

1 **Sensitivity of UK Butterflies to local climatic extremes:**

2 **Which life stages are most at risk?**

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14  
15 **Abstract**

- 16 1. There is growing recognition as to the importance of extreme climatic events  
17 (ECEs) in determining changes in species populations. In fact it's often the  
18 extent of climate variability that determines a population's ability to persist at a  
19 given site.
- 20 2. This study examined the impact of ECEs on the resident UK butterfly species  
21 (n=41) over a 37 year period. The study investigated the sensitivity of  
22 butterflies to four extremes (Drought, Extreme Precipitation, Extreme Heat,  
23 Extreme Cold), identified at the site level, across each species' life stages.  
24 Variations in the vulnerability of butterflies at the site level were also

25 compared based on 3 life history traits (voltinism, habitat requirement, and  
26 range).

27 3. This is the first study to examine the effects of ECEs at the site level across all  
28 life stages of a butterfly, identifying sensitive life stages and unravelling the  
29 role life history traits play in species sensitivity to ECEs.

30  
31 4. Butterfly population changes were found to be primarily driven by temperature  
32 extremes. Extreme heat was detrimental during overwintering periods and  
33 beneficial during adult periods and extreme cold had opposite impacts on both  
34 of these life stages. Previously undocumented detrimental effects were  
35 identified for extreme precipitation during the pupal life stage for univoltine  
36 species. Generalists were found to have significantly more negative  
37 associations with ECEs than specialists.

38 5. With future projections of warmer, wetter winters and more severe weather  
39 events, UK butterflies could come under severe pressure given the findings of  
40 this study.

41 **Key-words** Butterfly population changes, climate change, life history traits, linear  
42 mixed effects model, sensitivity

## 43 **Introduction**

44 Climate change is causing direct and substantial changes to biodiversity and to  
45 entire ecosystems (Cramer *et al.* 2014); species have been altering their growth,  
46 phenology, and distribution (Root *et al.* 2003; Møller, Rubolini & Lehikoinen 2008;  
47 Chen *et al.* 2011). While species are changing their distribution in an attempt to track  
48 the climatic conditions optimal for their survival, i.e. their climatic niche, their ability to  
49 do so is often limited. Some species are lagging behind the high velocity of climate

50 change (Loarie *et al.* 2009; Bertrand *et al.* 2011; Devictor *et al.* 2012) resulting in  
51 range contractions (Foden *et al.* 2007). Both widespread and range restricted  
52 species are projected to have range losses and/or increased extinction risks as a  
53 result of changes in mean climate (IPCC 2007; Warren 2011; Foden *et al.* 2013;  
54 Warren *et al.* 2013).

55 Most attribution of climate change impacts on biodiversity (Parmesan, Root & Willig  
56 2000; Root *et al.* 2003; Chen *et al.* 2011; Doney *et al.* 2012), and the projection of  
57 future impacts (Pereira *et al.* 2010; Bellard *et al.* 2012; Pacifici *et al.* 2015), is based  
58 upon the observed or projected change in mean climate, however the impacts of  
59 climatic extremes, such as heatwaves, heavy rainfall, and droughts are much less  
60 frequently studied and the rate and magnitude of these events is likely to increase in  
61 the future (IPCC 2012; Jones *et al.* 2014).

62 Extreme climate events (ECEs) have been shown to directly affect species  
63 populations by influencing reproductive and mortality rates (Jiguet, Brotons &  
64 Devictor 2011). Changes in climate variability, as a result of climate change, leading  
65 to changes in the magnitude and frequency of ECEs may be more important for  
66 determining whether a species can persist in a given location, than are modest  
67 increases in average temperature (Parmesan *et al.* 2000; Bauerfeind & Fischer  
68 2014).

69 Butterflies have been used to demonstrate ecological examples of species'  
70 responses to climate change (Parmesan *et al.* 1999; Warren *et al.* 2001; Wilson *et*  
71 *al.* 2005; Franco *et al.* 2006; Thomas, Franco & Hill 2006; Pöyry *et al.* 2009;  
72 Diamond *et al.* 2011) and due to their ectothermic characteristics are a good  
73 taxonomic group to look at effects of extreme climatic events. ECEs, such as  
74 drought and heavy precipitation events, have been shown to be detrimental to the

75 survival of butterflies, causing local extinction events (McLaughlin *et al.* 2002; Oliver  
76 *et al.* 2015) which highlights the importance of incorporating these ECEs in  
77 ecological studies (Easterling *et al.* 2000; Jentsch & Beierkuhnlein 2008; Smith 2011;  
78 Fischer, Klockmann & Reim 2014). Warmer, wetter winters have been negatively  
79 associated with changes in population growth rates as has heavy rainfall (Pollard  
80 1988; WallisDeVries, Baxter & Van Vliet 2011).

81 Univoltine and multivoltine species are under different selective pressures due to  
82 differing numbers and timings of life stages. Life stage can be incorporated into the  
83 analysis to allow identification of sensitive stages within a butterfly's lifecycle to  
84 particular extremes (WallisDeVries *et al.* 2011; Radchuk, Turlure & Schtickzelle  
85 2013).

86 Impacts of ECEs can be examined at a large scale (Pollard 1988; Roy *et al.* 2001;  
87 WallisDeVries *et al.* 2011) or take into account site specific information to avoid  
88 hiding population losses in one area due to gains in another (Wilbanks & Kates  
89 1999). By analysing the impacts of ECEs at site level these losses and gains can be  
90 unmasked, allowing for attributions to be identified that may not have been in a  
91 broader scale study (Pearce-Higgins 2011; Newson *et al.* 2014). Site specific  
92 differences may be a function of a species' local site adaption to regional climate  
93 variables (Ayres & Scriber 1994) and habitat availability and characteristics also  
94 affect species responses to ECEs. Oliver *et al.* (2015) showed that reducing habitat  
95 fragmentation was effective at countering negative drought effects on butterfly  
96 populations and reducing landscape-scale habitat fragmentation may influence a  
97 species ability to withstand weather-mediated population declines (Newson *et al.*  
98 2014).

99 ECEs have been defined using specific arbitrary thresholds (WallisDeVries *et al.*  
100 2011), such as extreme heat being anything above 30°C. This only identifies heat as  
101 an issue during the summer, excluding the possibility that heat may also play a role  
102 during other periods of the year and other stages of a species' life cycle.  
103 This study takes a new approach to identifying species responses to extremes,  
104 accounting for both the life stage and site specific effects thus providing a more  
105 dynamic and biologically relevant approach in identifying climatic extremes for an  
106 organism. This study aims to assess the impacts of ECEs on UK species over the 37  
107 year period from 1976- 2012. This study will (i) examine the influence of ECEs on  
108 butterfly population change over a 37 year period; (ii) determine which butterfly life  
109 stages are sensitive to which ECEs and (iii) determine whether butterfly population  
110 changes are more associated with extremes of temperature or precipitation?

