**Small Molecule Activation by Intermolecular Zr(IV)-Phosphine Frustrated Lewis Pairs**

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**ABSTRACT:** We report intermolecular transition metal frustrated Lewis pairs (FLPs) based on zirconocene aryloxide and phosphine moieties that exhibit a broad range of small molecule activation chemistry that has previously been the preserve of only intramolecular pairs. Reactions with $\text{D}_2$, $\text{CO}_2$, THF, and PhCCH are reported. By contrast with previous intramolecular examples, these systems allow facile access to a variety of steric and electronic characteristics at the Lewis acidic and Lewis basic components, with the three-step syntheses of 10 new intermolecular transition metal FLPs being reported. Systematic variation to the phosphine Lewis base is used to unravel steric considerations, with the surprising conclusion that phosphines with relatively small Tolman steric parameters not only give highly reactive FLPs but are often seen to have the highest selectivity for the desired product. DOSY NMR spectroscopic studies on these systems reveal for the first time the nature of the Lewis acid/ Lewis base interactions in transition metal FLPs of this type.

**1. INTRODUCTION**

Frustrated Lewis pairs (FLPs) have proved to be a powerful new concept in small molecule activation and catalysis. By controlling the steric and electronic architecture of certain combinations of Lewis acids and bases to preclude the formation of a classical Lewis adduct, a high latent reactivity is imparted on the system.\(^1\) This donor–acceptor ability is reminiscent of transition metal chemistry, and the ability of main group FLP systems to mimic reactivity normally associated with transition metals has been one of the remarkable features of this area. Initial investigations focused on the use of phosphine–borane FLPs and their ability to heterolytically cleave dihydrogen and facilitate hydrogenation reactions,\(^2\) in addition to the binding and activation of carbon dioxide (CO\(_2\)).\(^3\) Subsequently, it was shown that main group FLPs are also able to mediate a wider range of transformations, such as 1,2-addition to alkynes\(^4\) and the ring opening of cyclic ethers.\(^5\)

A wide selection of inter- and intramolecular main group FLPs based on diverse Lewis acid and base groups has now been reported.\(^6-8\) These pairs are able to mediate the heterolytic cleavage of $\text{H}_2$, $\text{CO}_2$ and isocyanate sequestration and deprotonation or 1,2 addition to terminal alkynes. Recent reports have also shown the utility of main group FLPs in catalytic hydrogenation reactions. Intermolecular main group FLP systems are ubiquitous despite the obvious entropic disadvantages of this approach.

We, and others, have extended FLP chemistry to transition metals in the hope that combining the powerful small molecule activation chemistry of FLPs with the well-known suite of catalytically relevant reactions of transition metals could lead to yet more new chemistry.\(^9\) Much of our initial focus has been on intramolecular systems in which the fluorinated borane fragment is replaced by an electrophilic group 4 metalloocene (Figure 1, A−C). The chemistry of these cationic zirconocene−phosphinoaryl oxide complexes in general mirrors main group systems (activation of $\text{H}_2$, $\text{CO}_2$, THF), but also demonstrates reactivity that is either unique or rarely observed in main group systems, such as C=Cl and C=F bond cleavage, and catalytic dehydrocoupling of amine-boranes.\(^10\) Other related intramolecular zirconocene-phosphine systems have also been reported by Erker et al. (Figure 1, D and E). These compounds are accessed through 1,1- or 1,2-carbozirconation reactions of alkynes to the zirconium(IV) cation $[\text{Cp}^*\text{ZrCH}=\text{CH}_2][\text{B}(\text{C}_6\text{F}_5)_4]$. As with our intramolecular systems, these Zr/P pairs react with...
Complex 1 is stabilized by solvent coordination to a multitude of small molecules (CO, CO₂, H₂, NO PhC(H)O, PhN₃O). In stark contrast to the wide, varied and selective reactivity of intramolecular Zr/P FLPs, transformations mediated by intermolecular Zr/P FLPs are extremely limited. There are only two examples reported to date with the substrates employed limited to relatively reactive molecules containing highly polarized C=O or N=O bonds (example in Scheme 1). This very narrow reactivity is doubly disappointing in that species 2, which exhibits an essentially linear Zr₁-O₁-Mes bond angle (153.2°) indicative of multiple Zr-O bonding. Solvent coordination in 2 is presumably precluded by the Cp* ligand. In 2, unlike in previously structurally characterized examples of cationic Zr aryloxide complexes, there is no evidence of an agostic Zr-O bonding.

