPOA0287	SPOA0286	SPOA0285	SPOA0284	SPOA0283	SPOA0282	SPOA0281	SPOA0280	SPOA0279	SPOA0278	SPOA0277	SPOA0276	SPOA0275	SPOA0274	SPOA0273	SPOA0272	SPOA0271	SPOA0270	SPOA0269	SPOA0268
					livG	livF-2		pmtA	pssA	psd				acdA-3			caiD-2									cscK	gabD-3	gabT						
phage integrase family site specific recombinase	GntR family transcriptional regulator	periplasmic branched-chain amino acid-binding protein	branched-chain amino acid ABC transporter,	permease protein annuo aciu ADC transporter,	branched-chain amino acid ABC transporter, ATP- binding protein branched chain amino acid ABC transporter	binding protein	hydantoin racemase (EC:5.1.99)	phosphatid yie thanolamine N-methyltransferase (EC:2.1.1.17)	CDP-dtacylglycerolserine O- phosphatidyltransferase (EC:2.7.8.8)	phosphatidylserine decarboxylase (EC:4.1.1.65)	branched-chain amino acid aminotransferase	hypothetical protein	AraC family transcriptional regulator	acyl-CoA dehydrogenase (EC:1.3.99.3)	acyl-CoA synthetase	ThiJ/PfpI family protein	carnitinyl-CoA dehydratase	LysR family transcriptional regulator	GMC family oxidoreductase	malonyl-CoA synthase	LysR family transcriptional regulator	TRAP dicarboxylate transporter, DctP subunit	TRAP dicarboxylate transporter, DctQ subunit	TRAP dicarboxylate transporter, DctM subunit	alpha/beta hydrolase	fructokinase (EC:2.7.1.4)	succinate-semialdehyde dehydrogenase (EC:1.2.1.16)	4-aminobutyrate aminotransferase (EC:2.6.1.19)	DNA-binding protein	glutathione-dependent formaldehyde dehydrogenase	methylamine utilization protein MauG	hypothetical protein	hypothetical protein	IcIR family transcriptional regulator
Х	-2.02	-8.77	-4.57	-4.10	-3.44	-3.20	-3.87	2.39	Х	2.06	-1.01	1.53	-2.38	15.00	35.90	6.36	8.93	-1.55	7.77	-1.03	-1.68	-4.06	-1.99	1.28	Х	-1.37	-1.56	2.27	-1.27	3.96	14.00	6.90	11.00	9.05
х	0.04	0.00	0.02	0.08	0.10	0.13	0.02	0.06	x	0.00	0.59	0.90	0.06	0.22	0.00	0.00	0.00	х	0.00	0.77	0.02	0.02	0.23	0.71	х	0.31	×	0.17	0.26	0.15	0.01	0.02	0.02	0.33
-1.11	1.02	-1.22	1.05	-1.18	-1.01	-1.16	-1.14	E	1.45	1.18	1.26	1.08	-1.25	1.47	x	1.19	1.55	1.17	-1.17	1.51	1.06	-1.33	1.13	-1.00	-1.27	1.08	1.32	1.66	-1.56	1.28	1.82	-1.04	1.36	2.09
0.60	0.88	0.21	0.25	0.53	0.99	0.88	0.86	0.69	0.08	0.21	0.26	0.83	0.38	0.79	×	0.24	0.01	0.70	×	0.13	0.73	0.12	0.17	0.96	0.60	0.54	0.03	0.24	0.04	0.67	0.04	0.96	0.45	0.17
N/A	-0.50	-5.00	-1.76	-2.64	-2.23	-2.18	-2.51	1.75	N/A	1.62	0.13	1.31	-1.82	8.24	N/A	3.78	5.24	-0.19	3.30	0.24	-0.31	-2.70	-0.43	0.14	N/A	-0.15	-0.12	1.97	-1.42	2.62	7.91	2.93	6.18	5.57
N/A	1.52	3.78	2.81	1.46	1.22	1.02	1.37	0.64	N/A	0.44	1.14	0.22	0.57	6.77	N/A	2.59	3.69	1.36	4.47	1.27	1.37	1.37	1.56	1.14	N/A	1.23	1.44	0.31	0.14	1.34	6.09	3.97	4.82	3.48
-1.41	-1.70	-26.30	-10.50	-12.50	-8.63	-7.58	-8.33	3.50	3.67	4.38	1.45	1.81	2.74	-3.62	-1.02	1.01	-1.06	-1.03	-1.17	1.64	-2.37	-6.94	-3.03	-1.27	-1.49	-1.07	-3.24	-1.09	9.33	0.99	-1.96	-2.40	-4.36	-1.88
0.97	0.06	0.01	0.00	0.02	0.06	0.09	0.02	0.01	0.01	0.00	0.04	0.77	0.04	0.53	х	0.50	0.83	0.91	0.15	0.32	0.06	0.01	0.04	0.99	0.62	0.50	0.04	0.79	0.00	0.88	0.14	0.20	0.10	0.09
-0.99	1.08	-1.21	x	-1.18	-1.23	-1.34	-1.33	-1.25	-1.85	-1.80	-1.11	1.06	-1.21	-1.08	х	1.10	1.24	-1.44	x	1.06	-1.43	-1.06	х	-1.04	-1.34	1.53	-1.53	-1.39	-1.85	1.40	-0.95	1.05	1.24	-1.23
0.29	0.54	0.59	×	0.62	0.11	0.69	0.19	0.15	0.04	0.01	0.29	0.83	0.16	0.97	х	0.23	0.27	0.59	x	0.54	0.24	0.85	Х	1.00	0.35	0.08	0.14	0.64	0.03	0.59	0.39	0.96	0.69	0.66
-1.20	-0.31	-13.76	N/A	-6.84	-4.93	-4.46	-4.83	1.13	0.91	1.29	0.17	1.44	0.77	-2.35	N/A	1.06	0.09	-1.24	N/A	1.35	-1.90	-4.00	N/A	-1.16	-1.42	0.23	-2.39	-1.24	3.74	1.20	-1.46	-0.68	-1.56	-1.56
0.21	1.39	12.55	N/A	5.66	3.70	3.12	3.50	2.38	2.76	3.09	1.28	0.38	1.98	1.27	N/A	0.05	1.15	0.21	N/A	0.29	0.47	2.94	N/A	0.12	0.08	1.30	0.86	0.15	5.59	0.20	0.50	1.73	2.80	0.33
-0.99	-1.66	1.17	-1.12	-1.19	-1.16	-1.18	-1.16	-1.50	-1.54	-1.31	1.23	1.32	-1.29	-1.32	x	-1.06	1.19	1.08	-1.09	1.14	1.25	1.06	1.67	1.20	-1.27	1.07	-1.25	-1.28	1.06	1.05	-1.43	-1.14	1.05	-1.56
0.30	0.00	0.51	0.73	0.72	0.35	0.85	0.86	0.02	0.03	0.08	0.19	0.38	0.21	0.88	×	0.13	0.05	0.82	0.74	0.06	0.60	0.71	0.13	0.73	0.39	0.67	0.27	0.74	0.49	0.97	0.08	0.87	0.88	0.16
х	-1.98	-1.06	×	-1.30	-1.28	1.06	1.23	-1.78	-3.19	-1.97	1.16	1.33	1.53	1.98	Х	-1.02	-1.04	-1.18	×	-1.21	-0.89	1.22	Х	1.23	1.13	1.32	-1.37	1.52	-1.62	1.09	-1.30	1.67	1.15	-1.23
х	0.06	0.96	×	х	x	0.92	0.70	0.13	0.01	0.02	0.16	0.48	0.09	0.70	х	1.00	0.98	0.95	х	0.12	0.73	0.47	x	Х	0.39	0.01	0.61	x	0.04	0.97	Х	0.55	0.70	0.84
N/A	-1.82	0.05	N/A	-1.25	-1.22	-0.06	0.04	-1.64	-2.37	-1.64	1.20	1.33	0.12	0.33	N/A	-1.04	0.08	-0.05	N/A	-0.04	0.18	1.14	N/A	1.22	-0.07	1.20	-1.31	0.12	-0.28	1.07	-1.37	0.27	1.10	-1.40
N/A	0.16	1.12	N/A	0.06	0.06	1.12	1.20	0.14	0.82	0.33	0.04	0.01	1.41	1.65	N/A	0.02	1.12	1.13	N/A	1.18	1.07	0.08	N/A	0.02	1.20	0.13	0.06	1.40	1.34	0.02	0.06	1.41	0.05	0.17

