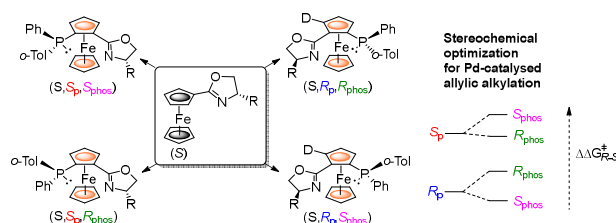


Stereoselective Synthesis of all Possible Phosferrox Ligand Diastereoisomers Displaying Three Elements of Chirality: Stereochemical Optimization for Asymmetric Catalysis

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Supporting Information Placeholder



ABSTRACT: All four possible diastereoisomers of phosphinoferrocyloxazoline (Phosferrox type) ligands containing three elements of chirality were synthesized as single enantiomers. The S_c configured oxazoline moiety ($R = \text{Me}, i\text{-Pr}$) was used to control the generation of planar chirality by lithiation, with the alternative diastereoisomer formed by use of a deuterium blocking group. In each case subsequent addition of PhPCl_2 followed by $o\text{-TolMgBr}$ resulted in a single P -stereogenic diastereoisomer (S_c,S_p,S_{phos} and S_c,R_p,R_{phos} respectively). The alternative diastereoisomers were formed selectively by addition of $o\text{-TolPCl}_2$ followed by PhMgBr (S_c,S_p,R_{phos} and S_c,R_p,S_{phos} respectively). Preliminary application of these four ligand diastereoisomers, together with (S_c,S_p) and (S_c,R_p) Phosferrox (PPh_2), to palladium catalysed allylic alkylation of *trans*-1,3-diphenylallyl acetate revealed a stepwise increase/decrease in ee, with the configuration of the matched/matched diastereoisomer as S_c,S_p,S_{phos} (97% ee).

INTRODUCTION

Metal-based asymmetric catalysis is a major branch of organic chemistry that is exploited widely for the synthesis of research-focused and economically important compounds.¹ To these ends the discovery of a new metal-catalysed reaction is frequently followed by work aimed at extending the reaction to the formation of enantioenriched chiral products. This process typically requires the screening of many chiral non-racemic ligands to achieve the goal of high enantioselectivity.² The choice of ligands screened may be guided by the success of related chemistry, and/or ligand availability, many of which may be sourced commercially. Certain ligand types have proven to be successful in a number of different reactions, and as a result these have been described as 'privileged'.³ Once identified as such many analogous ligands, based on the same core structure, are frequently synthesized by substituent variation. In this way multiple ligands closely related to, for example, BINAP,⁴ Josiphos⁵ and Walphos⁶ are known, and many are commercially available (Figure 1).

Although specific advantages have been identified for the use of C_2 -symmetric ligands in asymmetric catalysis (*e.g.* BINAP),⁷ C_1 -symmetric ligands have also found widespread

use. These frequently contain two elements of chirality (*e.g.* Josiphos and Walphos) such that two diastereoisomers are potentially available. Typically only one of the two is usually explored, this being primarily a consequence of synthetic availability.⁸ In cases where both diastereomeric ligands are tested in a metal-catalysed reaction, the matched and mismatched chirality pairings are usually identified readily.⁹

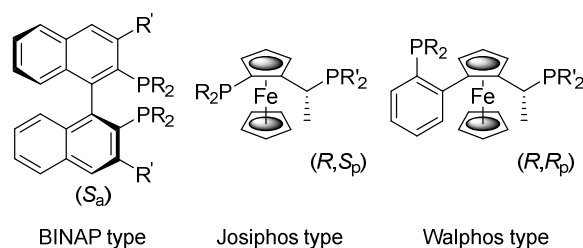


Figure 1. Representative 'privileged' and commercially available ligands for which multiple variation of the substituents (R/R') has been demonstrated.

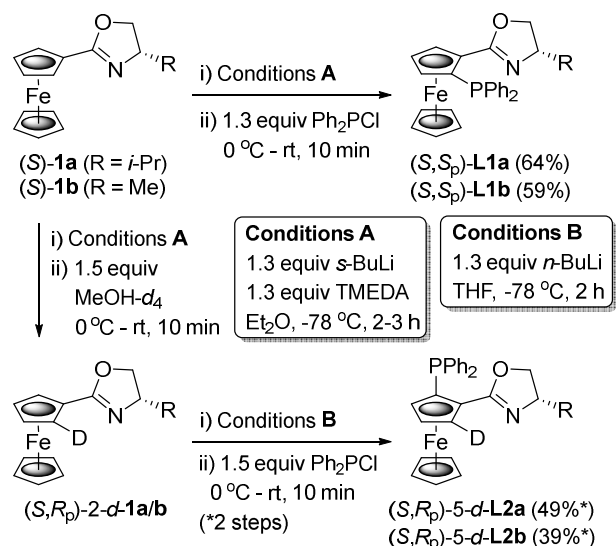
This basis of ligand selection may potentially be expanded to the use of all *four* possible diastereoisomers arising from a ligand containing three elements of chirality. The

configuration of each of these will likely have a significant influence on product enantioselectivity, and the optimum relative configuration may well be reaction dependent. This has the potential to provide an alternative pathway to ligand optimization in asymmetric catalysis. To the best of our knowledge such methodology has not been described previously.¹⁰

RESULTS AND DISCUSSION

Our approach to achieving the selective synthesis of all four possible enantiopure diastereoisomers of a chiral bidentate ligand started with ferrocenyloxazoline (*S*)-**1a** (R = *i*-Pr). This is known to undergo highly diastereoselective lithiation (dr > 100 : 1),¹¹ such that subsequent addition of Ph₂PCL results in bidentate ligand (*S,S*)-**L1a** (Scheme 1).¹² This ligand has been applied extensively in metal-catalysed asymmetric synthesis (Pd, Ru, Ir, Cu, Ag, Ni).¹³ We recently reported the adaptation of this lithiation/Ph₂PCL quench procedure for the selective synthesis of the corresponding diastereomeric ligand. Specifically, initial selective lithiation is followed by introduction of deuterium to give (*S,R_p*)-2-*d*-**1a**,¹⁴ such that a second lithiation, under non-selective conditions, is instead controlled by the introduced isotope ($k_H/k_D \sim 20$). Following addition of Ph₂PCL the target product (*S,R_p*-5-*d*-**L2a**) is obtained predominantly as one diastereoisomer (dr = 10 : 1 – Scheme 1).¹⁵ Corresponding ligands (*S,S*)-**L1b** and (*S,R_p*-5-*d*-**L2b**) have been obtained in the same way from ferrocenyloxazoline (*S*)-**1b** (R = Me).^{12c,15}

Scheme 1. Oxazoline auxiliary mediated control of planar chiral diastereoselectivity (*S,S_p*), and adaption by use of a deuterium blocking group (*S,R_p*).

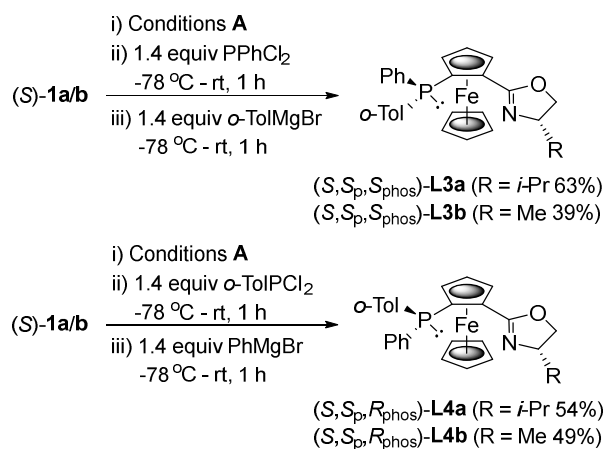


In order to extend this chemistry for the introduction of a third element of chirality we considered procedures for the selective creation of a phosphorus-based stereogenic center. Although many methods are known for the stereocontrolled synthesis of *P*-stereogenic compounds,¹⁶ in this instance efficiency would result if the lithiated planar chiral precursor to phosphorus introduction could be used to control diastereoselectivity. There are several reports in the literature of the reaction of chiral nucleophiles with RPX₂ (X =