2. RESULTS AND DISCUSSION

2.1. Synthesis of Cationic Zirconocene Aryloxide Lewis Acids. The required cationic Zr(IV) fragments shown in Scheme 2 were synthesized via two routes. The complex bearing Cp ligands was accessed through preparation of [Cu₂Zr(OMes)₂] (OMes = 2,4,6-trimethylphenoxide) by a modified literature procedure. Subsequent methyl abstraction using [Cu₂Zr(OMes)₂] in a noncoordinating (PhCl) solvent gave [Cu₂ZrOMes] [Cu₂ZrF₅] (1) in 94% yield. The Cu⁺ analogue was synthesized by an alternative route, as protonolysis of a methyl group from Cu₂ZrMe₂ by 2,4,6-trimethylphenol (MesOH) was found to be extremely sluggish (60% yield after >10 days, 20 °C, hexane). [Cu₂ZrOMes][Cu₂ZrF₅] (2) was therefore accessed by initial methyl abstraction from Cu₂ZrMe₂ using [Cu₂Zr(OMes)₂] prior to protonolysis of the remaining methyl group using MesOH (Scheme 3). This modification afforded the desired complex in 85% yield over two steps in minutes.

The molecular structure of Zr(IV) cations 1 and 2 are shown in Figure 2. Complex 1 is stabilized by solvent coordination (chlorobenzene) in the solid state causing a slight bending of the Zr-O₁-Mes bond angle (153.2°). This is contrary to species 2, which exhibits an essentially linear Zr-O₁-Mes angle (176.7°) indicative of multiple Zr-O bonding. Solvent coordination in 2 is presumably precluded by the additional steric bulk afforded by the Cu⁺ ligand. In 2, unlike in previously structurally characterized examples of cationic Zr aryloxide complexes, there is no evidence of an agostic interaction between the ortho-alkyl group and the electron deficient zirconium.

2.2. Reaction with Phosphines: Generation of Frus-treated Lewis Pairs (FLPs). Taking our inspiration from main group systems and our previous intramolecular examples, initial attempts to generate an FLP system from 1 and 2 were made by addition of the bulky phosphine P₃Bu₃. However, this always failed.
resulted in an uncharacterizable mixture of products including appreciable amounts of [HPBu3][B(C6F5)4] \( ^{31}P \) NMR \( \delta = 59.1 \) ppm) which precluded further clean reactivity. By contrast, addition of an equimolar amount of a less basic and less sterically hindered phosphine (PCy3 (a), PEt3 (b), PPPh3 (c), PMes5 (d), and P(C6F5)3 (e)) to 1 and 2 in chlorobenzene solution resulted in clean conversion to new species.

In this case \(^{31}P\) NMR spectroscopy is a useful probe for the nature of the Zr-P interaction, formation of Zr-P bond resulting in a large downfield shift (Table 1). In the case of 1, it was found that upon addition of PCy3, PEt3, and PPPh3, a Zr-P interaction was formed with 1a, 1b, and 1c all exhibiting large downfield shifts in their \(^{31}P\) NMR resonances, when compared to the free phosphine. The systems containing the more bulky PMes5 and P(C6F5)3, 1d and 1e, show no change in their \(^{31}P\) NMR chemical shift, suggestive of the absence of a Zr-P interaction. In contrast, none of the systems with the bulkier Cp* complex 2 (2a-e) show evidence of a Zr-P interaction in solution. This pattern is in good agreement with the Tolman steric parameters of the phosphines as shown in Table 1. Only the less bulky phosphines (with the less bulky zirconocene 1) possess a Zr-P interaction. The less basic nature of the fluoroaryl substituted phosphine is also likely to be an important electronic consideration.

A DOSY (Diffusion-Ordered Spectroscopy) NMR study was undertaken to detect potential secondary interactions present between the Lewis acid and Lewis base, as well as to further explore the nature of the interaction present in 1a-e. A similar study has been carried out on main group PR3/B(C6F5)3 (R = Bu and Mes) FLPs confirming secondary interactions are present between the fluorines on the B(C6F5)3 and the protons on PR3.17

Our study shows that in 1a-e the interaction observed by \(^{31}P\) NMR spectroscopy is in fact dynamic and not a persistent Zr-P bond. For example, under our conditions (0.06 mol dm\(^{-3}\), 0.5-PBr) 18 was found to possess a diffusion coefficient (D) of \(6.0 \times 10^{-10} \) m\(^2\) s\(^{-1}\) and for PEt3 (b) a value of D = 19.5 \( \times 10^{-10} \) m\(^2\) s\(^{-1}\). Upon combination of 1 with 1 equiv of PEt3 to form 1b, the values of D obtained for the two components were found to be 5.5 \( \times 10^{-10} \) m\(^2\) s\(^{-1}\) (1) and 7.3 \( \times 10^{-10} \) m\(^2\) s\(^{-1}\) (PEt3). The smaller diffusion coefficients in both cases indicate an interaction in solution consistent with the \(^{31}P\) NMR spectrum, however if the interaction was a persistent Zr-P bond the values of D for the two components should be equal. The nature of the interaction is therefore dynamic, with the equilibrium positioned toward the “bound” pair. Similar observations were made in the case of 1a and 1c (data in the Supporting Information).