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SPOA0341	SPOA0340	SPOA0339	SPOA0338	SPOA0337	SPOA0336	SPOA0335	SPOA0334	SPOA0333	SPOA0332	SPOA0331	SPOA0330	SPOA0329	SPOA0328	SPOA0327	SPOA0326	SPOA0325	SPOA0324	SPOA0323	SPOA0322	SPOA0321	SPOA0320	SPOA0319	SPOA0318	SPOA0317	SPOA0316	SPOA0315	SPOA0314	SPOA0313	SPOA0312	SPOA0311	SPOA0310	SPOA0309	SPOA0306	SPOA0305	SPOA0304	SPOA0303
											eda-2	mgsA									hpcD		megL			icd									repA	
hypothetical protein	tautomerase	HAD family hydrolase	LysR family transcriptional regulator	hypothetical protein	LacI family transcription regulator	subunit	Subunit TP AD dicarboxylata family transporter DetD	TRAP dicarboxylate transporter, DctM subunit	dihydroxy-acid dehydratase (EC:4.2.1.9)	2-dehydro-3-deoxygluconokinase	2-denydro-3-deoxypnospnogluconate aldolase (EC:4,1.2,14,4,1.3,16)	methylglyoxal synthase (EC:4,2,3,3)	hypothetical protein	rzcC protein	LysR family transcriptional regulator	2-nitropropane dioxygenase	hypothetical protein	cyclic nucleotide-binding protein	hypothetical protein	glyoxalase family protein	5-carboXymetny1-2-nydroXymuconate dena- isomerase (EC:5.3.3.10)	transketolase	methionine gamma-lyase (EC:4.4.1.11)	AsnC family transcriptional regulator	amidohydrolase	ISOCITTATE denydrogenase, NADF-dependent (EC:1.1.1.42)	LysR family transcriptional regulator	trimethylamine methyltransferase family protein	glutamate-1-semialdehyde 2,1-aminomutase	FAD-dependent oxidoreductase	glycine cleavage system protein T	sulphoacetaldehyde acetyltransferase	S1 RNA-binding domain-containing protein	hypothetical protein	replication protein	replication protein
1.64	2.81	3.23	1.12	-1.70	-1.53	-1.53	-1.35	-1.56	-1.70	-0.98	-2.30	-1.14	Х	-1.70	1.68	2.05	-1.13	1.02	-1.44	-1.37	-1.30	1.39	1.10	1.71	2.02	4.68	-1.11	-1.68	х	-2.81	-1.16	1.21	1.27	1.04	1.40	1.30
0.06	0.00	0.09	0.91	0.04	0.63	0.03	0.03	0.81	×	0.73	0.02	0.48	×	0.14	0.01	0.21	0.06	0.96	0.15	0.49	0.06	0.64	0.99	0.75	0.02	0.00	0.79	×	×	x	0.73	0.40	0.30	0.96	0.26	0.18
-8.35	-6.74	-6.80	-1.42	-1.58	-1.86	-2.16	-1.93	-1.62	-1.46	-1.25	-1.51	-1.37	-1.26	-1.03	1.51	4.04	1.54	1.38	1.35	-1.25	-1.07	-1.01	-1.32	-1.11	1.12	2.12	1.01	-1.14	-1.02	-1.17	-1.09	-1.09	1.00	-1.07	-1.19	-1.20
0.02	0.01	0.01	0.55	0.18	0.13	0.09	0.04	0.68	0.60	0.66	0.46	0.10	0.09	0.81	0.02	0.00	0.05	0.22	0.07	0.19	0.95	0.99	0.66	0.38	0.57	0.05	0.98	0.72	0.98	0.86	0.54	0.28	0.96	0.77	0.10	0.69
-3.36	-1.97	-1.79	-0.15	-1.64	-1.70	-1.85	-1.64	-1.59	-1.58	-1.12	-1.91	-1.26	N/A	-1.37	1.60	3.05	0.21	1.20	-0.04	-1.31	-1.19	0.19	-0.11	0.30	1.57	3.40	-0.05	-1.41	N/A	-1.99	-1.13	0.06	1.14	-0.02	0.11	0.05