Cl, NEt₂) where differential replacement of both leaving groups X leads to a product in high diastereoselectivity.¹⁷ A further example is the use of lithiated planar chiral Ugi's amine as the nucleophile for the first step, followed by addition of an aryl Grignard reagent, to give *P*-stereogenic *P*-N ligands in high dr.¹⁸

Accordingly, diastereoselective lithiation at -78 °C of (*S*)-**1a** (conditions A) was followed by addition of PPhCl₂ and warming of the reaction mixture to room temperature. After re-cooling to -78 °C, subsequent addition of *o*-TolMgBr, followed by warming of the reaction mixture to room temperature and work-up, resulted in the isolation of a new Phosferrox ligand. After purification by column chromatography and recrystallisation, the product was identified as (*S,S_p*,*S_{phos}*)-**L3a** following determination of the X-ray crystal structure (Scheme 2, Figure 2).¹⁹ Use of the same methodology, but instead with *o*-TolPCL₂ and PhMgBr as reagents, resulted in the formation of the alternative *P*-stereogenic diastereoisomer (*S,S_p*,*R_{phos}*)-**L4a**, an outcome again confirmed by determination of the X-ray crystal structure (Figure 3).¹⁹ In both cases the new phosphorus-based stereogenic center is formed with high diastereoselectivity (*vide infra*). As lithiation of (*S*)-**1b** is also known to proceed with high diastereoselectivity,^{12c,15} this enabled the application of this methodology to the synthesis of (*S,S_p*,*S_{phos}*)-**L3b** and (*S,S_p*,*R_{phos}*)-**L4b**, both isolated as single diastereoisomers following column chromatography.

Scheme 2. Stereoselective synthesis of *P*-stereogenic diastereoisomers (*S,S_p*,*S_{phos}*)-**L3a/b** and (*S,S_p*,*R_{phos}*)-**L4a/b**.



The extension of this methodology to the synthesis of the remaining two planar chiral diastereoisomers started with (*S,R_p*)-2-*d*-**1a** (>90% D incorporation).¹⁵ Use of lithiation conditions B, followed by addition of *o*-TolPCL₂ and then PhMgBr as before, led to isolation of a single Phosferrox derivative containing *n*-butyl and *ortho*-tolyl groups (14% yield). The competitive and undesired reaction of *n*-BuLi led us to evaluate *s*-BuLi/THF as an alternative base/solvent combination for this reaction. These conditions (C) applied previously to the lithiation of (*S*)-**1a** resulted in an 8 : 1 dr,¹¹ therefore allowing the larger kinetic isotope effect associated with lithiation²⁰ to dominate planar chiral diastereoselectivity. To confirm this, application of conditions C to the lithiation of (*S,R_p*)-2-*d*-**1a** followed by addition of TMSCl

resulted in a 5 : 1 ratio of the resulting silylated (S,R_p) : (S,S_p) diastereoisomers (68% yield). Furthermore, lithiation in the same way, followed by addition of *o*-TolPCl₂ and then BuMgCl, led to the predominant formation of the same Phosferrox derivative formed when using *n*-BuLi as base, assigned as (S,R_p,S_{phos})-**2a** by analogy to the chemistry described above (Scheme 3).

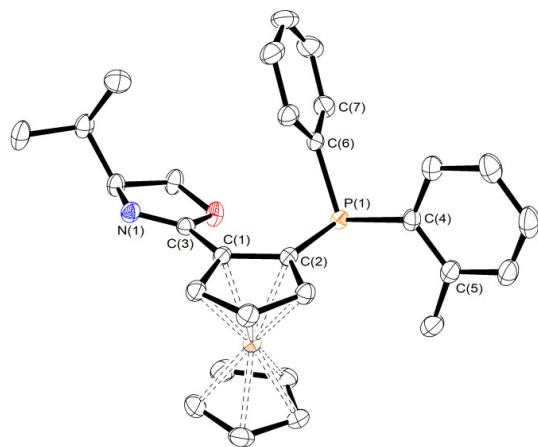


Figure 2. Representation of the X-ray crystal structure of (S,S_p,S_{phos})-**L3a** (hydrogen atoms omitted for clarity). Principal bond lengths [Å] include: C(1)-C(3) 1.460(2), C(2)-P(1) 1.8283(18). Principal torsions [°] include: C(2)-C(1)-C(3)-N(1) -159.38(19), C(1)-C(2)-P(1)-C(6) 58.74(17), C(1)-C(2)-P(1)-C(4) 162.54(16), C(2)-P(1)-C(6)-C(7) -154.48(15), C(2)-P(1)-C(4)-C(5) 86.89(15). Thermal ellipsoids are drawn at the 50% probability level.

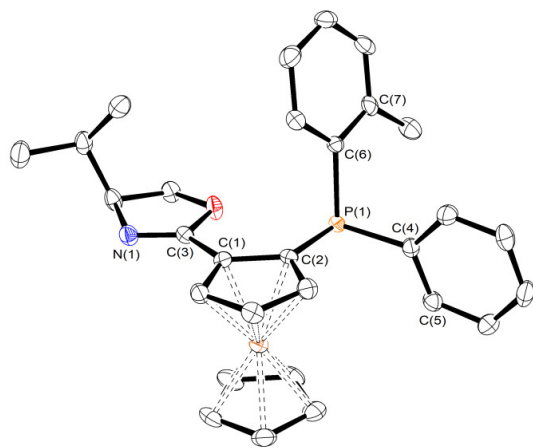
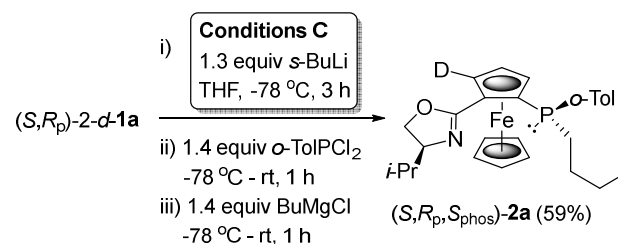
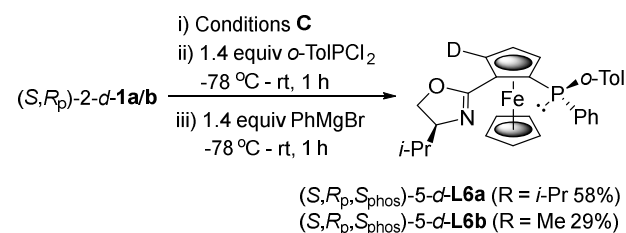
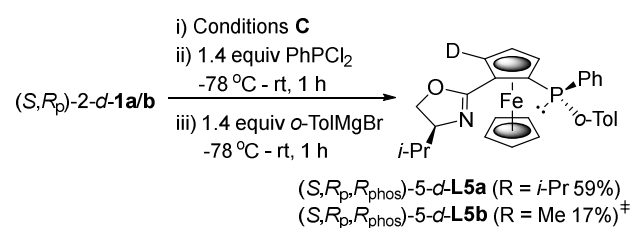


Figure 3. Representation of the X-ray crystal structure of (S,S_p,R_{phos})-**L4a** (hydrogen atoms omitted for clarity). Principal bond lengths [Å] include: C(1)-C(3) 1.458(4), C(2)-P(1) 1.826(3). Principal torsions [°] include: C(2)-C(1)-C(3)-N(1) -175.6(3), C(1)-C(2)-P(1)-C(6) 83.7(3), C(1)-C(2)-P(1)-C(4) -172.2(3), C(2)-P(1)-C(6)-C(7) 176.3(2), C(2)-P(1)-C(4)-C(5) 92.6(3). Thermal ellipsoids are drawn at the 50% probability level.

Scheme 3. Demonstration of deuterium mediated planar chirality reversal with configurational control of a phosphorus based stereogenic center.



Scheme 4. Stereoselective synthesis of *P*-stereogenic diastereoisomers (S,R_p,R_{phos})-**L5** and (S,R_p,S_{phos})-**L6**, †(dr = 15 : 1 for **L5b**).