## 111 **Materials and Methods**

### 112 ***The Datasets***

#### 113 ***The butterfly dataset – UKBMS***

114 Site level butterfly population indices were obtained from The UK Butterfly Monitoring  
115 Scheme (UKBMS), a comprehensive dataset for UK Butterflies consisting of records  
116 from thousands of volunteers across the UK. This data covers a period from 1976  
117 (38 monitored sites) to 2012 (878 monitored sites). In total over the 37 year period  
118 there have been 1,802 different recording sites. At monitored sites, weekly counts of  
119 adult butterflies were made over a 26 week period between the beginning of April  
120 and the end of September on fixed routes provided the weather conditions were  
121 favourable for butterfly activity (Pollard & Yates 1993). This procedure is repeated  
122 yearly allowing for comparisons between years at that particular site but also  
123 between sites. Full details of the sampling methodology can be found in (Pollard,

124 Hall & Bibby 1986). Population indices are based upon all generations that fall within  
125 the recording period, the indices are not split by generation.

126 Species with fewer than 10 sites and/or less than 15 years of data were removed  
127 from the analysis as in (WallisDeVries *et al.* 2011) limiting the number of species  
128 included in the analyses to 41 of the 59 regularly occurring UK butterflies. A separate  
129 model was created for each species to account for different lifecycle timings,  
130 numbers of generations and overwintering strategies.

131 Information on life history traits (voltinism: univoltine / multivoltine, species range:  
132 Northern range limited / widespread species, habitat generalist / habitat specialist  
133 species) were collated using (Asher *et al.* 2001).

134

#### 135 ***The weather observations dataset***

136 Daily maximum, minimum temperature and precipitation data on a 0.25 degree  
137 regular lat/long grid were obtained from the E-OBS dataset for the UK between 1950  
138 and 2012 (Haylock *et al.* 2008). Site specific daily data was extracted using the  
139 latitude and longitude of the survey sites from the UKBMS dataset. For more  
140 information on how the data is interpolated into its gridded format see (Haylock *et al.*  
141 2008).

#### 142 ***Identification of Extreme Weather Events and their biological relevance***

143 Calendar dates were identified for all life stages of each butterfly (Ovum, Larvae,  
144 Pupae, Adult and Overwintering) according to their phenology (Eeles 2014).  
145 Overwintering period was set as a fixed period for all species (WallisDeVries *et al.*  
146 2011), starting on the 1<sup>st</sup> of November and finishing on the 28<sup>th</sup> of February. The  
147 phenology of each species can vary from year to year in addition to the site to site  
148 variation (Van Strien *et al.* 2008; WallisDeVries *et al.* 2011). In this study we use

149 fixed phenology dates for the butterflies to identify the start and end of each lifecycle  
150 for 37 years of data which the UKBMS covers.

151 Once the phenologies of each life stage for each species were identified, the climate  
152 data set was used to detect and extract any extreme climate events occurring during  
153 each life stage for each species at each site based on all 63 years covered by the  
154 climate data. Four types of ECEs were defined using site and species-specific  
155 thresholds, and the number of days exceeding that threshold was calculated  
156 (WallisDeVries *et al.* 2011), Table 1. Two standard deviations was chosen to set the  
157 extremes for temperature (Beaumont *et al.* 2011) and the 97.5 percentile to set  
158 extremes for precipitation as they were hypothesised to identify temperatures and  
159 precipitation beyond the climatic norm for species in each area. This was carried out  
160 at the site level over the 63 year period covered by the E-OBS dataset. All extremes  
161 were defined as the number of days exceeding the threshold criteria identified by the  
162 above methods for a given butterfly's life cycle stage.

163 The ECE definitions adopted give more flexibility, biological application and meaning  
164 in relation to time of the year and location of the extreme impacts than arbitrary  
165 thresholds. Each extreme is tailored specifically to each individual species. In  
166 addition to this it accounts for the historical climate a species has experienced at a  
167 given site for a given life stage. Arbitrary thresholds of temperature, such as 30°C  
168 used in previous studies, limit our capacity to understand how temperature may  
169 affect life stages that do not fall during the hottest periods of the year. This study  
170 uses site and species specific life stage climatic extremes enabling an understanding  
171 of how extremes occurring in different stages of the life cycle may impact on  
172 population change.

173

174 **Statistical Analysis**

175 **Species-specific models**

176 Species-specific linear mixed models were built which relate the annual adult  
177 butterfly abundance of a particular species to the ECEs previously identified for the  
178 different stages of that butterfly species' life cycle: ovum, larva, pupa, adult,  
179 (repeating in multivoltine species) and overwintering period. These models assess  
180 the impacts that identified extremes during each butterfly's life stages had on the  
181 butterfly's adult population across the UK. The dependent variable was chosen as  
182 the log of the indices of adult abundance from one year to the next and was used  
183 rather than just the indices for adult abundance in order to satisfy model  
184 assumptions of normality. The log transformation has been used as in similar studies  
185 (Roy *et al.* 2001; WallisDeVries *et al.* 2011) to account for the varying numbers of  
186 butterflies present at a site (Freeman 2009). Site was included as a random variable  
187 (Mair *et al.* 2014) to account for site specific adaptation between different  
188 populations of the same species due to issues such as habitat differences amongst  
189 sites. Counts of the number of ECEs identified for the different stages of that butterfly  
190 species' life cycle: ovum, larva, pupa, adult, and overwintering period were  
191 incorporated as fixed explanatory variables. Backwards stepwise selection using  
192 Akaike's Information Criterion (AIC) as recommended by (Thiele 2012) was used to  
193 remove variables that don't explain the variation in butterfly populations. Due to the  
194 possibility that several models may fit our data suitably well, the Pdredge function in  
195 the *MuMIn* package in R statistical software was used to dredge for all the possible  
196 model options using the variables selected for by the backwards stepwise selection.  
197 Any model with a  $\Delta$  AIC of less than 4 was deemed similar to the best fit model and  
198 was incorporated in the model averaging which has been increasingly backed and

199 applied in similar studies and is recommended for prediction and forecasting (Thiele  
200 2012).

### 201 ***Combined univoltine and multivoltine models***

202 Linear models were created by separating univoltine from multivoltine species and  
203 combining all species in each group to run a combined model for univoltine and  
204 multivoltine species. It displays the differences in response of the butterflies based  
205 on their voltinism. It also helps to understand the relative importance of variables  
206 found as being significant in the individual species models when looking at them  
207 from a univoltine and multivoltine perspective. The relative importance of each  
208 variable within the combined models was assessed using the package *relaimpo*  
209 (Grömping 2006) in R and defined as the percentage contribution of each predictor  
210 to the  $R^2$  of the model. It allows us to give statistical support relevance to counts of  
211 variables gained from species-specific models.

### 212 ***Life history traits sensitivity to ECEs comparison: Welch t tests.***

213 Welch t tests were used to make comparisons between species with different life  
214 history traits and their response to ECEs. Comparisons were based on the mean  
215 percentage of negative responses in relation to total number of possible variables  
216 from the individual species models when divided and grouped based on their life  
217 history traits.

## 218 **Results**

### 219 ***Which life stages are affected by which ECEs?***

220 The percentage of species for which an extreme affected a certain life stage varied  
221 depending on voltinism. Thus results are presented for univoltine and multivoltine  
222 species separately. All quoted percentages in the results for species affected are  
223 based on significant effects in the individual species models.