5.00	4.78	5.02	1.27	0.06	0.17	0.31	0.29	0.03	0.12	0.13	0.40	0.12	N/A	0.34	0.09	1.00	1.34	0.18	1.40	0.06	0.12	1.20	1.21	1.41	0.45	1.28	1.06	0.27	N/A	0.82	0.03	1.15	0.14	1.06	1.30	1.25
2.93	4.79	5.81	1.29	-2.18	-2.61	-3.91	-2.02	-2.05	-1.67	-1.14	-3.27	-1.39	-0.85	-1.17	-1.47	-1.60	-1.14	1.47	-1.16	-1.53	-1.38	-1.60	-1.30	-2.68	-3.73	4.28	-1.31	-1.43	-1.41	-1.49	-1.15	-1.24	2.13	1.40	1.38	1.32
0.00	0.00	0.02	0.38	0.05	0.44	0.04	0.16	0.75	0.71	0.76	0.00	0.54	0.09	0.86	0.03	0.04	0.55	0.62	0.82	0.13	0.41	0.67	0.81	0.50	0.00	0.01	0.91	0.85	0.89	0.87	0.62	0.81	0.01	0.39	0.03	0.12
-144.00	-118.00	-177.00	-1.98	-1.33	-2.64	-1.56	-2.27	-1.31	-1.90	-1.82	-1.36	-1.54	-1.06	1.10	1.09	1.43	-1.14	-1.12	-1.66	1.15	1.11	-1.08	-1.00	1.07	1.58	2.33	-1.12	-1.24	-1.12	-1.42	-1.18	1.07	-1.13	-1.07	-1.35	-1.20
) 0.00	0.00	0.00	x	0.17	0.10	0.05	0.03	0.82	0.38	0.20	0.35	0.14	0.74	0.84	0.29	0.02	0.39	0.51	0.05	0.46	0.35	0.93	1.00	0.81	0.01	0.01	0.76	0.46	0.83	x	0.21	0.45	0.26	0.68	0.03	0.73
-70.54	-56.61	-85.60	-0.35	-1.76	-2.63	-2.74	-2.15	-1.68	-1.79	-1.48	-2.32	-1.47	-0.95	-0.03	-0.19	-0.09	-1.14	0.18	-1.41	-0.19	-0.14	-1.34	-1.15	-0.81	-1.08	3.31	-1.22	-1.34	-1.27	-1.46	-1.17	-0.09	0.50	0.17	0.01	0.06
73.47	61.40	91.41	1.64	0.43	0.02	1.18	0.13	0.37	0.12	0.34	0.96	0.08	0.11	1.14	1.28	1.52	0.00	1.30	0.25	1.34	1.25	0.26	0.15	1.88	2.66	0.98	0.10	0.10	0.14	0.04	0.02	1.16	1.63	1.24	1.37	1.26
7 1.13	1.68	1.80	1.37	-1.47	-1.24	-1.19	-1.33	-1.32	-1.29	-1.45	-1.30	-1.41	-1.04	1.21	1.20	-1.56	1.10	1.10	1.02	-1.03	1.08	3.83	6.50	-1.51	-1.13	1.14	-0.98	1.33	-1.20	1.34	-1.41	-1.71	-1.09	1.31		-1.21
0.85	<sup>8</sup> 0.00	_	0.36	7 0.01	4 0.48	0.24	3 0.13	2 0.81	9 0.63	5 0.40	0.20	0.24	4 0.80	0.45	0.40	5 0.04	0.36	0.58	0.61	3 0.98	0.33	0.12	0.04	0.19	3 0.28	0.60	0.88	0.28	0.61	0.56	0.02		9 0.08			0.21
1.74	1.72	1.98	-1.14	-0.97	3 1.08	-1.10	-1.36	1.69	3 1.09	-1.05	1.16	-1.02	-1.19	1.08	1.33	1.18	-1.28	3 1.06		3 1.33	3 2.94	36.90	\$ 59.60	-1.09	3 1.26	-1.25	3 1.27	3 1.26	1.01	1.22	-1.23		-1.13			1.15
4 0.14	2 0.21	3 0.19	4 0.98	7 0.49	3 0.83	0 0.71	6 0.13	9 0.63	€ 0.71	5 0.84	5 0.65	2 0.81	9 0.68	3 0.85	3 0.24	3 0.04	8 0.21	5 0.66	5 0.03	3 X	4 0.07	0 0.01	0 0.02	9 0.76	5 0.23	5 0.10	0.79	5 0.21	X	2 X	3 0.41					0.68
4 1.44	1 1.70	9 1.89	8 0.12	9 -1.22	3 -0.08	1 -1.15	3 -1.35	3 0.19	1 -0.10	4 -1.25	5 -0.07	-1.22	8 -1.12	5 1.15	4 1.27	4 -0.19	1 -0.09	6 1.08	3 -0.47	0.15	7 2.01	1 20.37	2 33.05	6 -1.30	3 0.07	0 -0.06	9 0.14	1 1.30	-0.10	1.28	-1.32					8 -0.03
4 0.31	0.02	0.09	2 1.26	2 0.25	8 1.16	5 0.04	5 0.02	9 1.51	0 1.19	5 0.20	7 1.23	2 0.20	2 0.08	0.06	0.07	9 1.37	9 1.19	3 0.02	7 1.49	5 1.18	0.93	7 16.54	5 26.55	0 0.21	7 1.20	6 1.20	4 1.13	0.04	0 1.11	3 0.06	2 0.09	3 0.19	1 0.02			3 1.18