Conditions C were then used for the synthesis of (S,R_p,R_{phos})-**5-d-L5a** and (S,R_p,S_{phos})-**5-d-L6a** (Scheme 4). For both reactions comparison of the ³¹P NMR spectra obtained after work-up again revealed excellent configurational control of the phosphorus-based stereogenic center. Separation of the product from the alternative planar chiral diastereoisomer was achieved readily by column chromatography. This chemistry was extended to the synthesis of (S,R_p,R_{phos})-**5-d-L5b** and (S,R_p,S_{phos})-**5-d-L6b**, both isolated predominantly as a single diastereoisomer following column chromatography, and in the case of (S,R_p,S_{phos})-**L6b** as a single isomer by a subsequent recrystallisation. The configuration of (S,R_p,S_{phos})-**L6b** was established by an X-ray crystal structure determination (Figure 4),¹⁹ confirming that it is the planar chirality, and not the oxazoline-based stereogenic center, that exclusively controls the configuration of the phosphorus based stereogenic center.

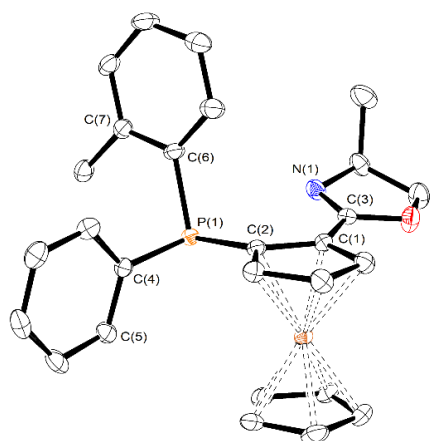
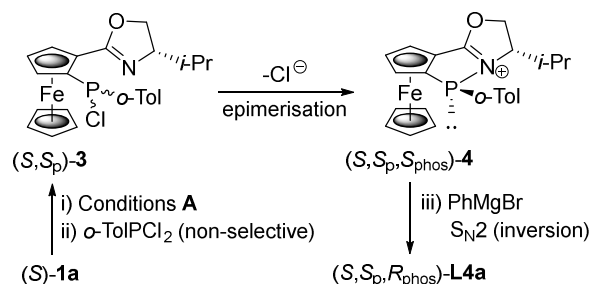


Figure 4. Representation of the X-ray crystal structure of (S,R_p,S_{phos}) -**L6b** (hydrogen and deuterium atoms omitted for clarity). Principal bond lengths [Å] include: C(1)-C(3) 1.460(5), C(2)-P(1) 1.828(4). Principal torsions [°] include: C(2)-C(1)-C(3)-N(1) 7.5(7), C(1)-C(2)-P(1)-C(6) -89.2(4), C(1)-C(2)-P(1)-C(4) 169.7(3), C(2)-P(1)-C(6)-C(7) -173.0(3), C(2)-P(1)-C(4)-C(5) -93.3(3). Thermal ellipsoids are drawn at the 50% probability level.

All four *P*-stereogenic diastereoisomers (**L3a/b-L6a/b**) were obtained by column chromatography on silica. In some instances epimerization was noted, as has been observed previously for other *P*-stereogenic compounds containing a ferrocenyl substituent.^{18a,21,22} To investigate this further, a 1 : 0.2 ratio of (S,S_p,S_{phos}) -**L3a**/ (S,S_p,R_{phos}) -**L4a** was heated for 24 h at 65 °C in hexane containing chromatography grade SiO₂ (40-65 μm). This resulted in a change in ratio to 1 : 1, whereas the in the absence of SiO₂, or with added neutral Al₂O₃ (10-200 μm), no change was observed. Similarly, heating initially pure (S,R_p,R_{phos}) -5-d-**L5a** in silica doped hexane for 24 h at 65 °C gave a 1 : 0.5 ratio of (S,R_p,R_{phos}) -5-d-**L5a**/ (S,R_p,S_{phos}) -5-d-**L6a**. The promotion of epimerization by SiO₂ is likely related to its acidity.^{23,24}

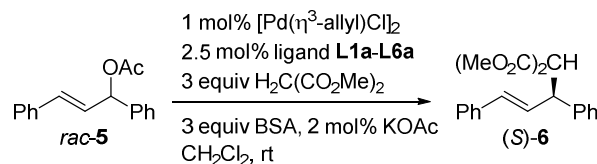
The excellent diastereoselectivity observed for the formation of the *P*-stereogenic ligands could be a consequence of highly selective differentiation, on substitution, of the two enantiotopic chlorine components of ArPCl₂ to give **3** (kinetic control). Alternatively, this first step in the reaction could proceed without selectivity, requiring a subsequent epimerization step to produce a single isomer (thermodynamic control). To investigate this further the synthesis of (S,S_p,R_{phos}) -**L4a** was repeated, with examination by ³¹P NMR spectroscopy of the reaction mixture after addition *o*-TolPCl₂ and warming to room temperature (Scheme 5). This revealed a major signal at 67.7 ppm, and following addition of PhMgBr this signal disappeared and was replaced by the ³¹P NMR signal for (S,S_p,R_{phos}) -**L4a** (-23.9 ppm). No signal was observed for (S,S_p,S_{phos}) -**L3a** (ca. -28 ppm), from which the diastereoselectivity is estimated as > 100 : 1.

Scheme 5. Planar chiral stereocontrol in the synthesis of *P*-stereogenic diastereoisomer (S,S_p,R_{phos}) -**L4a**.



Attempts to isolate and further characterize the intermediate were unsuccessful. There are three pointers to the intermediate being (S,S_p,S_{phos}) -**4**,¹⁸ although these do not rule out other possibilities. Firstly, aryl/lone-pair differentiation, with the aryl group oriented away from ferrocene in this cyclic structure, provides a basis for epimerization selectivity. Thermodynamic control of this sort has been used to explain the diastereoselective formation of other auxiliary mediated *P*-stereogenic derivatives.^{17,25} Secondly, stereospecific S_N2 reaction with PhMgBr accounts for the configuration of the phosphorus-based stereogenic center in (S,S_p,R_{phos}) -**L4a**. Finally, the observed ³¹P NMR chemical shift of 67.7 ppm is in reasonable agreement with the value estimated for this intermediate (ca. 72 ppm). This was determined starting with the reported value of 88 ppm²⁶ for (DMAP)Ph₂P⁺TfO⁻ corrected for the replacement of both phenyls by a ferrocenyl (-9.4 ppm) and an *ortho*-tolyl (-6.4 ppm) group.²⁷

Table 1. Palladium catalysed allylic alkylation with ligands **L1a-L6a**.



Entry	Ligand	Time (h)	Conversion (%) ^[a]	ee% ^[b]
1	(S,S_p) - L1a	<1	100	95
2	(S,R_p) -5-d- L2a	24	92	71
3	(S,S_p,S_{phos}) - L3a	5	100	97
4	(S,S_p,R_{phos}) - L4a	7	100	95
5	(S,R_p,R_{phos}) -5-d- L5a	24	90	84
6	(S,R_p,S_{phos}) -5-d- L6a	24	86	63

[a] Determined by ¹H NMR spectroscopy. [b] Determined by chiral HPLC. (S) -**6** was the major enantiomer in all cases.

The availability of all four ligand diastereoisomers enabled a preliminary investigation into the influence of ligand configuration on enantioselectivity. To this end we chose as a model reaction palladium catalysed allylic alkylation of *rac*-**5** as *t*-Bu substituted Phosferrox ligands are known to work well in this chemistry.^{13a,28} In addition to *i*-Pr

substituted diastereomeric ligands **L3a-L6a**, precursor ligands **L1a** and **L2a** were also employed. All ligands resulted in the formation of (*S*)-**6** as the major enantiomer, and the ee values obtained (Table 1) revealed a clear preference for the *S,S_p* diastereoisomers, with the *S_p,R_p,S_{phos}* ligand **L3a** being the matched/matched diastereoisomer. A pictorial representation of these ee values highlights the additive/subtractive influence of the phosphorus-based stereogenic center (Figure 5a). Assuming the availability of only two diastereomeric transition states,²⁹ an alternative representation of these results is as $\Delta\Delta G_{R,S}^\ddagger$ values (calculated with *rt* = 22 °C - Figure 5b). In both cases the phosphine based stereogenic center adds to $\Delta\Delta G_{R,S}^\ddagger$ a value of ~ 2 kJ mol⁻¹.