224 ***Univoltine Species***

225 The adult and overwintering life stages are the most sensitive for 29 univoltine  
226 species (Fig 1.). Extreme heat during the overwintering life stage and extreme cold  
227 during the adult life stage are the most frequently occurring negative extreme  
228 variables both causing population declines (affecting 45% and 35% of species  
229 respectively). Adult and overwintering life stages have opposing population  
230 responses to temperature extremes, extreme heat during the adult life stage is  
231 causes positive population change for 21% of species, while during overwintering it  
232 is associated with negative population change in 45% of species. Another extremely  
233 important variable to which univoltine species are vulnerable to is extreme  
234 precipitation during the pupal life stage affecting 28% of species. Drought appears to  
235 impact on the adult stage most negatively, 24% of the species, but appears to be  
236 beneficial during the ovum life stage also for 24% of species which is shown in the  
237 combined species model to be more importance for univoltine butterfly population  
238 change than its negative impacts, Table. 2. The combined model, including all  
239 univoltine species, identifies which of the variables from the species specific models  
240 to focus on when considering response of univoltine species. The first 5 variables  
241 account for 73.6% of the predictive power of the combined model (Table. 2).  
242 Extreme heat in the overwintering stage and precipitation in the pupal stage have  
243 strong negative effects on univoltine butterfly population trends. Extreme heat in the  
244 adult and pupal life stage drive positive population change in univoltine species. In  
245 summary, univoltine species seem particularly sensitive to temperature extremes at  
246 both ends of the scale (Heat or Cold) and it is the adult and overwintering phases  
247 that are vulnerable to these extremes. In addition to this, extreme precipitation during  
248 the pupal life stage is a detrimental driver of population change in a number of  
249 univoltine species.

250 ***Multivoltine Species***

251 Extreme heat during overwintering and extreme precipitation during 1<sup>st</sup> and 2<sup>nd</sup>  
252 generation adult life stages are the most frequently occurring extreme variables  
253 causing population declines in multivoltine species (67%, 58% and 50% of all  
254 multivoltine species affected respectively, Figure 1). As in univoltine species, adult  
255 and overwintering life stages have opposite population responses to temperature  
256 extremes. Extreme heat during the adult life stage is associated with positive  
257 population change in 42% of species. Drought plays a much more important role in  
258 multivoltine species than univoltine species. Drought negatively affects 50% of  
259 species during their 2<sup>nd</sup> larval life stage but has a positive impact on 25% of the  
260 species during their 1<sup>st</sup> ovum life stage. In the model combining all multivoltine  
261 species, the 9 most important variables account for 73% of the predictive power of  
262 the combined multivoltine model (Table 3). The multivoltine model is clearly driven  
263 by extremes of temperature, five were extremes in heat and one a cold extreme.  
264 Unlike univoltine species however, multivoltine seem to be susceptible across all life  
265 stages with ovum, larvae, pupae, adult and overwintering all being represented in the  
266 nine most important variables in the combined model. Species' vulnerability to  
267 extremes appears to be most prominent in the 1st generation and is primarily driven  
268 by exposure to extreme heat with the exception of the negative impacts of  
269 precipitation during the adult stage. Multivoltine species have a significantly higher  
270 proportion of negative responses to ECEs across their life stages than univoltine  
271 species ( $t_{(25)}=-2.86$ ,  $p=0.008$ ), Table 4. The results suggest that multivoltine species  
272 are more sensitive to extremes than univoltine species.

273 Within univoltine species there is no significant difference in the number of negative  
274 responses when comparing specialist with generalist species ( $t_{(20)}=-1.6$ ,  $p=0.122$ )

275 Table 4.

276 There is no significant difference between widespread and northern range limited  
277 species nested in univoltine species, ( $t_{(20)}= 1.69$ ,  $p=0.102$ ) Table 4. However when  
278 nested in multivoltine species, widespread species show more responses to  
279 extremes across their life stage than northern range limited species ( $t_{(8)}=3.76$ ,  
280  $p=0.004$ ) Table 4.

## 281 **Discussion**

282 UK butterfly populations are influenced by extreme climatic events. Extreme  
283 temperature events play a significant role in determining the population changes in  
284 species from year to year in both multivoltine and univoltine species. Previous  
285 studies found that cold weather during the adult phase negatively affect population  
286 change, while warm weather has positive associations to population (Calvert,  
287 Zuchowski & Brower 1983; Roy *et al.* 2001; Warren *et al.* 2001; WallisDeVries *et al.*  
288 2011). The benefit of heat on butterfly populations is to be expected given their  
289 poikilothermic nature. This study examined the effects of extreme temperature and  
290 precipitation variables on all butterfly life stages, for both univoltine and multivoltine  
291 species. For UK butterflies the overwintering stage was found to be particularly  
292 sensitive to extremes. Butterfly populations are negatively affected by hotter  
293 temperatures while overwintering and benefit from colder winters. This concurs with  
294 previous studies such as (Radchuk *et al.* 2013; Oliver *et al.* 2015) who found in their  
295 laboratory experiments that the overwintering larval stage was extremely sensitive to  
296 increases in temperature. This study identified negative associations of high  
297 temperatures during the overwintering stage but did not find that this sensitivity was

298 confined to species overwintering in their larval stage. Radchuk *et al.* (2013) argue  
299 that elevated temperatures during the overwintering period increase rates of  
300 mortality due to increased incidences of disease and fungi both of which are more  
301 abundant in milder winters (Harvell 2002). Whilst this may be the case, we  
302 hypothesise that in the case of butterflies overwintering as larvae or adults it may be  
303 due to extreme hot temperatures acting as a cue for butterflies or their larvae to  
304 come out from overwintering too early, decoupling from photoperiod cues, (Wiklund,  
305 Lindfors & Forsberg 1996) and subsequently killed off by temperatures returning to  
306 colder conditions or potentially the destruction of their food plant due to similar  
307 mechanisms (McLaughlin *et al.* 2002).

308 This study did not account for annual variation in butterfly phenology (Van Strien *et*  
309 *al.* 2008), the life stage periods were fixed based on the average of the last 37 years  
310 thus life stage exposure to extremes may have been less well quantified in years or  
311 sites with advanced or delayed phenology. Overall our approach is likely to be robust  
312 since it accounts site variability (by including the effects of climatic extremes at the  
313 site level), and includes a long-term data set (37 years) to quantify country wide  
314 species population responses to ECEs. These results should not be extrapolated  
315 beyond the UK due to issues such as local adaptation, it is prudent to expect  
316 potential differences in the responses of continental European populations of the  
317 same butterflies.

### 318 ***Single generation vs multi-generation species***

319 All life stages for univoltine species showed sensitivity to ECEs during the  
320 overwintering stage, with extreme cold events being beneficial and extreme heat  
321 detrimental on butterfly populations. One of the more prominent and consistent  
322 negative contributors to univoltine species' population change is precipitation events  
323 during the pupal and larval periods. This is an important finding as it hasn't been

324 identified in previous studies but would be expected from heavy rainfall events  
325 (Pollard 1988). Indeed, Hill *et al.* (2003) have previously hypothesised the potential  
326 importance of precipitation having a detrimental impact on both the larval and pupal  
327 stage, which is clearly supported by our analysis of univoltine species. The impacts  
328 of drought are difficult to interpret in this study as species do not seem to respond as  
329 uniformly to this extreme as the other extremes. However, during the ovum life stage  
330 our combined species models have indicated it plays an important and significant  
331 role in determining increases in population size.