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### Chapter 10: Appendix

-137.00

-66.54

70.47

1.22 1.13

0.73 0.77

1.88 1.77

1.55

1.45

0.33 0.32 0.22 0.35

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2.04

179.00

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-87.75

91.26

-158.00

1.19

0.20 0.00

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120.00

-58.81 -53.81

61.20 57.20

1.46 2.27

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SPOA0380	SPOA0379	SPOA0378	SPOA0377	SPOA0376	SPOA0375	SPOA0374	SPOA0373	SPOA0372	SPOA0371	SPOA0370	SPOA0369	SPOA0368	SPOA0367	SPOA0366	SPOA0365	SPOA0364	SPOA0363	SPOA0362	SPOA0361	SPOA0360	SPOA0359	SPOA0358	SPOA0357	SPOA0356	SPOA0355	SPOA0354	SPOA0353	SPOA0352	SPOA0351	SPOA0350	SPOA0349	SPOA0348	SPOA0347	SPOA0346	SPOA0345	SPOA0344	SPOA0343	SPOA0342
										copA	copB																											
hypothetical protein	GntR family transcriptional regulator	mandelate racemase	aldehyde dehydrogenase	epoxide hydrolase	GntR family transcriptional regulator	TRAP dicarboxylate transporter, DctP subunit	TRAP dicarboxylate transporter, DctM subunit	TRAP dicarboxylate transporter, DctQ subunit	hypothetical protein	copper resistance protein A	copper resistance protein B	hypothetical protein	ABC transporter, permease protein	amino acid permease	copper-translocating P-type ATPase (EC:3.6.3.4)	hypothetical protein	hypothetical protein	hypothetical protein	cytochrome c family protein	multicopper oxidase	cytochrome c family protein	sensor histidine kinase (EC:2.7.3)	DNA-binding response regulator	hypothetical protein	hypothetical protein	aminotransferase, class III	aldehyde dehydrogenase	hypothetical protein	protein	aldo/keto reductase carboxymuconolactone decarboxylase family	LysR family transcriptional regulator	hypothetical protein	hypothetical protein					
-1.35	-1.91	-1.17	-1.64	-1.42	1.44	х	-1.45	-1.51	1.43	х	1.08	-1.10	х	-1.01	-0.94	2.13	1.15	1.41	1.21	-1.07	1.11	-1.03	-1.94	-1.07	-1.05	-1.32	-0.98	1.13	-1.96	х	-1.69	1.87	2.60	1.95	1.99	-1.31	1.63	1.39
0.49	0																																					
	0.08	0.76	x	0.79	0.34	x	0.47	0.86	0.60	Х	0.85	0.16	×	0.99	0.77	0.04	x	0.89	0.21	0.96	0.38	0.98	0.04	0.84	0.12	0.09	0.51	0.67	0.61	Х	0.01	0.02	0.04	0.07	0.16	0.44	0.54	0.75
-1.41	.08 -1.46	0.76 1.12	X 1.08	0.79 -1.41	0.34 1.18	X -1.57	0.47 -1.21	0.86 -1.10	0.60 1.09	X -1.11	0.85 -1.37	0.16 -1.48	X X	0.99 -1.20	0.77 1.01	0.04 -1.62	X -1.49	0.89 -1.12	0.21 -1.05	0.96 -1.06	0.38 -1.08	0.98 -1.19	0.04 -1.43	0.84 -1.21	0.12 -1.44	0.09 -1.42	0.51 -1.51	0.67 1.12	0.61 -1.09	X 1.05	0.01 1.44	0.02 -4.78	0.04 -4.37	0.07 -5.10	0.16 -4.87	0.44 -1.39	0.54 -9.67	0.75 -9.09
-1.41 0.33	-				-																-	-	-													-		
	-1.46	1.12	1.08	-1.41	1.18	-1.57	-1.21	-1.10	1.09	-1.11	-1.37	-1.48	х	-1.20	1.01	-1.62	-1.49	-1.12	-1.05	-1.06	-1.08	-1.19	-1.43	-1.21	-1.44	-1.42	-1.51	1.12	-1.09	1.05	1.44	-4.78	-4.37	-5.10	-4.87	-1.39	-9.67	-9.09
0.33	-1.46 0.06	1.12 X	1.08 0.86	-1.41 0.77	1.18 0.42	-1.57 0.46	-1.21 0.88	-1.10 0.96	1.09 0.71	-1.11 0.87	-1.37 0.22	-1.48 0.01	X X	-1.20 0.95	1.01 0.85	-1.62 0.01	-1.49 0.10	-1.12 0.89	-1.05 0.63	-1.06 0.96	-1.08 0.66	-1.19 0.12	-1.43 0.41	-1.21 0.17	-1.44 0.03	-1.42 0.02	-1.51 0.36	1.12 0.44	-1.09 0.86	1.05 X	1.44 0.03	-4.78 0.00	-4.37 0.01	-5.10 0.00	-4.87 0.01	-1.39 0.11	-9.67 0.01	-9.09 0.02
0.33 -1.38	-1.46 0.06 -1.69	1.12 X -0.02	1.08 0.86 -0.28	-1.41 0.77 -1.42	1.18 0.42 1.31	-1.57 0.46 N/A	-1.21 0.88 -1.33	-1.10 0.96 -1.31	1.09 0.71 1.26	-1.11 0.87 N/A	-1.37 0.22 -0.15	-1.48 0.01 -1.29	X X N/A	-1.20 0.95 -1.11	1.01 0.85 0.04	-1.62 0.01 0.26	-1.49 0.10 -0.17	-1.12 0.89 0.15	-1.05 0.63 0.08	-1.06 0.96 -1.07	-1.08 0.66 0.02	-1.19 0.12 -1.11	-1.43 0.41 -1.69	-1.21 0.17 -1.14	-1.44 0.03 -1.25	-1.42 0.02 -1.37	-1.51 0.36 -1.24	1.12 0.44 1.13	-1.09 0.86 -1.53	1.05 X N/A	1.44 0.03 -0.13	-4.78 0.00 -1.46	-4.37 0.01 -0.89	-5.10 0.00 -1.58	-4.87 0.01 -1.44	-1.39 0.11 -1.35	-9.67 0.01 -4.02	-9.09 0.02 -3.85