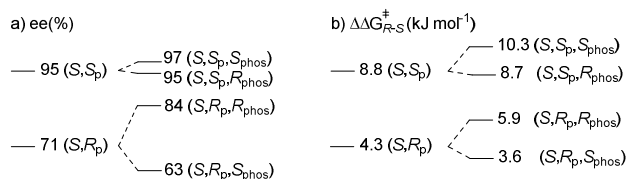


Figure 5. Outcomes of palladium catalysed allylic alkylation with ligands **L1a-L6a**.

CONCLUSION

We have demonstrated that sequential addition to a lithiated planar chiral ferrocenyloxazoline of Ar¹PCl₂ followed by Ar²MgBr results in a diarylferrocenyl phosphine with excellent control of diastereoselectivity. Coupled with the high diastereoselectivity of lithiation, as controlled by the (*S*)-valine or (*S*)-alanine derived oxazoline auxiliary, and the use of a deuterium blocking group to reverse lithiation diastereoselectivity, this enables the selective synthesis of all possible Phosferrox ligand diastereoisomers containing three elements of chirality. To the best of our knowledge this is the first time four diastereoisomers of a bidentate ligand have been synthesized. On application to palladium-catalysed allylic alkylation of *trans*-1,3-diphenylallyl acetate the configuration of the matched/matched *i*-Pr substituted ligand was identified as *S,S_p,S_{phos}*. Furthermore, comparison in catalysis to the simpler diphenylphosphine containing Phosferrox ligands reveals a positive and negative influence of the phosphorus-based stereogenic center. The results of this study concur with our previous literature analysis on the addition of an element of chirality to a parent ferrocene-based ligand to give both possible diastereomeric offspring ligands.³⁰ In nearly all cases this analysis revealed (in 20 out of 23 examples), as in this work, a systematic increase and decrease in ee, providing further support for the iterative use of this approach in ligand optimization studies. Application of the new ligands in this way is currently in progress.

EXPERIMENTAL SECTION

General remarks. Diethyl ether and tetrahydrofuran were distilled over sodium and benzophenone ketyl. Tetramethylethylenediamine (TMEDA) was dried with 4 Å MS. All lithiation reactions were carried out under an inert atmosphere of either nitrogen or argon. Alkylolithiums and Grignards were not titrated prior to use. Silica gel (60 Å pore size, 40 - 63 μm technical grade) was used for chromatography. All protons and carbons were assigned using 2D NMR techniques including HSQC, HMBC, NOESY and COESY.

Preparation of (S)-2-[(S_p)-2-((R_{phos})-*ortho*-tolylphenylphosphino)ferrocenyl]-4-(1-methylethyl)oxazoline, (S,S_p,S_{phos})-L3a.** (*S*)-**1a** (0.100 g, 0.34 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry diethyl ether (4 mL). TMEDA (0.07 mL, 0.44 mmol) was added and solution was cooled to -78 °C and stirred for 5 min after which *s*-butyllithium (1.4 M in hexanes) (0.31 mL, 0.44 mmol) was slowly added. After stirring for 3 hours, dichlorophenylphosphine (64 μL, 0.47 mmol) was added and the reaction allowed to stir at room temperature for 1 h. The reaction was re-cooled to -78 °C and *o*-tolylmagnesium bromide (2.0 M in THF) (0.24 mL, 0.47 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 10% EtOAc/hexane) yielded an orange solid (0.11 g, 63%). *R_f* 0.2 (10% EtOAc/hexane). Mp 150 - 151 °C. [α]_D^{25.5°C} = +120 (*c* 0.20, CHCl₃). IR (film): 3053, 2955, 2926, 2873, 1656 (CN). ¹H NMR (500 MHz, CDCl₃): 7.29 - 7.22 (2H, m, *o*-TolH), 7.23 - 7.15 (5H, m, PhH), 7.08 - 7.01 (2H, m, *o*-TolH), 5.09 (1H, brs, CpH), 4.40 (1H, brs, CpH), 4.27 (1H, apt, ²⁺*J*_{H/H} = 8.9 Hz, CHH), 4.22 (5H, s, CpH), 3.85 (1H, td, ²*J*_{H/H} = 9.0, ³*J*_{H/H} = 6.2 Hz, CH), 3.72 (1H, s, CpH), 3.41 (1H, apt, ²⁺*J*_{H/H} = 8.5 Hz, CHH), 2.85 (3H, s, CH₃), 1.62 (1H, apoct, ³⁺*J*_{H/H} = 6.6 Hz, CH), 0.83 (3H, d, ³*J*_{H/H} = 6.8 Hz, CH₃), 0.66 (3H, d, ³*J*_{H/H} = 6.8 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 166.4 (C=N), 143.5 (d, ²*J*_{CP} = 27.4 Hz, *o*-TolC), 140.5 (d, ¹*J*_{CP} = 13.0 Hz, PhC), 136.8 (d, ¹*J*_{CP} = 15.4 Hz, *o*-TolC), 136.0 (d, ²*J*_{CP} = 3.7 Hz, *o*-TolC), 132.4 (d, ²*J*_{CP} = 20.4 Hz, PhC), 130.1 (d, ³*J*_{CP} = 4.9 Hz, *o*-TolC), 129.2 (*o*-TolC), 128.2 (d, ³*J*_{CP} = 6.8 Hz, PhC), 127.8 (PhC), 126.1 (d, ³*J*_{CP} = 2.1 Hz, *o*-TolC), 79.8 (d, ²*J*_{CP} = 17.6 Hz, CpC), 74.5 (CpC), 74.2 (CpC), 71.9 (CpC), 71.6 (CH), 71.0 (2CpC), 70.1 (CH₂), 32.0 (CH), 22.2 (d, ³*J*_{CP} = 22.9 Hz, CH₃), 19.1 (CH₃), 17.7 (CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -26.51 (PPhO-Tol). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₉H₃₁FeNOP 496.1493; Found 496.1493.**

Preparation of dichloro*ortho*-tolylphosphine. Bis(diethylamino)chlorophosphine (5.00 g, 23.7 mmol) was added to a flame dried Schlenk tube, dissolved in diethyl ether (25 mL) and cooled to -78 °C. *Ortho*-tolylmagnesium bromide (2M in THF) (17.80 mL 35.6 mmol) was then added slowly and the reaction allowed to warm to room temperature and stirred vigorously for 1 h. The resulting suspension was allowed to settle and the mother liquor transferred to a dry flask by cannula filtration, and the remaining white solid washed with diethyl ether (3 x 10 mL). The solvent was then removed *in vacuo* and the resulting oil re-dissolved in fresh diethyl ether (10 mL). The flask was cooled to 0 °C and hydrogen chloride (2 M in Et₂O) (59.33 mL, 118.7 mmol) was added and the reaction allowed to stir vigorously at room temperature for 1 hour. Again, the resulting suspension was allowed to settle and the mother liquor transferred to a dry flask by cannula filtration, and the remaining white solid washed with diethyl ether (3 x 10 mL). The solvent was removed *in vacuo* and the resulting oil purified by Kugelrohr distillation (130 - 140 °C @ 7 mbar) to give a clear oil (2.53 g, 55 %): ¹H NMR (500 MHz, CDCl₃): 8.05 (1H, brd, ³*J*_{H/H} = 7.0 Hz, ArH), 7.44 (1H, apt, ³⁺*J*_{H/H} = 7.4 Hz, ArH), 7.38 (1H, apt, ³⁺*J*_{H/H} = 7.3 Hz, ArH), 7.23 (1H, brd, ³*J*_{H/H} = 7.2 Hz, ArH), 2.65 (3H, s, ArCH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 140.6 (d, ²*J*_{CP} = 35.5 Hz, ArC), 137.9 (d, ¹*J*_{CP} = 57.5 Hz, ArC), 132.7 (ArC), 130.9 (ArC), 130.4 (d, ²*J*_{CP} = 11.1 Hz, ArC), 127.1 (ArC), 20.0 (d, ³*J*_{CP} = 24.9 Hz, CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): 163.35 (ClPAr₂). Matches previously reported data.³¹