332 It would appear that univoltine species prefer warmer, drier climates outside of winter  
333 periods. Current predictions forecast that the UK will have a warmer climate with  
334 drier summers (Jenkins *et al.* 2009) which on the face of it would seem to benefit  
335 most univoltine species however this may not be the case as warmer, wetter winters  
336 could potentially be a driving force behind many population changes as in (Radchuk  
337 *et al.* 2013).

338 Temperature extremes are the primary driving factor when analysing the impact of  
339 ECEs on multivoltine butterfly populations. As in the univoltine species, hot weather  
340 during overwintering period is negative with extreme cold being beneficial. The adult  
341 stage is extremely sensitive to extremes in temperature but primarily the second  
342 generation stage, Table 3. This is probably due to the timing of the second  
343 generation for most multivoltine species, which have their flight period during  
344 summer. Temperature has been shown to be extremely important during these  
345 summer periods (Roy *et al.* 2001). Similar to the univoltine species, multivoltine  
346 appear to be positively impacted by drought conditions during the 1<sup>st</sup> generation  
347 ovum and adult stages. This apparent benefit of drought may indicate that the levels  
348 of drought identified in this study are not at a level that is detrimental to butterflies.

349 Our analysis shows that univoltine species are less sensitive to ECEs than  
350 multivoltine species. These results need to be interpreted with caution taking into  
351 account the small number of multivoltine (n=12) species included in the analysis.  
352 This may be a due to exposure to extremes during more life stages, more  
353 generations in a year may put more selection pressures on a species. (Radchuk *et*  
354 *al.* 2013) emphasise the importance of a resource based habitat approach and it is  
355 clear that more life stages would put more selection pressures on the species or  
356 potentially due to the fact that an extreme in one year can affect two consecutive  
357 generations when life stages overlap.

### 358 ***Generalists vs specialists***

359 Generalist species have more significant negative associations with ECEs than  
360 specialist species. This suggest that ECEs may affect population change in  
361 generalist species, especially in populations on the edge of their climatic range  
362 (Hellmann *et al.* 2008), while population change of habitat specialists species is  
363 controlled by other factors (e.g. habitat loss and degradation) (Warren *et al.* 2001).  
364 We hypothesise that generalist species are more vulnerable as they are filling their  
365 climatic niche and hence many populations within the species range may be situated  
366 on the climatic range edge and be more vulnerable to increased climate variability  
367 outside of their comfort zone. In contrast specialist species are confined to particular  
368 host plants which may not ubiquitous across the specialist species' climatic niche,  
369 hence those specialist species are not filling their climatic niche and are effectively in  
370 or close to their core range and are not subjected to ECEs that are outside their  
371 ability to adapt and cope. It is also possible that specialist species are being buffered  
372 by their habitats where they have been able to persist (Oliver, Brereton & Roy 2013).

373 ***Widespread vs Northern range limited species***

374 No significant difference in the number of negative associations between widespread  
375 and northern range limited species was found when nested within univoltine species.  
376 The opposite was found for multivoltine species with widespread species having  
377 significantly more negative associations when nested in multivoltine species. These  
378 results need to be interpreted with caution as mentioned previously. If validated this  
379 result may indicate that widespread species may be subjected to a much higher  
380 variation in climatic conditions than northern range limited species and as such may  
381 be subject to temperatures and precipitation levels that are detrimental.

382 ***Conclusion***

383 This study has identified a hitherto unknown sensitivity of univoltine species to  
384 extreme precipitation during their pupal life stage. In addition, this study although  
385 using novel ECE definitions, found an agreement with previous studies, indicating  
386 that warm and even climatically extreme hot summers are beneficial to butterfly  
387 populations, while extremely wet cold summers are detrimental to their populations.  
388 The detrimental effect of extreme heat during overwintering has been evidenced  
389 previously but fewer studies have shown the sensitivity of the pupal stage to extreme  
390 precipitation events and warrants further attention. Interestingly the perceived  
391 sensitivity of butterflies to drought (Oliver *et al.* 2015) was not evidenced in our  
392 analysis but this could be due to limitations in our definition of drought.  
393 Sensitivity to ECEs in butterflies was primarily dominated by temperature extremes  
394 which would support our hypothesis that butterfly population changes are more  
395 dependent on heat extremes as shown by both the combined species models and  
396 the proportion of species affected in the species specific models. This study has  
397 identified scope for future work. An interesting augmentation of this study would be  
398 to identify dramatic species decline events and examine the extent to which they are

399 associated with ECEs. Finally, building on the work of (Oliver *et al.* 2015), further  
400 analysis is warranted on the ability of habitats to buffer extremes other than drought  
401 that have been identified as being detrimental by this study. Extreme wind could be  
402 factored into future studies also. Unfortunately, the appropriate data was not  
403 available through the weather sources used in this paper.

404 The novel identification of the sensitivity of the pupal life stage to extreme  
405 precipitation supports our decision to address the impacts of extremes at a finer  
406 scale than previous studies and has also shown the importance of looking at ECEs  
407 across all life stages given these relatively new findings.

408 This study has shown that butterflies could potentially benefit from increasing  
409 temperatures in the UK in the future but warmer and wetter winters and increases in  
410 severe weather events that have also been predicted (Defra 2009; Jenkins *et al.*  
411 2009) could be detrimental to the survival of many of its butterfly species and further  
412 research is needed regarding the balance of importance that these variables could  
413 have and whether the benefits of warmer summers will be outweighed by the  
414 detrimental winter effects. Based on the results of this study, future conservation  
415 efforts hoping to mitigate against ECEs in the future should focus their efforts on the  
416 adult and overwintering life stages of UK butterflies.

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418

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428 (<http://www.ecad.eu>).

429

### 430 **Data accessibility**

431 Weather data (E-OBS dataset) available from

432 <http://www.ecad.eu/download/ensembles/download.php> (Haylock *et al.* 2008).

433 The UKBMS (Butterfly) database is managed and maintained by the Biological

434 Records Centre, based at the Centre for Ecology & Hydrology (CEH). Access to

435 population indices available from the CEH Data catalogue

436 <http://doi.org/10.5285/378f0f77-1842-4789-ba15-6fbdf7d02299> (Botham *et al.* 2016).

### 437 **References:**

438 Asher, J., Warren, M., Fox, R., Harding, P., Jeffcoate, G. & Jeffcoate, S. (2001) *The*  
439 *Millenium Atlas of Butterflies in Britain and Ireland*. Oxford University Press,  
440 Oxford.

441 Ayres, M.P. & Scriber, J.M. (1994) Local adaptation to regional climates in *Papilio*  
442 *canadensis* (Lepidoptera: Papilionidae). *Ecological Monographs*, **64**, 465–482.

443 Bauerfeind, S.S. & Fischer, K. (2014) Simulating climate change: Temperature  
444 extremes but not means diminish performance in a widespread butterfly.  
445 *Population Ecology*, **56**, 239–250.