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0.17 0.05 N/A

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0.95 0.59 0.50

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0.94 0.72

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1.23

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0.05 0.01 0.16 N/A

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SPOA0416	SPOA0415	SPOA0414	SPOA0413	SPOA0412	SPOA0411	SPOA0410	SPOA0409	SPOA0408	SPOA0407	SPOA0406	SPOA0405	SPOA0404	SPOA0403	SPOA0402	SPOA0401	SPOA0400	SPOA0399	SPOA0398	SPOA0397	SPOA0396	SPOA0395	SPOA0394	SPOA0393	SPOA0392	SPOA0391	SPOA0390	SPOA0389	SPOA0388	SPOA0387	SPOA0386	SPOA0385	SPOA0384	SPOA0383	SPOA0382	SPOA0381
															badA-2																				
hypothetical protein	hypothetical protein	xanthine dehydrogenase tamily protein, large subunit	xanthine dehydrogenase tamily protein, small subunit	subunit	hypothetical protein	hypothetical protein	tryptophan 2,3-dioxygenase	hypothetical protein	salicylyl-CoA 5-hydroxylase (EC:1.14.13.40)	D-beta-hydroxybutyrate dehydrogenase	MarR family transcriptional regulator	enoyl-CoA hydratase (EC:4.2.1.17)	acyl-CoA dehydrogenase	cupin	benzoate-coenzyme A ligase (EC:6.2.1.25)	endoribonuclease L-PSP	R body protein RebB-like protein	R body protein RebB-like protein	hypothetical protein	hypothetical protein	hypothetical protein	hypothetical protein	R body protein RebB-like protein	RNA polymerase sigma-70 factor	cyclic nucleotide-binding protein	phosphonate monoester hydrolase	MerR family transcriptional regulator	acetyl-CoA synthetase	LysR family transcriptional regulator	GMC family oxidoreductase	hypothetical protein	protein	r	spermidine/putrescine ABC transporter, permease	spermidine/putrescine ABC transporter, periplasmic substrate-binding protein spermidine/nutrescine ABC transporter. ATP-
1.68	3.06	4.06	3.16	2.92	-1.15	-1.76	1.73	1.43	1.25	1.07	-1.67	1.52	1.24	-1.19	-1.26	1.05	1.67	1.80	1.94	2.56	2.22	х	1.56	-1.02	1.41	1.50	-1.34	-1.15	-1.76	1.14	1.13	1.00	-1.09	-1.36	-1.41
0.06	0.21	0.01	0.44	0.01	0.13	0.00	0.07	0.22	0.47	0.34	0.40	0.02	0.51	0.31	0.15	0.92	0.04	0.00	0.00	0.00	0.10	×	0.16	0.55	0.84	0.11	0.02	0.82	0.46	0.68	0.86	0.98	0.91	0.53	0.04
1.24	1.55	1.85	1.25	1.35	1.20	-1.62	1.17	1.26	-1.14	-1.10	-1.26	1.04	-1.26	-1.61	-2.25	-1.10	1.98	1.37	1.06	1.69	-1.00	1.11	1.17	-0.99	-1.08	1.00	1.15	-1.43	-1.45	-1.06	-1.27	-1.32	-1.06	-1.22	-1.74
0.42	0.28	0.05	0.34	0.04	0.25	0.24	0.29	0.27	0.26	0.30	0.31	0.83	0.74	0.25	0.14	0.72	0.08	0.22	0.86	0.06	0.98	×	0.58	0.78	0.86	0.98	0.32	0.41	0.27	0.91	0.53	0.80	0.92	0.73	0.02