Preparation of (S)-2-[(S_p)-2-((R_{phos})-*ortho*-tolylphenylphosphino)ferrocenyl]-4-(1-methylethyl)oxazoline, (S,S_p,R_{phos})-L4a.** (*S*)-**1a** (0.100 g, 0.34 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry diethyl ether (4 mL). TMEDA (0.07 mL, 0.44 mmol) was added and solution was cooled to -78 °C and stirred for 5 min after which *s*-butyllithium (1.4 M in hexanes) (0.31 mL, 0.44 mmol) was slowly added. After stirring for 3 hours, *ortho*-tolyl dichlorophosphine (64 μL, 0.47 mmol) was added and the reaction allowed to stir at room**

temperature for 1 h. The reaction was re-cooled to -78 °C and phenylmagnesium bromide (1.0 M in THF) (0.47 mL, 0.47 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 10% EtOAc/hexane) yielded an orange solid (0.09 g, 54%). *R*_f 0.3 (10% EtOAc/hexane). Mp 104 - 105 °C. [α]_D^{25.5°C} = +20 (c 0.20, CHCl₃). IR (film): 3053, 2959, 2902, 2869, 1660 (CN). ¹H NMR (500 MHz, CDCl₃): 7.46 (2H, aptd, ³J_{HP} = 7.3, ³J_{HH} = 7.3, ⁴J_{HH} = 2.2 Hz, PhH), 7.38 - 7.34 (3H, m, PhH), 7.14 (1H, aptd, ³J_{HP} = 7.4, ³J_{HH} = 7.4, ⁴J_{HH} = 1.3 Hz, *o*-TolH), 7.10 - 7.05 (1H, m, *o*-TolH), 7.01 (1H, apt, ³⁺³J_{HH} = 7.4 Hz, *o*-TolH), 6.84 - 6.79 (1H, m, *o*-TolH), 5.13 (1H, brs, CpH), 4.42 (1H, apt, ³⁺³J_{HH} = 2.0 Hz, CpH), 4.30 (1H, apt, ²⁺³J_{HH} = 8.9 Hz, CHH), 4.24 (5H, s, CpH), 3.92 - 3.86 (1H, m, CH), 3.74 (1H, apt, ²⁺³J_{HH} = 8.1 Hz, CHH), 3.59 (1H, s, CpH), 2.28 (3H, d, ⁴J_{HP} = 1.5 Hz, CH₃), 1.71 (1H, apoct, ³⁺³⁺³J_{HH} = 6.7 Hz, CH), 0.85 (3H, d, ³J_{HH} = 6.8 Hz, CH₃), 0.71 (3H, d, ³J_{HH} = 6.8 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 165.3 (C=N), 140.9 (d, ²J_{CP} = 25.5 Hz, *o*-TolC), 138.3 (d, ¹J_{CP} = 14.2 Hz, *o*-TolC), 137.3 (d, ¹J_{CP} = 12.5 Hz, PhC), 134.9 (d, ²J_{CP} = 21.1 Hz, PhC), 131.9 (*o*-TolC), 129.7 (d, ³J_{CP} = 4.5 Hz, *o*-TolC), 129.1 (PhC), 128.4 (d, ³J_{CP} = 7.1 Hz, PhC), 128.1 (*o*-TolC), 125.7 (*o*-TolC), 78.9 (d, ²J_{CP} = 14.7 Hz, CpC), 75.8 (d, ¹J_{CP} = 16.6 Hz, CpC), 74.4 (d, ²J_{CP} = 4.2 Hz, CpC), 72.4 (d, ³J_{CP} = 1.9 Hz, CpC), 72.2 (CH), 71.0 (CpC), 70.9 (d, ³J_{CP} = 1.0 Hz, CpC), 69.8 (CH₂), 32.4 (CH), 21.1 (d, ³J_{CP} = 21.7 Hz, CH₃), 18.8 (CH₃), 17.8 (CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -23.90 (Po-TolPh). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₉H₃₁FeNOP 496.1493; Found 496.1497.

Preparation of (S)-2-[(S_P)-2-((S_{phos})-*ortho*-tolylphenylphosphino)ferrocenyl]-4-methyloxazoline, (S,S_P,S_{phos})-L3b. (S)-1b (0.120 g, 0.45 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry diethyl ether (5 mL). TMEDA (0.09 mL, 0.58 mmol) was added and solution was cooled to -78 °C and stirred for 5 min after which *s*-butyllithium (1.4 M in hexanes) (0.41 mL, 0.58 mmol) was slowly added. After stirring for 2 hours, dichlorophenylphosphine (85 μL, 0.62 mmol) was added and the reaction allowed to stir at room temperature for 1 h. The reaction was re-cooled to -78 °C and *o*-tolylmagnesium bromide (2.0 M in THF) (0.31 mL, 0.62 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 30% EtOAc/hexane) yielded an orange solid (0.08 g, 39%). *R*_f 0.2 (30% EtOAc/hexane). Mp 56 - 57 °C. [α]_D^{22.4°C} = +182 (c 0.20, CHCl₃). IR (film): 3052, 3003, 2964, 2929, 2890, 1649 (CN). ¹H NMR (500 MHz, CDCl₃): 7.29 - 7.17 (7H, m, PhH + *o*-TolH), 7.08 - 6.98 (2H, m, *o*-TolH), 4.95 (1H, ddd, ³J_{HH} = 2.4, ⁴J_{HH} = 1.5, ⁴J_{HP} = 0.5 Hz, CpH), 4.38 - 4.31 (2H, m, CpH + CHH), 4.23 (5H, s, CpH), 4.09 - 4.00 (1H, m, CH), 3.69 (1H, ddd, ³J_{HH} = 2.4, ³J_{HP} = 1.4, ⁴J_{HH} = 0.8 Hz, CpH), 3.02 (1H, dd, ²J_{HH} = 9.0, ³J_{HH} = 8.1 Hz, CHH), 2.86 (3H, s, CH₃), 1.03 (3H, d, ³J_{HH} = 6.5 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 166.1 (C=N), 143.5 (d, ²J_{CP} = 27.4 Hz, *o*-TolC), 140.4 (d, ¹J_{CP} = 12.8 Hz, PhC), 136.9 (d, ¹J_{CP} = 15.4 Hz, *o*-TolC), 135.8 (d, ²J_{CP} = 3.3 Hz, *o*-TolC), 132.6 (d, ²J_{CP} = 20.6 Hz, PhC), 130.1 (d, ³J_{CP} = 4.9 Hz, *o*-TolC), 129.2 (*o*-TolC), 128.2 (d, ³J_{CP} = 6.9 Hz, PhC), 127.9 (PhC), 126.1 (d, ³J_{CP} = 2.0 Hz, *o*-TolC), 79.7 (d, ²J_{CP} = 15.5 Hz, CpC), 74.6 (CpC), 74.2 (CpC + CH₂), 71.2 (CpC), 70.9 (d, ³J_{CP} = 0.8 Hz, CpC), 70.6 (CpC), 61.5 (CH), 22.2 (d, ³J_{CP} = 22.7 Hz, CH₃), 21.0 (CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -25.89 (PPho-Tol). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₇H₂₇FeNOP 468.1180; Found = 468.1174.