446 Beaumont, L.J., Pitman, A., Perkins, S., Zimmermann, N.E. & Yoccoz, N.G. (2011)  
447 Impacts of climate change on the world 's most exceptional ecoregions.  
448 *Proceedings of the National Academy of Sciences of the United States of*  
449 *America*, **108**, 2306–2311.

450 Bellard, C., Bertelsmeier, C., Leadley, P., Thuiller, W. & Courchamp, F. (2012)  
451 Impacts of climate change on the future of biodiversity. *Ecology Letters*, **15**,  
452 365–377.

453 Bertrand, R., Lenoir, J., Piedallu, C., Riofrío-Dillon, G., de Ruffray, P., Vidal, C.,  
454 Pierrat, J.-C.J.-C., Gegout, J.-C., Riofrío-Dillon, G. & Gégout, J.-C. (2011)  
455 Changes in plant community composition lag behind climate warming in lowland

- 456 forests. *Nature*, **479**, 517–20.
- 457 Botham, M., Roy, D.B., Brereton, T. & Middlebrook, I. (2016) United Kingdom  
458 Butterfly Monitoring Scheme: site indices data 2015.
- 459 Calvert, W., Zuchowski, W. & Brower, L. (1983) The Effect of Rain, Snow and  
460 Freezing Temperatures on Overwintering Monarch Butterflies in Mexico.  
461 *Biotropica*, **15**, 42–47.
- 462 Chen, I., Hill, J.K., Ohlemüller, R., Roy, D.B. & Thomas, C.D. (2011) Rapid range  
463 shifts of species of climate warming. *Science*, **333**, 1024–1026.
- 464 Cramer, W., Yohe, G.W., Auffhammer, M., Huggel, C., Molau, U., da Silva Dias, M.  
465 a. F., Solow, a., Stone, D. a. & Tibig, L. (2014) Detection and Attribution of  
466 Observed Impacts. In: Climate Change 2014: Impacts, Adaptation and  
467 Vulnerability. Part A: Global and Sectoral Aspects. Contribution of Working  
468 Group II to the Fifth Assessment Report of the Intergovernmental Panel on  
469 Climate Ch. *Climate Change 2014: Impacts, Adaptation and Vulnerability.*  
470 *Contribution of Working Group II to the Fifth Assessment Report of the*  
471 *Intergovernmental Panel on Climate Change*, p. Cambridge University Press,  
472 Cambridge, UK and New York, NY, USA.
- 473 Defra. (2009) Adapting to climate change UK Climate Projections. *Uk Climate*  
474 *Projections*, 52.
- 475 Devictor, V., van Swaay, C., Brereton, T., Brotons, L., Chamberlain, D., Heliölä, J.,  
476 Herrando, S., Julliard, R., Kuussaari, M., Lindström, Å., Reif, J., Roy, D.B.,  
477 Schweiger, O., Settele, J., Stefanescu, C., van Strein, A., van Turnhout, C.,  
478 Vermouzek, Z., WallisDeVries, M.F., Wynhoff, I. & Jiguet, F. (2012) Differences  
479 in the climatic debts of birds and butterflies at a continental scale. *Nature climate*  
480 *change*, **2**, 121–124.
- 481 Diamond, S., Frame, A., Martin, R. & Buckley, L. (2011) Species' traits predict  
482 phenological responses to climate change in butterflies. *Ecology*, **92**, 1005–  
483 1012.
- 484 Diaz, H.F. & Murnane, R.J. (2008) *Climate Extremes and Society* (eds HF Diaz and  
485 RJ Murnane). Cambridge University Press.
- 486 Doney, S.C., Ruckelshaus, M., Duffy, J.E., Barry, J.P., Chan, F., English, C.A.,  
487 Galindo, H.M., Grebmeier, J.M., Hollowed, A.B., Knowlton, N., Polovina, J.,  
488 Rabalais, N.N., Sydeman, W.J. & Talley, L.D. (2012) Climate change impacts on  
489 marine ecosystems. *Ann Rev Mar Sci*, **4**, 11–37.
- 490 Easterling, D.R., Meehl, G.A., Parmesan, C., Changnon, S.A., Karl, T.R. & Mearns,  
491 L.O. (2000) Climate Extremes: Observations, Modeling, and Impacts. *Science*,  
492 **289**, 2068–2074.
- 493 Eeles, P. (2014) UK Butterflies - Phenologies. URL  
494 [http://www.ukbutterflies.co.uk/species\\_phenologies.php](http://www.ukbutterflies.co.uk/species_phenologies.php) [accessed 4 February  
495 2015]
- 496 Fischer, K., Klockmann, M. & Reim, E. (2014) Strong negative effects of simulated  
497 heat waves in a tropical butterfly. *The Journal of experimental biology*, **217**,  
498 2892–8.
- 499 Foden, W.B., Butchart, S.H.M., Stuart, S.N., Vié, J.-C., Akçakaya, H.R., Angulo, A.,  
500 Devantier, L.M., Gutsche, A., Turak, E., Cao, L., Donner, S.D., Katariya, V.,

501 Bernard, R., Holland, R. a, Hughes, A.F., O'Hanlon, S.E., Garnett, S.T.,  
502 Sekercioğlu, C.H. & Mace, G.M. (2013) Identifying the World's Most Climate  
503 Change Vulnerable Species: A Systematic Trait-Based Assessment of all Birds,  
504 Amphibians and Corals. *PloS one*, **8**, e65427.

505 Foden, W., Midgley, G.F., Hughes, G., Bond, W.J., Thuiller, W., Hoffman, M.T.,  
506 Kaleme, P., Underhill, L.G., Rebelo, A. & Hannah, L. (2007) A changing climate  
507 is eroding the geographical range of the Namib Desert tree Aloe through  
508 population declines and dispersal lags. *Diversity and Distributions*, **13**, 645–653.

509 Franco, A.M.A., Hill, J.K., Kitschke, C., Collingham, Y.C., Roy, D.B., Fox, R.,  
510 Huntley, B. & Thomas, C.D. (2006) Impacts of climate warming and habitat loss  
511 on extinctions at species' low-latitude range boundaries. *Global Change Biology*,  
512 **12**, 1545–1553.

513 Freeman, S. (2009) Towards a method for the estimation and use of averaged multi-  
514 species trends , as indicators of patterns of change in butterfly populations.  
515 *Centre for Ecology and Hydrology*.

516 Grömping, U. (2006) Relative Importance for Linear Regression in R : The Package  
517 relaimpo. *Journal of Statistical Software*, **17**.

518 Harvell, C.D. (2002) Climate Warming and Disease Risks for Terrestrial and Marine  
519 Biota. *Science*, **296**, 2158–2162.

520 Haylock, M.R., Hofstra, N., Klein Tank, a. M.G., Klok, E.J., Jones, P.D. & New, M.  
521 (2008) A European daily high-resolution gridded data set of surface temperature  
522 and precipitation for 1950-2006. *Journal of Geophysical Research Atmospheres*,  
523 **113**, D20119.

524 Hellmann, J.J., Pelini, S.L., Prior, K.M. & Dzurisin, J.D.K. (2008) The response of two  
525 butterfly species to climatic variation at the edge of their range and the  
526 implications for poleward range shifts. *Oecologia*, **157**, 583–592.