1.46	2.31	2.96	2.21	2.14	0.03	-1.69	1.45	1.35	0.06	-0.02	-1.47	1.28	-0.01	-1.40	-1.76	-0.03	1.83	1.59	1.50	2.13	0.61	N/A	1.37	-1.01	0.17	1.25	-0.10	-1.29	-1.61	0.04	-0.07	-0.16	-1.08	-1.29	-1.58
0.22	0.75	1.11	0.96	0.79	1.18	0.07	0.28	0.09	1.20	1.09	0.21	0.24	1.25	0.21	0.50	1.08	0.16	0.22	0.44	0.44	1.61	N/A	0.20	0.01	1.25	0.25	1.25	0.14	0.16	1.10	1.20	1.16	0.02	0.07	0.17
-1.61	2.76	1.37	1.17	2.45	-1.13	0.68	-1.27	-1.59	-2.13	-2.10	-1.80	-1.51	-1.85	-2.99	-5.13	-2.71	-5.36	-6.80	-4.71	-3.46	-3.10	х	-4.74	-3.33	1.40	1.60	-1.42	-2.00	-1.82	-1.26	-1.58	1.07	-1.43	-1.86	-2.06
0.30	0.36	0.31	0.98	0.01	0.53	0.00	0.64	0.38	0.03	0.02	0.50	0.18	0.28	0.29	0.00	0.01	0.00	0.00	0.00	0.00	0.06	×	0.00	0.02	0.68	0.02	0.19	0.36	0.53	0.96	0.67	0.92	0.84	0.48	0.41
-1.08	1.00	1.47	1.61	1.40	-1.49	1.41	-1.10	-1.11	-1.45	-1.42	-1.01	1.09	-1.18	1.42	1.13	1.26	3.19	3.18	2.36	2.11	1.66	x	1.88	1.92	-1.34	1.07	1.20	-1.37	-1.42	1.14	-1.17	-1.13	-1.83	-1.37	-1.05
0.57	0.91	0.18	0.04	0.00	0.19	0.09	0.11	0.80	0.07	0.02	0.84	0.57	0.77	0.06	0.65	0.19	0.02	0.02	0.05	0.02	0.22	×	0.12	0.01	0.36	0.87	0.24	0.08	0.19	0.58	0.60	0.91	0.38	0.48	0.41
-1.35	1.88	1.42	1.39	1.93	-1.31	1.05	-1.19	-1.35	-1.79	-1.76	-1.41	-0.21	-1.52	-0.79	-2.00	-0.73	-1.09	-1.81	-1.18	-0.68	-0.72	N/A	-1.43	-0.71	0.03	1.34	-0.11	-1.69	-1.62	-0.06	-1.38	-0.03	-1.63	-1.62	-1.56
0.27	0.88	0.05	0.22	0.53	0.18	0.36	0.09	0.24	0.34	0.34	0.40	1.30	0.33	2.21	3.13	1.99	4.28	4.99	3.54	2.79	2.38	N/A	3.31	2.63	1.37	0.27	1.31	0.31	0.20	1.20	0.21	1.10	0.20	0.24	0.50
-1.28	1.79	2.11	1.10	1.27	1.11	0.95	-1.29	-1.01	-1.12	-1.10	-1.24	-1.02	-1.02	1.19	-1.52	1.12	1.22	-1.26	-1.79	-1.01	-1.40	Х	-1.29	-1.37	-1.06	-1.71	1.24	-1.04	1.07	1.30	-1.16	1.17	1.40	1.34	1.29
0.11	0.16	0.04	0.86	0.18	0.25	0.10	0.19	0.99	0.22	0.33	0.10	0.86	0.97	0.34	0.09	0.16	0.43	0.31	0.02	0.91	0.45	х	0.57	0.02	0.90	0.02	0.24	0.97	0.76	0.51	0.56	0.88	0.50	0.44	0.08
-1.31	1.37	1.92	1.52	1.48	-0.96	1.56	-1.42	-1.33	-1.01	-1.13	1.04	-1.07	-1.15	1.46	-1.04	1.06	-1.49	-1.37	-1.72	-1.10	1.19	×	1.24	x	1.40	-1.77	-0.98	2.34	1.15	-1.60	-1.14	1.34	1.44	1.16	1.22
0.12	0.24	0.14	0.28	0.05	0.46	0.25	0.14	0.26	0.54	0.98	0.75	0.70	0.86	0.06	0.93	0.45	0.09	0.11	0.54	0.84	0.63	x	0.13	х	0.49	0.06	0.56	0.15	0.43	x	0.95	0.79	0.57	0.66	0.24
-1.30	1.58	2.02	1.31	1.38	0.07	1.26	-1.36	-1.17	-1.07	-1.12	-0.10	-1.05	-1.09	1.33	-1.28	1.09	-0.14	-1.32	-1.76	-1.06	-0.11	N/A	-0.03	N/A	0.17	-1.74	0.13	0.65	1.11	-0.15	-1.15	1.26	1.42	1.25	1.26
0.02	0.21	0.10	0.21	0.11	1.04	0.31	0.06	0.16	0.06	0.01	1.14	0.03	0.06	0.14	0.24	0.03	1.36	0.06	0.04	0.05	1.30	N/A	1.27	N/A	1.23	0.03	1.11	1.69	0.04	1.45	0.01	0.09	0.02	0.09	0.04