Preparation of (S)-2-[(S_P)-2-((R_{phos})-*ortho*-tolylphenylphosphino)ferrocenyl]-4-methyloxazoline, (S,S_P,R_{phos})-L4b. (S)-1b (0.120 g, 0.45 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry diethyl ether (5 mL). TMEDA (0.09 mL, 0.58 mmol) was added and solution was cooled to -78 °C and stirred for 5 min after which *s*-

butyllithium (1.4 M in hexanes) (0.41 mL, 0.58 mmol) was slowly added. After stirring for 2 hours, *ortho*-tolylidichlorophosphine (85 μL, 0.62 mmol) was added and the reaction allowed to stir at room temperature for 1 h. The reaction was re-cooled to -78 °C and phenylmagnesium bromide (1.0 M in THF) (0.62 mL, 0.62 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 30% EtOAc/hexane) yielded an orange solid (0.10 g, 49%). *R*_f 0.3 (30% EtOAc/hexane). Mp 188 - 190 °C. [α]_D^{22.2°C} = +84 (c 0.27, CHCl₃). IR (film): 3049, 2961, 2904, 1642 (CN). ¹H NMR (500 MHz, CDCl₃): 7.49 - 7.44 (2H, m, PhH), 7.39 - 7.33 (3H, m, PhH), 7.16 (1H, apt, ³J_{HH} = 6.8, ³J_{HP} = 6.8 Hz, *o*-TolH), 7.11 - 7.06 (1H, m, *o*-TolH), 7.04 (1H, apt, ³⁺³J_{HH} = 7.4 Hz, *o*-TolH), 6.84 (1H, apdd, ³⁺³J_{HH} = 6.7, ⁴J_{HH} = 4.3 Hz, *o*-TolH), 5.03 (1H, brs, CpH), 4.43 - 4.34 (2H, m, CpH + CHH), 4.23 (5H, s, CpH), 4.15 - 4.03 (1H, m, CH), 3.56 (1H, brs, CpH), 3.39 (1H, apt, ²⁺³J_{HH} = 8.3 Hz, CHH), 2.27 (3H, s, CH₃), 1.13 (1H, d, ³J_{HH} = 6.6 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 166.1 (C=N), 141.0 (d, ²J_{CP} = 25.3 Hz, *o*-TolC), 138.2 (d, ¹J_{CP} = 14.3 Hz, *o*-TolC), 137.1 (d, ¹J_{CP} = 11.9 Hz, PhC), 134.8 (d, ²J_{CP} = 21.0 Hz, PhC), 131.7 (*o*-TolC), 129.8 (d, ³J_{CP} = 4.4 Hz, *o*-TolC), 129.1 (PhC), 128.4 (d, ³J_{CP} = 7.3 Hz, PhC), 128.1 (*o*-TolC), 125.7 (*o*-TolC), 78.7 (d, ²J_{CP} = 14.5 Hz, CpC), 75.4 (d, ¹J_{CP} = 16.3 Hz, CpC), 74.5 (d, ³J_{CP} = 3.8 Hz, CpC), 74.2 (CH₂), 72.4 (d, ³J_{CP} = 1.8 Hz, CpC), 71.2 (CpC), 70.9 (d, ³J_{CP} = 0.9 Hz, CpC), 61.7 (CH), 21.3 (CH₃), 21.1 (CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -23.32 (Po-TolPh). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₇H₂₇FeNOP 468.1180; Found 468.1181.

Preparation of (S)-2-[(R_P)-2-((S_{phos})-butyl*ortho*-tolylphosphino)-5-deuterioferrocenyl]-4-(1-methylethyl)oxazoline, (S,R_P,S_{phos})-2a. (S)-2-*d*-1a¹⁵ (0.040 g, 0.13 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry THF (1 mL). The solution was cooled to -78 °C and stirred for 5 min after which *s*-butyllithium (1.4 M in hexanes) (0.13 mL, 0.17 mmol) was slowly added. After stirring for 2 hours, *ortho*-tolylidichlorophosphine (27.5 μL, 0.19 mmol) was added and the reaction allowed to stir at room temperature for 1 h. The reaction was re-cooled to -78 °C and butylmagnesium chloride (2.0 M in THF) (0.09 mL, 0.19 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 10% EtOAc/hexane) yielded an orange oil (0.038 g, 59%). *R*_f 0.2 (10% EtOAc/hexane). [α]_D^{23.5°C} = -153 (c 0.16, CHCl₃). IR (film): 3056, 2957, 2929, 2873, 1656 (CN). ¹H NMR (500 MHz, CDCl₃): 7.07 - 7.05 (2H, m, *o*-TolH), 6.97 - 6.89 (2H, m, *o*-TolH), 4.53 (1H, d, ³J_{HH} = 2.5 Hz, CpH), 4.51 (1H, d, ³J_{HH} = 2.4 Hz, CpH), 4.25 (5H, s, CpH), 4.01 (1H, dd, ²J_{HH} = 9.7, ³J_{HH} = 8.2 Hz, CHH), 3.95 (1H, apt, ²⁺³J_{HH} = 7.4 Hz, CHH), 3.89 - 3.84 (1H, m, CH), 2.63 (3H, d, ⁴J_{HP} = 1.1 Hz, CH₃), 2.04 - 1.96 (1H, m, CHH), 1.83 - 1.76 (1H, m, CHH), 1.66 - 1.56 (2H, m, CH + CHH), 0.51 - 1.42 (3H, m, CHH + CH₂), 0.93 (3H, t, ³J_{HH} = 7.2 Hz, CH₃), 0.70 (3H, d, ³J_{HH} = 6.8 Hz, CH₃), 0.66 (3H, d, ³J_{HH} = 6.8 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 165.1 (C=N), 142.1 (d, ²J_{CP} = 27.8 Hz, *o*-TolC), 140.5 (d, ¹J_{CP} = 17.3 Hz, *o*-TolC), 131.5 (d, ²J_{CP} = 2.0 Hz, *o*-TolC), 129.3 (d, ³J_{CP} = 5.4 Hz, *o*-TolC), 128.0 (*o*-TolC), 125.8 (*o*-TolC), 79.9 (d, ²J_{CP} = 16.7 Hz, CpC), 74.7 (CpC), 72.1 (CH), 71.5 (d, ²J_{CP} = 3.4 Hz, CpC), 70.8 (CpC), 70.4 (CpC), 69.3 (CH₂), 32.4 (CH), 29.0 (d, ²J_{CP} = 1.8 Hz, CH₂), 28.8 (d, ¹J_{CP} = 4.9 Hz, CH₂), 24.7 (d, ³J_{CP} = 12.5 Hz, CH₂), 21.9 (d, ³J_{CP} = 23.1 Hz, CH₃), 18.3 (CH₃), 17.8 (CH₃), 14.0 (CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -40.38 (Po-TolBu). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₇H₃₄DFeNOP 477.1869; Found 477.1866.

Preparation of (S)-2-[(R_P)-2-((R_{phos})-*ortho*-tolylphenylphosphino)-5-deuterioferrocenyl]-4-(1-methylethyl)oxazoline, (S,R_P,R_{phos})-2-*d*-L5a. (S)-2-*d*-1a (0.073 g, 0.25 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry THF (3 mL). The solution was