527 Hill, J.K., Hamer, K.C., Dawood, M.M., Tangah, J. & Chey, V.K. (2003) Rainfall but  
528 not selective logging affect changes in abundance of a tropical forest butterfly in  
529 Sabah, Borneo. *Journal of Tropical Ecology*, **19**, 35–42.

530 IPCC. (2007) *Climate Change 2007: Impacts, Adaptation and Vulnerability.*  
531 *Contribution of Working Group II to the Fourth Assessment Report of the*  
532 *Intergovernmental Panel on Climate Change* (eds ML Parry, OF Canziani, JP  
533 Palutikof, CE Hanson, and PJ van der Linden). Cambridge University Press,  
534 Cambridge.

535 IPCC. (2012) *2012: Managing the Risks of Extreme Events and Disasters to*  
536 *Advance Climate Change Adaptation. A Special Report of Working Groups I and*  
537 *II of the Intergovernmental Panel on Climate Change [Field, C.B., V. Barros,*  
538 *T.F. Stocker, D. Qin, D.J. Dokken, K.L. Eb* (eds CB Field, V Barros, TF Stocker,  
539 and Q Dahe). Cambridge University Press, Cambridge, UK and New York, NY,  
540 USA.

541 Jenkins, G.J., Murphy, J.M., Sexton, D.M.H., Lowe, J. a., Jones, P. & Kilsby, C.G.  
542 (2009) *UK Climate Projections: Briefing Report*. Exeter, UK.

543 Jentsch, A. & Beierkuhnlein, C. (2008) Research frontiers in climate change: Effects  
544 of extreme meteorological events on ecosystems. *Comptes Rendus -*  
545 *Geoscience*, **340**, 621–628.

- 546 Jiguet, F., Brotons, L. & Devictor, V. (2011) Community responses to extreme  
547 climatic conditions. *Current Zoology*, **57**, 406–413.
- 548 Jones, M.R., Blenkinsop, S., Fowler, H.J. & Kilsby, C.G. (2014) Objective  
549 classification of extreme rainfall regions for the UK and updated estimates of  
550 trends in regional extreme rainfall. *International Journal of Climatology*, **34**, 751–  
551 765.
- 552 Loarie, S.R., Duffy, P.B., Hamilton, H., Asner, G.P., Field, C.B. & Ackerly, D.D.  
553 (2009) The velocity of climate change. *Nature*, **462**, 1052–1055.
- 554 Mair, L., Hill, J.K., Fox, R., Botham, M., Brereton, T. & Thomas, C.D. (2014)  
555 Abundance changes and habitat availability drive species' responses to climate  
556 change. *Nature Climate Change*, **4**, 127–131.
- 557 McLaughlin, J.F., Hellmann, J.J., Boggs, C.L. & Ehrlich, P.R. (2002) Climate change  
558 hastens population extinctions. *Proceedings of the National Academy of  
559 Sciences of the United States of America*, **99**, 6070–6074.
- 560 Møller, A.P., Rubolini, D. & Lehikoinen, E. (2008) Populations of migratory bird  
561 species that did not show a phenological response to climate change are  
562 declining. *Proceedings of the National Academy of Sciences of the United  
563 States of America*, **105**, 16195–16200.
- 564 Newson, S.E., Oliver, T.H., Gillings, S., Crick, H.Q.P., Morecroft, M.D., Duffield, S.J.,  
565 Macgregor, N. a. & Pearce-Higgins, J.W. (2014) Can site and landscape-scale  
566 environmental attributes buffer bird populations against weather events?  
567 *Ecography*, **37**, 872–882.
- 568 Oliver, T.H., Brereton, T. & Roy, D.B. (2013) Population resilience to an extreme  
569 drought is influenced by habitat area and fragmentation in the local landscape.  
570 *Ecography*, **36**, 579–586.
- 571 Oliver, T.H., Marshall, H.H., Morecroft, M.D., Brereton, T., Prudhomme, C. &  
572 Huntingford, C. (2015) Interacting effects of climate change and habitat  
573 fragmentation on drought-sensitive butterflies. *Nature Climate Change*, **5**, 1–6.
- 574 Pacifici, M., Foden, W.B., Visconti, P., Watson, J.E.M., Butchart, S.H.M., Kovacs,  
575 K.M., Scheffers, B.R., Hole, D.G., Martin, T.G., Akçakaya, H.R., Corlett, R.T.,  
576 Huntley, B., Bickford, D., Carr, J. a., Hoffmann, A. a., Midgley, G.F., P., P.-K.,  
577 Pearson, R.G., Williams, S.E., Willis, S.G., Young, B. & Rondinini, C. (2015)  
578 Assessing species vulnerability to climate change. *Nature Climate Change*, **5**,  
579 215–225.
- 580 Parmesan, C., Root, T.L. & Willig, M.R. (2000) Impacts of Extreme Weather and  
581 Climate on Terrestrial Biota\*. *Bulletin of the American Meteorological Society*,  
582 **81**, 443.
- 583 Parmesan, C., Ryrholm, N., Stefanescu, C., Hill, J.K., Thomas, C.D., Descimon, H.,  
584 Huntley, B., Kaila, L., Kullberg, J., Tammaru, T., Tennent, W.J., Thomas, J. a &  
585 Warren, M. (1999) Poleward shifts in geographical ranges of butterfly species  
586 associated with regional warming. *Nature*, **399**, 579–583.
- 587 Pearce-Higgins, J.W. (2011) Modelling conservation management options for a  
588 southern range-margin population of Golden Plover *Pluvialis apricaria*  
589 vulnerable to climate change. *Ibis*, **153**, 345–356.
- 590 Pereira, H.M., Leadley, P.W., Proença, V., Alkemade, R., Scharlemann, J.P.W.,