SPOA0452	SPOA0451	SPOA0450	SPOA0449	SPOA0448	SPOA0447	SPOA0446	SPOA0445	SPOA0444	SPOA0443	SPOA0442	SPOA0441	SPOA0440	SPOA0439	SPOA0438	SPOA0437	SPOA0436	SPOA0435	SPOA0434	SPOA0433	SPOA0432	SPOA0431	SPOA0430	SPOA0429	SPOA0428	SPOA0427	SPOA0426	SPOA0425	SPOA0424	SPOA0423	SPOA0422	SPOA0421	SPOA0420	SPOA0419	SPOA0418	SPOA0417
				hutG				fabA				fabB	fabI-3		dehII			catD		pcaB								fabJ-2							
MarR family transcriptional regulator	MarR family transcriptional regulator	hypothetical protein	phytanoyl-CoA dioxygenase family protein	N-formylglutamate amidohydrolase (EC:3.5.1.68)	branched-chain amino acid ABC transporter ATP- binding protein	LysR family transcriptional regulator	FUR family transcriptional regulator	3-hydroxydecanoyl-ACP dehydratase (EC:4.2.1.60)	hypothetical protein	hypothetical protein	hypothetical protein	3-oxoacyl-ACP synthase (EC:2.3.1.41)	enoyl-ACP reductase (EC:1.3.1.9)	peptidyl-prolyl cis-trans isomerase, FKBP-type	haloacid dehalogenase (EC:3.8.1.2)	alpha/beta hydrolase	pyridoxal-phosphate dependent enzyme	3-oxoadipate enol-lactone hydrolase (EC:3.1.1.24)	esterase	3-carboxy-cis, cis-muconate cycloisomerase (EC:5.5.1.2)	tetracycline resistance protein	zinc-binding dehydrogenase family oxidoreductase	hypothetical protein	mandelate racemase	ArsR family transcriptional regulator	AraC family transcriptional regulator	acetyl-CoA acetyltransferase (EC:2.3.1.9)	fatty oxidation complex, alpha subunit	TetR family transcriptional regulator	hypothetical protein	pyridoxamine 5'-phosphate oxidase family protein	hypothetical protein	PhzF family phenazine biosynthesis protein	acetolactate synthase, large subunit	GntR family transcriptional regulator
-1.26	-1.31	-0.97	2.34	1.05	-1.37	-1.03	-1.73	-1.77	-1.39	-1.47	-1.84	1.80	1.55	-2.08	1.65	1.24	1.87	0.99	1.58	1.64	1.12	2.33	Х	7.01	-1.16	1.07	1.51	1.59	0.97	1.20	-1.64	1.84	1.69	-1.07	-1.72
0.04	0.03	0.98	0.08	0.50	0.72	0.99	0.03	0.08	0.03	0.37	0.34	0.03	0.13	0.00	0.75	0.74	0.57	0.01	0.30	0.02	х	0.01	x	0.07	0.02	0.28	0.72	0.06	0.32	0.44	0.39	0.16	0.03	0.88	0.20
-1.04	1.00	1.10	1.41	-1.03	-1.33	-1.19	1.19	1.09	1.12	-1.08	-1.18	1.26	1.43	-1.17	1.33	-1.15	1.56	1.40	1.07	1.40	1.05	2.83	х	1.26	-1.33	-1.08	1.07	-1.00	-1.24	-1.30	-1.33	-1.52	-1.55	-1.15	-1.17
0.81	0.92	0.75	0.07	0.84	0.61	0.86	0.08	0.80	0.38	0.81	0.67	0.61	0.31	0.15	0.42	0.11	0.04	0.15	0.89	0.03	0.58	0.01	х	0.23	0.17	0.32	0.82	0.96	0.30	0.25	0.53	0.05	0.03	0.45	0.28
-1.15	-0.16	0.07	1.88	0.01	-1.35	-1.11	-0.27	-0.34	-0.14	-1.28	-1.51	1.53	1.49	-1.63	1.49	0.05	1.72	1.20	1.33	1.52	1.09	2.58	N/A	4.14	-1.25	-0.01	1.29	0.30	-0.13	-0.05	-1.49	0.16	0.07	-1.11	-1.45
0.11	1.16	1.04	0.47	1.04	0.02	0.08	1.46	1.43	1.26	0.20	0.33	0.27	0.06	0.46	0.16	1.20	0.16	0.20	0.25	0.12	0.04	0.25	N/A	2.88	0.09	1.08	0.22	1.30	1.11	1.25	0.16	1.68	1.62	0.04	0.28