cooled to -78 °C and stirred for 5 min after which *s*-butyllithium (1.4 M in hexanes) (0.23 mL, 0.32 mmol) was slowly added. After stirring for 2 hours, dichlorophenylphosphine (47 μ L, 0.34 mmol) was added and the reaction allowed to stir at room temperature for 1 h. The reaction was re-cooled to -78 °C and *ortho*-tolylmagnesium bromide (2.0 M in THF) (0.17 mL, 0.34 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 10% EtOAc/hexane) yielded an orange solid (0.07 g, 59%). *R*_f 0.2 (10% EtOAc/hexane). Mp 100 - 101 °C. $[\alpha]_D^{25.5^\circ\text{C}} = -80$ (c 0.22, CHCl₃). IR (film): 3052, 2957, 2873, 1656 (CN). ¹H NMR (500 MHz, CDCl₃): 7.28 - 7.22 (2H, m, *o*-TolH), 7.21 - 7.13 (5H, m, PhH), 7.08 - 7.02 (2H, m, *o*-TolH), 4.37 (1H, d, ³J_{HH} = 2.4 Hz, CpH), 4.20 (5H, s, CpH), 3.99 (1H, dd, ³J_{HH} = 7.0, ³J_{HH} = 5.0 Hz, CHH), 3.87 - 3.75 (2H, m, CH + CHH), 3.71 (1H, d, ³J_{HH} = 2.3 Hz, CpH), 2.84 (3H, s, CH₃), 1.57 (1H, apoct, ³⁺³J = 5.9 Hz, CH), 0.72 (3H, d, ³J_{HH} = 6.8 Hz, CH₃), 0.71 (3H, d, ³J_{HH} = 6.8 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 165.1 (C=N), 143.7 (d, ²J_{CP} = 26.6 Hz, *o*-TolC), 140.7 (d, ¹J_{CP} = 12.4 Hz, PhC), 137.0 (d, ¹J_{CP} = 15.5 Hz, *o*-TolC), 136.5 (d, ²J_{CP} = 4.9 Hz, *o*-TolC), 132.3 (d, ²J_{CP} = 20.0 Hz, PhC), 130.1 (d, ³J_{CP} = 4.8 Hz, *o*-TolC), 129.2 (*o*-TolC), 128.0 (d, ³J_{CP} = 6.7 Hz, PhC), 127.6 (PhC), 126.0 (d, ³J_{CP} = 2.5 Hz, *o*-TolC), 79.0 (d, ²J_{CP} = 15.3 Hz, CpC), 74.4 (d, ²J_{CP} = 2.8 Hz, CpC), 72.1 (CH), 72.0 (d, ¹J_{CP} = 20.8 Hz, CpC), 70.8 (CpC), 70.3 (CpC), 69.5 (CH₂), 32.5 (CH), 22.4 (d, ³J_{CP} = 22.2 Hz, CH₃), 18.3 (CH₃), 17.9 (CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -26.69 (PPho-Tol). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₉H₃₀DFeNOP 497.1556; Found 497.1556.

Preparation of (S)-2-[(R_p)-2-((S_{phos})-*ortho*-tolylphenylphosphino)-5-deuteroferrocenyl]-4-(1-methylethyl)oxazoline, (S,R_p,S_{phos})-2-d-L6a. (S)-2-d-1a¹⁵ (0.100 g, 0.34 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry THF (4 mL). The solution was cooled to -78 °C and stirred for 5 min after which *s*-butyllithium (1.4 M in hexanes) (0.31 mL, 0.44 mmol) was slowly added. After stirring for 2 hours, *ortho*-tolylidichlorophosphine (63 μ L, 0.47 mmol) was added and the reaction allowed to stir at room temperature for 1 h. The reaction was re-cooled to -78 °C and phenylmagnesium bromide (1.0 M in THF) (0.47 mL, 0.47 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 10% EtOAc/hexane) yielded an orange solid (0.10 g, 58%). *R*_f 0.3 (10% EtOAc/hexane). Mp 97 - 98 °C. $[\alpha]_D^{22.4^\circ\text{C}} = +85.5$ (c 0.22, CHCl₃). IR (film): 3052, 2954, 2873, 1656 (CN). ¹H NMR (500 MHz, CDCl₃): 7.49 (2H, aptd, ³J_{HH} = 7.3, ³J_{HP} = 7.3, ⁴J_{HH} = 2.2 Hz, PhH), 7.38 - 7.34 (3H, m, PhH), 7.13 (1H, aptd, ³J_{HH} = 7.4, ³J_{HP} = 7.4, ⁴J_{HH} = 1.3 Hz, *o*-TolH), 7.08 - 7.04 (1H, m, *o*-TolH), 6.99 (1H, apt, ³⁺³J_{HH} = 7.7 Hz, *o*-TolH), 6.78 (1H, ddd, ³J_{HH} = 7.6, ⁴J_{HH} = 4.3, ⁵J_{HP} = 1.1 Hz, *o*-TolH), 4.37 (1H, d, ³J_{HH} = 2.5 Hz, CpH), 4.23 (5H, s, CpH), 4.08 (1H, dd, ³J_{HH} = 9.6, ²J_{HH} = 8.2 Hz, CHH), 3.98 (1H, dd, ²J_{HH} = 8.2, ³J_{HH} = 6.6 Hz, CHH), 3.90 (1H, aptd, ³⁺³J_{HH} = 9.6, ³J_{HH} = 6.2 Hz, CH), 3.57 (1H, dd, ³J_{HH} = 2.5, ²J_{HH} = 0.6 Hz, CpH), 2.29 (3H, d, ⁴J_{HP} = 1.4 Hz, CH₃), 1.48 (1H, apoct, ³⁺³J_{HH} = 6.3 Hz, CH), 0.64 (3H, d, ³J_{HH} = 6.8 Hz, CH₃), 0.59 (3H, d, ³J_{HH} = 6.8 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 164.7 (d, ³J_{CP} = 3.6 Hz, C=N), 141.0 (d, ²J_{CP} = 25.4 Hz, *o*-TolC), 138.1 (d, ¹J_{CP} = 12.8 Hz, *o*-TolC), 137.4 (d, ¹J_{CP} = 12.2 Hz, PhC), 135.0 (d, ²J_{CP} = 20.8 Hz, PhC), 131.8 (*o*-TolC), 129.7 (d, ³J_{CP} = 4.7 Hz, *o*-TolC), 129.0 (PhC), 128.3 (d, ³J_{CP} = 7.2 Hz, PhC), 128.0 (*o*-TolC), 125.5 (*o*-TolC), 78.6 (d, ²J_{CP} = 13.5 Hz, CpC), 75.3 (d, ¹J_{CP} = 16.1 Hz, CpC), 74.6 (d, ²J_{CP} = 4.4 Hz, CpC), 72.3 (CH), 70.8 (CpC), 70.6 (CpC), 69.8 (CH₂), 32.8 (CH), 21.1 (d, ³J_{CP} = 21.7 Hz, CH₃), 18.2 (CH₃), 17.9 (CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -24.70 (Po-TolPh). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₉H₃₀DFeNOP 497.1556; Found 497.1557.

Preparation of (S)-2-[(R_p)-2-((R_{phos})-*ortho*-tolylphenylphosphino)-5-deuteroferrocenyl]-4-methyloxazoline, (S,R_p,R_{phos})-2-d-L5b. (S)-2-d-1b¹⁵ (0.100 g, 0.37 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry diethyl ether (5 mL). The solution was cooled to -78 °C and stirred for 5 min after which *s*-butyllithium (1.4 M in hexanes) (0.34 mL, 0.48 mmol) was slowly added. After stirring for 2 hours, dichlorophenylphosphine (70 μ L, 0.52 mmol) was added and the reaction allowed to stir at room temperature for 1 h. The reaction was re-cooled to -78 °C and *o*-tolylmagnesium bromide (2.0 M in THF) (0.26 mL, 0.52 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 30% EtOAc/hexane) yielded an orange oil (0.04 g, 17%). *R*_f 0.2 (30% EtOAc/hexane). $[\alpha]_D^{23.5^\circ\text{C}} = -67$ (c 0.60, CHCl₃). IR (film): 3052, 2962, 2923, 2873, 1652 (CN). ¹H NMR (500 MHz, CDCl₃): 7.27 - 7.22 (2H, m, *o*-TolH), 7.20 - 7.14 (5H, m, PhH), 7.07 - 7.04 (2H, m, *o*-TolH), 4.37 (1H, d, ³J_{HH} = 2.5 Hz, CpH), 4.20 (5H, s, CpH), 4.08 - 4.02 (1H, m, CH), 3.91 - 3.86 (1H, m, CHH), 3.79 (1H, dd, ²J_{HH} = 8.0, ³J_{HH} = 5.8 Hz, CHH), 3.70 (1H, dd, ³J_{HH} = 2.5, ⁴J_{HH} = 0.7 Hz, CpH), 2.85 (3H, s, CH₃), 0.96 (3H, d, ³J_{HH} = 6.6 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 165.3 (d, ³J_{CP} = 2.9 Hz, C=N), 143.6 (d, ²J_{CP} = 27.0 Hz, *o*-TolC), 140.7 (d, ¹J_{CP} = 12.4 Hz, PhC), 136.9 (d, ¹J_{CP} = 15.4 Hz, *o*-TolC), 136.4 (d, ²J_{CP} = 4.3 Hz, *o*-TolC), 132.3 (d, ²J_{CP} = 20.2 Hz, PhC), 130.1 (d, ³J_{CP} = 4.7 Hz, *o*-TolC), 129.2 (*o*-TolC), 128.0 (d, ³J_{CP} = 6.7 Hz, PhC), 127.5 (PhC), 126.0 (d, ³J_{CP} = 2.2 Hz, *o*-TolC), 79.4 (d, ²J_{CP} = 15.7 Hz, CpC), 74.4 (CpC), 74.3 (d, ¹J_{CP} = 10.3 Hz, CpC), 73.6 (CH₂), 70.9 (CpC), 70.4 (CpC), 61.6 (CH), 22.3 (d, ³J_{CP} = 22.4 Hz, CH₃), 21.3 (CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -26.51 (PPho-Tol). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₇H₂₆DFeNOP 469.1243; Found 469.1242.