- 591 Fernandez-Manjarrés, J.F., Araújo, M.B., Balvanera, P., Biggs, R., Cheung,  
592 W.W.L., Chini, L., Cooper, H.D., Gilman, E.L., Guénette, S., Hurtt, G.C.,  
593 Huntington, H.P., Mace, G.M., Oberdorff, T., Revenga, C., Rodrigues, P.,  
594 Scholes, R.J., Sumaila, U.R. & Walpole, M. (2010) Scenarios for global  
595 biodiversity in the 21st century. *Science*, **330**, 1496–1501.
- 596 Pollard, E. (1988) Temperature, rainfall and butterfly numbers. *Journal of Applied*  
597 *Ecology*, **25**, 819–828.
- 598 Pollard, E., Hall, M.. & Bibby, T.. (1986) *Monitoring the Abundance of Butterflies*  
599 *1976-1985*.
- 600 Pollard, E. & Yates, T. (1993) *Monitoring Butterflies for Ecology and Conservation*.
- 601 Pöyry, J., Luoto, M., Heikkinen, R.K., Kuussaari, M. & Saarinen, K. (2009) Species  
602 traits explain recent range shifts of Finnish butterflies. *Global Change Biology*,  
603 **15**, 732–743.
- 604 Radchuk, V., Turlure, C. & Schtickzelle, N. (2013) Each life stage matters: The  
605 importance of assessing the response to climate change over the complete life  
606 cycle in butterflies. *Journal of Animal Ecology*, **82**, 275–285.
- 607 Renwick, A.R., Massimino, D., Newson, S.E., Chamberlain, D.E., Pearce-Higgins,  
608 J.W. & Johnston, A. (2012) Modelling changes in species' abundance in  
609 response to projected climate change. *Diversity and Distributions*, **18**, 121–132.
- 610 Root, T.L., Price, J.T., Hall, K.R., Schneider, S.H., Rosenzweig, C. & Pounds, J.A.  
611 (2003) Fingerprints of global warming on wild animals and plants. *Nature*, **421**,  
612 57–60.
- 613 Roy, D.B., Rothery, P., Moss, D., Pollard, E. & Thomas, J.A. (2001) Butterfly  
614 numbers and weather: Predicting historical trends in abundance and the future  
615 effects of climate change. *Journal of Animal Ecology*, **70**, 201–217.
- 616 Smith, M.D. (2011) An ecological perspective on extreme climatic events: A synthetic  
617 definition and framework to guide future research. *Journal of Ecology*, **99**, 656–  
618 663.
- 619 Van Strien, A.J., Plantenga, W.F., Soldaat, L.L., Van Swaay, C.A.M. &  
620 WallisDeVries, M.F. (2008) Bias in phenology assessments based on first  
621 appearance data of butterflies. *Oecologia*, **156**, 227–235.
- 622 Thiele, J. (2012) Potential of GLMM in modelling invasive spread. *CAB Reviews:*  
623 *Perspectives in Agriculture, Veterinary Science, Nutrition and Natural*  
624 *Resources*, **7**, 1–10.
- 625 Thomas, C.D., Franco, A.M. a & Hill, J.K. (2006) Range retractions and extinction in  
626 the face of climate warming. *Trends in Ecology and Evolution*, **21**, 415–416.
- 627 WallisDeVries, M.F., Baxter, W. & van Vliet, A.J.H. (2011) Beyond climate  
628 envelopes: Effects of weather on regional population trends in butterflies.  
629 *Oecologia*, **167**, 559–571.
- 630 Warren, R. (2011) The role of interactions in a world implementing adaptation and  
631 mitigation solutions to climate change. *Philosophical transactions. Series A,*  
632 *Mathematical, physical, and engineering sciences*, **369**, 217–241.
- 633 Warren, M., Hill, J.K., Thomas, J.A., Asher, J., Fox, R., Huntley, B., Roy, D.B., Telfer,  
634 M.G., Jeffcoate, S., Harding, P., Jeffcoate, G., Willis, S.G., Greatorex-Davies,  
635 J.N.N., Moss, D. & Thomas, C.D. (2001) Rapid responses of British butterflies to

- 636 opposing forces of climate and habitat change. *Nature*, **414**, 65–69.
- 637 Warren, R., VanDerWal, J., Price, J., Welbergen, J. a., Atkinson, I., Ramirez-  
638 Villegas, J., Osborn, T.J., Jarvis, A., Shoo, L.P., Williams, S.E. & Lowe, J.  
639 (2013) Quantifying the benefit of early climate change mitigation in avoiding  
640 biodiversity loss. *Nature Climate Change*, **3**, 1–5.
- 641 Wiklund, C., Lindfors, V. & Forsberg, J. (1996) Early male emergence and  
642 reproductive phenology of the adult overwintering butterfly *Gonepteryx rhamni* in  
643 Sweden. *Oikos*, **75**, 227–240.
- 644 Wilbanks, T.J. & Kates, R.W. (1999) Global change in local places: How scale  
645 matters. *Climatic Change*, **43**, 601–628.
- 646 Wilson, R.J., Gutiérrez, D., Gutiérrez, J., Martínez, D., Agudo, R. & Monserrat, V.J.  
647 (2005) Changes to the elevational limits and extent of species ranges  
648 associated with climate change. *Ecology Letters*, **8**, 1138–1146.
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**Tables**

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Table 1 Extreme Climatic Events (ECEs) included in this study and their definitions (Diaz & Murnane 2008; Beaumont *et al.* 2011)

Extreme	Definition
<b>Extreme Heat</b>	Number of days above 2 standard deviations above the mean daily maximum temperature for the life cycle period of the species in question at a particular site
<b>Extreme Cold</b>	As for extreme heat but 2 standard deviations below the mean of the minimum daily temperature
<b>Drought</b>	15 days with a combined total of less than 0.02 mm of rain with each day on top of this being counted as an extra day of drought
<b>Extreme Precipitation</b>	Number of days above the 97.5 percentile for rainfall during the life cycle period in question for a particular species at that particular site. 2 standard deviations were not used in this case due to the shape of precipitation data (non-normal).

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654 Table 2 Significant variables obtained from the combined univoltine species linear model. Bonferroni corrections applied and variables ordered  
 655 by relative importance in the model using the *relaimpo* package. Variables bolded show a negative relationship with univoltine populations.

Univoltine Species			Estimate	Std. Error	t value	p-value	Relative Importance
Variable							
<b>Extr. Heat</b>	<b>during</b>	<b>Overwintering</b>	<b>-0.064</b>	<b>0.004</b>	<b>-17.681</b>	<b>&lt;0.0001</b>	<b>19.93%</b>
Extr. Heat	during	Adult stage	0.052	0.005	11.068	<0.0001	17.54%
Extr. Heat	during	Pupal stage	0.040	0.005	8.309	<0.0001	14.24%
<b>Extr. Precipitation</b>	<b>during</b>	<b>Pupal stage</b>	<b>-0.051</b>	<b>0.004</b>	<b>-12.915</b>	<b>&lt;0.0001</b>	<b>12.74%</b>
Drought	during	Ovum stage	0.044	0.004	11.365	<0.0001	9.14%
<b>Extr. Cold</b>	<b>during</b>	<b>Adult stage</b>	<b>-0.040</b>	<b>0.004</b>	<b>-10.593</b>	<b>&lt;0.0001</b>	<b>4.93%</b>
<b>Extr. Precipitation</b>	<b>during</b>	<b>Larval stage</b>	<b>-0.026</b>	<b>0.004</b>	<b>-6.476</b>	<b>&lt;0.0001</b>	<b>3.99%</b>
Drought	during	Pupal stage	0.031	0.004	7.259	<0.0001	3.96%
Extr. Cold	during	Overwintering	0.030	0.004	8.104	<0.0001	3.96%
<b>Extr. Heat</b>	<b>during</b>	<b>Ovum stage</b>	<b>-0.023</b>	<b>0.005</b>	<b>-4.560</b>	<b>&lt;0.0001</b>	<b>2.79%</b>
<b>Extr. Precipitation</b>	<b>during</b>	<b>Adult stage</b>	<b>-0.009</b>	<b>0.004</b>	<b>-2.399</b>	<b>0.0165</b>	<b>2.01%</b>
<b>Extr. Precipitation</b>	<b>during</b>	<b>Ovum stage</b>	<b>-0.019</b>	<b>0.004</b>	<b>-5.031</b>	<b>&lt;0.0001</b>	<b>1.98%</b>
<b>Extr. Heat</b>	<b>during</b>	<b>Larval stage</b>	<b>-0.017</b>	<b>0.005</b>	<b>-3.308</b>	<b>0.0009</b>	<b>1.38%</b>
<b>Drought</b>	<b>during</b>	<b>Adult stage</b>	<b>-0.011</b>	<b>0.004</b>	<b>-2.663</b>	<b>0.0077</b>	<b>0.74%</b>
<b>Extr. Precipitation</b>	<b>during</b>	<b>Overwintering</b>	<b>-0.015</b>	<b>0.004</b>	<b>-3.954</b>	<b>0.0001</b>	<b>0.69%</b>

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Table 3 Significant variables obtained from the combined multivoltine species linear model. Bonferroni corrections applied and variables ordered by relative importance in the model using the *relaimpo* package. Variables bolded show a negative relationship with univoltine populations.