-1.35	-1.21	-1.25	-1.18	-2.54	-1.81	1.13	-1.55	0.81	-1.46	-3.48	1.51	1.08	1.21	2.08	1.38	-1.72	1.52	1.43	1.16	2.38	3.05	-1.14	х	1.25	-1.19	1.64	-1.93	-1.71	1.47	-1.35	-2.02	-1.46	-1.86	1.11	-2.35
0.03	0.02	0.90	0.87	0.02	0.67	0.42	0.01	0.46	0.06	0.02	0.74	0.52	0.39	0.04	0.22	0.19	0.16	0.20	0.99	0.02	0.00	0.12	×	0.70	0.03	0.05	0.03	0.04	0.08	0.63	0.68	0.44	0.03	0.76	0.21
1.04	1.27	1.11	1.10	-1.31	-1.04	- 1.46	1.07	1.01	1.49	1.02	1.62	1.16	0.99	1.73	1.61	1.24	1.35	1.10	1.49	1.09	-1.34	2.35	Х	1.44	1.15	-1.56	1.32	1.40	1.22	-1.03	1.07	-1.04	1.05	-1.66	-1.48
0.95	0.36	0.10	0.53	0.30	0.96	0.53	0.69	0.69	0.04	x	0.41	0.57	0.53	0.02	0.09	0.55	0.24	0.91	0.26	0.84	0.04	0.00	х	0.12	0.44	0.07	0.14	0.17	0.02	0.90	0.68	0.75	0.78	0.06	0.06
-0.16	0.03	-0.07	-0.04	-1.93	-1.43	-0.17	-0.24	0.91	0.02	-1.23	1.57	1.12	1.10	1.91	1.50	-0.24	1.44	1.27	1.33	1.74	0.86	0.61	N/A	1.35	-0.02	0.04	-0.31	-0.16	1.35	-1.19	-0.48	-1.25	-0.41	-0.28	-1.92
1.20	1.24	1.18	1.14	0.62	0.39	1.30	1.31	0.10	1.48	2.25	0.06	0.04	0.11	0.18	0.12	1.48	0.09	0.16	0.16	0.65	2.20	1.75	N/A	0.10	1.17	1.60	1.63	1.56	0.13	0.16	1.55	0.21	1.46	1.39	0.44
1.13	1.17	-1.15	1.76	-1.54	-1.24	1.04	1.70	1.13	1.23	-1.62	-1.13	1.23	1.50	1.10	1.05	-1.38	1.37	2.07	1.23	1.43	-1.12	1.06	х	1.28	1.47	-1.09	1.55	1.26	1.03	-1.53	-1.54	1.18	-1.25	-1.15	-1.48
0.51	0.32	0.31	0.04	0.10	0.76	0.75	0.00	0.48	0.03	0.17	0.82	0.40	0.02	0.10	0.75	0.15	0.09	0.01	0.46	0.29	0.19	0.56	х	0.28	0.01	0.45	0.06	0.03	0.94	0.13	0.17	0.21	0.28	0.36	0.11
1.33	1.61	-1.40	2.31	-1.44	1.09	-0.96	1.41	1.21	1.43	1.05	-1.10	1.14	0.99	-1.05	1.15	1.70	-1.01	1.07	0.95	-1.35	1.12	1.72	Х	1.44	1.03	-1.32	2.15	1.69	1.13	-1.42	1.09	1.83	1.20	-1.34	-1.70
0.59	0.09	0.10	0.04	0.43	0.86	0.81	0.08	0.36	0.02	x	0.97	0.98	0.36	0.36	0.35	0.17	0.96	0.50	0.50	0.03	0.49	0.05	х	0.07	0.89	0.09	0.05	0.03	0.47	0.64	0.51	0.06	0.18	0.64	0.04
1.23	1.39	-1.28	2.04	-1.49	-0.08	0.04	1.56	1.17	1.33	-0.29	-1.12	1.19	1.25	0.03	1.10	0.16	0.18	1.57	1.09	0.04	0.00	1.39	N/A	1.36	1.25	-1.21	1.85	1.48	1.08	-1.48	-0.23	1.51	-0.03	-1.25	-1.59
0.10	0.22	0.13	0.27	0.05	1.17	1.00	0.15	0.04	0.10	1.34	0.01	0.05	0.26	1.08	0.05	1.54	1.19	0.50	0.14	1.39	1.12	0.33	N/A	0.08	0.22	0.12	0.30	0.21	0.05	0.06	1.32	0.33	1.23	0.10	0.11

are highlighted in yellow. Due to the normalisation procedure, some genes have an "X" label in the fold-change columns, and N/A indicates where a error. Up- or down-regulated gene expression values are highlighted in green or red, respectively. P values < 0.05 (and thus, statistically significant) P-value was absent following normalisation. each of the two duplicate arrays, for each inducer, is given, followed by the P value (P). The calculated mean is provided, along with the standard the gene (compared to uninduced cells) in the different co-inducers; 5 mM DMSP (Dp), 2.5 mM acrylate (A), 5 mM DMS (Ds). The fold change for Columns show the gene number (SPO), name (if applicable), function of the gene product (as predicted by KEGG) and fold change in expression of