Preparation of (S)-2-[(R_p)-2-((S_{phos})-*ortho*-tolylphenylphosphino)-5-deuteroferrocenyl]-4-methyloxazoline, (S,R_p,S_{phos})-2-d-L6b. (S)-2-d-1b¹⁵ (0.100 g, 0.37 mmol) was added to a flame dried Schlenk tube under an inert atmosphere and dissolved in dry tetrahydrofuran (4 mL). The solution was cooled to -78 °C and stirred for 5 min after which *s*-butyllithium (1.4 M in hexanes) (0.34 mL, 0.48 mmol) was slowly added. After stirring for 2 hours, *ortho*-tolylidichlorophosphine (76 μ L, 0.52 mmol) was added and the reaction allowed to stir at room temperature for 1 h. The reaction was re-cooled to -78 °C and phenylmagnesium bromide (1.0 M in THF) (0.52 mL, 0.52 mmol) was added and the reaction allowed to warm to room temperature and stir for an additional hour. The reaction was cooled to 0 °C quenched with saturated sodium carbonate solution and separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. Purification by column chromatography (SiO₂, 30% EtOAc in Hexane) yielded an orange solid as a 16:1 ratio of diastereoisomers (0.055 g, 32%). This was then recrystallized by dissolving in a minimum amount of dichloromethane and layering hexane on top and allowing to stand for 2 days in the freezer to give orange crystals (0.05 g, 29%). *R*_f 0.3 (30% EtOAc/hexane). Mp 178 - 179 °C. $[\alpha]_D^{24.3^\circ\text{C}} = -14$ (c 0.56, CHCl₃). IR (film): 3051, 2965, 2897, 1650 (CN). ¹H NMR (500 MHz, CDCl₃): 7.51 - 7.45 (2H, m, PhH), 7.38 - 7.33 (3H, m, PhH), 7.15 (1H, aptd, ³⁺³J_{HH} = 7.4, ⁴J_{HH} = 1.4 Hz, *o*-TolH), 7.09 - 7.05 (1H, m, *o*-TolH), 7.01 (1H, apdd, ³⁺³J_{HH} = 11.3, ⁴J_{HH} = 4.1 Hz, *o*-TolH), 6.80 (1H, ddd, ³J_{HH} = 7.6, ³J_{HP} = 4.3, ⁴J_{HH} = 1.2 Hz, *o*-TolH), 4.39 (1H, dd, ³J_{HH} = 2.5, ⁴J_{HP} = 0.5 Hz, CpH), 4.22 (5H, s, CpH), 4.13 - 4.07 (2H, m, CH + CHH), 3.86 - 3.78 (1H, m, CHH), 3.57 (1H, dd, ³J_{HH} = 2.5, ³J_{HP} = 0.7 Hz, CpH), 2.28 (3H, d, ⁴J_{HP} = 1.4 Hz, CH₃), 1.04 (1H, d, ³J_{HH} = 6.5 Hz, CH₃). ¹³C{¹H} NMR (125 MHz, CDCl₃): 165.4 (d, ³J_{CP} = 3.1 Hz, C=N), 141.0 (d, ²J_{CP} = 25.2 Hz, *o*-TolC), 138.1 (d, ¹J_{CP} = 13.5 Hz, *o*-TolC), 137.3 (d, ¹J_{CP} = 12.0 Hz, PhC), 134.9 (d, ²J_{CP} = 20.9 Hz, PhC), 131.8 (*o*-TolC), 129.7 (d, ³J_{CP} = 4.6 Hz, *o*-TolC), 129.1 (PhC), 128.4 (d, ³J_{CP} = 7.2 Hz, PhC), 128.0 (*o*-TolC), 125.5 (*o*-TolC), 78.7 (d, ²J_{CP} = 14.2 Hz, CpC), 75.0 (d, ¹J_{CP} = 16.4 Hz, CpC), 74.6 (d, ²J_{CP} = 4.2 Hz, CpC), 73.8 (CH₂), 71.1 (d, ³J_{CP} = 0.9 Hz, CpC), 61.7 (CH),

21.5 (CH₃), 21.2 (d, ³J_{CP} = 21.2 Hz, CH₃). ³¹P{¹H} NMR (202 MHz, CDCl₃): -24.02 (P_O-TolPh). HRMS (ASAP-TOF) *m/z*: [M+H]⁺ Calcd for C₂₇H₂₅DFeNOP+H⁺ 469.1243; Found 469.1236.

General Procedure for Allylic Alkylation. Ligand (0.0125 mmol) and [Pd(allyl)Cl]₂ (0.0019 g, 0.005 mmol) were added to a flame dried Schlenk tube, dissolved in dichloromethane (0.85 mL) and allowed to stir for 30 min at room temperature. In a separate flask, racemic 1,3-diphenylallyl acetate (0.126 g, 0.50 mmol) was weighed out and subsequently dissolved in dichloromethane (0.85 mL). After 30 min, the acetate solution was added to the reaction vessel followed by dimethyl malonate (0.17 mL, 1.5 mmol), *N,O*-bis(trimethylsilyl)acetamide (0.37 mL, 1.5 mmol) and KOAc (0.001 g, 0.01 mmol) in that order. The reaction was stirred at room temperature and monitored by TLC (SiO₂, 5% EtOAc/hexane). After completion or 24 h (whichever came first), the reaction was quenched with saturated ammonium chloride, separated with diethyl ether, dried with magnesium sulphate and the solvent removed *in vacuo*. A crude ¹H NMR was performed at this point to determine the conversion. Purification by column chromatography (SiO₂, 5% EtOAc/hexane) gave the product as a colourless oil. Enantiomeric excess measurements was determined by chiral HPLC analysis using a CHIRALCEL OD-H; Eluent = 98:2 (Hexane:IPA); Flow rate = 0.5 mL min⁻¹; Concentration = 0.0015 g mL⁻¹, Injection volume = 5 μL. Major enantiomer RT = 18.8 min, Minor enantiomer RT = 17.3 min. ¹H NMR (500 MHz, CDCl₃) δ 7.34 - 7.27 (8H, m, PhH), 7.26 - 7.18 (2H, m, PhH), 6.48 (1H, d, ³J_{HH} = 15.7 Hz, HC=CH), 6.33 (1H, dd, ³J_{HH} = 15.7, ³J_{HH} = 8.6 Hz, HC=CH), 4.27 (1H, dd, ³J_{HH} = 10.8, ³J_{HH} = 8.7 Hz, CH), 3.96 (1H, d, ³J_{HH} = 10.9 Hz, CH), 3.71 (3H, s, CH₃), 3.52 (3H, s, CH₃). ¹³C{¹H}NMR (125 MHz, CDCl₃): 168.4 (C=O), 167.9 (C=O), 140.3 (PhC), 137.0 (PhC), 132.0 (HC=CH), 129.3 (HC=CH), 128.9 (PhC), 128.6 (PhC), 128.0 (PhC), 127.7 (PhC), 127.3 (PhC), 126.5 (PhC), 57.8 (CH), 52.8 (CH₃), 52.6 (CH₃), 49.3 (CH). Matches previously reported data.³²

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website.

Copies of the ¹H, ¹³C, ³¹P NMR spectra, HPLC traces and details of the X-ray crystal structure determinations (PDF), together with the associated CIF files.

Accession Codes

CCDC 1916753, 1916752, and 1916751 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

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Notes

The authors declare no competing financial interest.

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