Multivoltine Species							
	Variable		Estimate	Std. Error	t value	Pr(> t )	Relative Importance
Extr. Heat	during 2nd generation	Adult stage	0.105	0.006	17.921	<0.001	14.81%
Drought	during 1st generation	Adult stage	0.076	0.006	13.599	<0.001	8.45%
Extr. Cold	during 2nd generation	Larval stage	0.083	0.005	15.740	<0.001	8.31%
<b>Extr. Heat</b>	<b>during</b>	<b>Overwintering</b>	<b>-0.100</b>	<b>0.007</b>	<b>-14.427</b>	<b>&lt;0.001</b>	<b>8.22%</b>
Extr. Heat	during 2nd generation	Ovum stage	0.064	0.006	11.262	<0.001	7.82%
Drought	during 1st generation	Ovum stage	0.086	0.005	16.283	<0.001	7.12%
<b>Extr. Heat</b>	<b>during 1st generation</b>	<b>Pupal stage</b>	<b>-0.066</b>	<b>0.006</b>	<b>-10.533</b>	<b>&lt;0.001</b>	<b>6.59%</b>
<b>Extr. Heat</b>	<b>during 1st generation</b>	<b>Ovum stage</b>	<b>-0.034</b>	<b>0.006</b>	<b>-5.253</b>	<b>&lt;0.001</b>	<b>6.33%</b>
<b>Extr. Precipitation</b>	<b>during 1st generation</b>	<b>Adult stage</b>	<b>-0.050</b>	<b>0.006</b>	<b>-8.701</b>	<b>&lt;0.001</b>	<b>5.48%</b>
Extr. Cold	during	Overwintering	0.080	0.006	13.284	<0.001	4.25%
<b>Extr. Precipitation</b>	<b>during 2nd generation</b>	<b>Ovum stage</b>	<b>-0.018</b>	<b>0.006</b>	<b>-2.849</b>	<b>0.004</b>	<b>2.98%</b>
<b>Extr. Precipitation</b>	<b>during 2nd generation</b>	<b>Larval stage</b>	<b>-0.027</b>	<b>0.007</b>	<b>-3.813</b>	<b>0.000</b>	<b>2.88%</b>
<b>Extr. Cold</b>	<b>during 2nd generation</b>	<b>Ovum stage</b>	<b>-0.042</b>	<b>0.005</b>	<b>-7.846</b>	<b>&lt;0.001</b>	<b>2.28%</b>
<b>Drought</b>	<b>during 2nd generation</b>	<b>Larval stage</b>	<b>-0.053</b>	<b>0.007</b>	<b>-7.992</b>	<b>&lt;0.001</b>	<b>1.80%</b>
Drought	during 2nd generation	Ovum stage	0.016	0.006	2.400	0.016	1.69%
<b>Drought</b>	<b>during</b>	<b>Overwintering</b>	<b>-0.031</b>	<b>0.005</b>	<b>-5.700</b>	<b>&lt;0.001</b>	<b>1.61%</b>
<b>Extr. Cold</b>	<b>during 1st generation</b>	<b>Pupal stage</b>	<b>-0.052</b>	<b>0.005</b>	<b>-9.946</b>	<b>&lt;0.001</b>	<b>1.44%</b>
<b>Extr. Heat</b>	<b>during 1st generation</b>	<b>Adult stage</b>	<b>-0.021</b>	<b>0.006</b>	<b>-3.468</b>	<b>0.001</b>	<b>1.38%</b>
<b>Extr. Precipitation</b>	<b>during 1st generation</b>	<b>Pupal stage</b>	<b>-0.036</b>	<b>0.006</b>	<b>-6.144</b>	<b>&lt;0.001</b>	<b>1.37%</b>
<b>Extr. Precipitation</b>	<b>during 1st generation</b>	<b>Larval stage</b>	<b>-0.032</b>	<b>0.005</b>	<b>-6.089</b>	<b>&lt;0.001</b>	<b>1.37%</b>
<b>Extr. Cold</b>	<b>during 2nd generation</b>	<b>Adult stage</b>	<b>-0.023</b>	<b>0.005</b>	<b>-4.526</b>	<b>&lt;0.001</b>	<b>1.29%</b>
<b>Extr. Cold</b>	<b>during 1st generation</b>	<b>Adult stage</b>	<b>-0.031</b>	<b>0.005</b>	<b>-5.788</b>	<b>&lt;0.001</b>	<b>0.62%</b>
Extr. Precipitation	during 2nd generation	Pupal stage	0.027	0.006	4.280	<0.001	0.61%
<b>Drought</b>	<b>during 2nd generation</b>	<b>Adult stage</b>	<b>-0.027</b>	<b>0.006</b>	<b>-4.370</b>	<b>&lt;0.001</b>	<b>0.51%</b>
Extr. Precipitation	during	Overwintering	0.012	0.006	2.183	0.029	0.32%
Drought	during 2nd generation	Pupal stage	0.014	0.007	2.106	0.035	0.25%

659

660 Table 4 Welch T tests results *comparing the mean percentage of negative responses in relation to total number of possible*  
 661 *variables from the individual species models when divided based on their life history traits.*

Life history Group (Traits being tested tested)	t Statistic	Degrees of freedom	Means (% vs %)	<i>p-value</i>
Voltinism (Univoltine versus Multivoltine)	-2.86	25.66	(13.62 vs 22.22)	0.008
Requirement (Specialist versus Generalist)	-3.00	35.99	(10.95 vs 19.81)	0.004
Within Univoltine Species (Widespread versus Northern Range limited)	1.69	25.57	(17.5, 11.25)	0.102
Within Multivoltine Species (Widespread versus Northern Range limited)	3.76	8.77	(26.98 vs 15.56)	0.005

662

663 **Figure Legends**

664 Figure 1 Percentage of species, from the species specific models, for each life stage which there was a significant ( $p < 0.05$ ) positive  
665 or negative relationship with an Extreme Climatic Event (ECE) related to temperature or precipitation. Univoltine (A and B) and  
666 multivoltine (C and D) species are shown separately. Impact of temperature extremes (A and C) and precipitation extremes (B and  
667 D) on univoltine and multivoltine species are also shown separately. Columns above the 0 line in the y axis indicate the % of  
668 species positively impacted by ECEs while below indicates the % of species positively impacted by ECEs.

# Figures