For FLP systems 1d-e and 2a-e, data obtained from DOSY experiments again shows the two components possessing smaller diffusion coefficients when in combination than when measured separately (Figures S15–S31). Taking 2b as an example, the values of D for the separate components are 8.6 \( \times 10^{-10} \) m\(^2\) s\(^{-1}\) (2) and 19.5 \( \times 10^{-10} \) m\(^2\) s\(^{-1}\) (PEt3), but upon combination these shift to 8.0 \( \times 10^{-10} \) m\(^2\) s\(^{-1}\) (2) and 16.5 \( \times 10^{-10} \) m\(^2\) s\(^{-1}\) (PEt3, (b)). This suggests again that a dynamic equilibrium may be present with encounter complexes forming and separating in solution. In this case, however, the increased steric bulk of the Cp* ligands means that a classical metal-phosphine bond cannot form; therefore, the dynamic equilibrium must arise from other weaker secondary interactions perhaps between the ancillary ligands. In conclusion, the DOSY study does indicate some degree of preorganization of the FLP prior to further reactions.

2.3. Reactivity of Pairs with Dihydrogen (D2). The heterolytic cleavage of H2 is perhaps the most typical example of small molecule activation mediated by FLPs and was a logical starting point here. For experimental expedience, D2 was used in place of H2 to allow more precise monitoring by \(^{2}H\) NMR spectroscopy.

When PhCl solutions of 1a-e were pressurized with 1 bar D2 no reaction was observed (Scheme 4). This is consistent with previous work where intramolecular Zr-P FLP systems bearing the Cp ligand set showed no reaction with H2 under similar conditions. Previous work has indicated the necessity for at least one Cp* ligand to achieve heterolytic hydrogen cleavage, attributed to the more electron rich ligand facilitating transient binding of H2 to the Zr metal center and allowing deprotonation of this now more acidic species by the internal phosphine base. Consistent with this previous observation, 2a and 2b both showed an instantaneous reactions with 1 bar D2. In the case of 2a, a new species is observed by \(^{31}P\) NMR spectroscopy (\( \delta = 33.7 \) ppm, \( \delta_{PD} = 67 \) Hz) displaying a characteristic 1:1:1 splitting pattern indicative of the formation of a P-D bond. The \(^{1}H\) NMR spectrum confirms this assignment, with a doublet (\( \delta = 4.25 \) ppm, \( \delta_{PD} = 67 \) Hz) corresponding to the phosphonium deuterium and a sharp singlet (\( \delta = 5.96 \) ppm) assigned as the Zr-deuteride. Treatment of 2b with 1 bar D2 results in a similar downfield shift in the \(^{31}P\) NMR spectrum to give a 1:1:1 triplet again symptomatic of a P-D bond (\( \delta = 21.1 \) ppm, \( \delta_{PD} = 67 \) Hz). Species 2c-e display no reactivity under the same conditions, with the lower basicity of these aryl-substituted phosphines being our working hypothesis for this observation.

In an attempt to probe the mechanism of the hydrogen cleavage reaction, a chlorobenzene solution of 2 was pressurized with D2 and cooled to -35 °C, at which point no evidence for a Zr-D2 complex was evident. These findings suggest a mechanism akin to that proposed by computational studies carried out on main-group FLP systems, in particular PH3Bu3/B(C6F5)3. In this case, it is proposed that preorganization of the FLP occurs prior to activation of the H2. This is corroborated by our DOSY data discussed above which
indicates the presence of transient encounter complexes in solution.

2.4. Reactivity of Pairs with Carbon Dioxide (CO₂). A range of main group and transition metal-based FLPs have shown the ability to sequester CO₂. The pairs 1a–e and 2a–e were treated with CO₂ by pressurizing chlorobenzene solutions of the species with 1 bar CO₂. Upon pressurizing with CO₂, systems 1a and 1b showed quantitative conversion to new species assigned as the CO₂ activation product by 31P NMR spectroscopy (3 δ = 27.9 ppm, 4 δ = 28.0 ppm). Compound 1c was found to yield two new species upon treatment with 1 bar CO₂ with 31P NMR chemical shifts of 5.4 and 19.9 ppm, however the 13C NMR spectrum showed no evidence of the carbonyl carbon. 1d and 1e were found to be inactive in the activation of CO₂ (Scheme 5).

Scheme 5. Reactivity of FLP Systems 1a–e and 2a–e with 1 bar CO₂

2a and 2b also react rapidly and quantitatively with 1 bar CO₂ giving rise to new species observed in the 31P NMR spectrum at δ = 24.9 ppm (5) and δ = 24.1 ppm (6). As with 1c–e, 2c gives a mixture of products when treated with CO₂, with no carbonyl peak visible in the 13C NMR spectrum, and 2d and 2e display no reactivity (Scheme 5). It surprised us to some extent that the cleanest results were obtained with the relatively nonbulky alkyl phosphines a and b, even in the cases where a Zr–P interaction is observed (1a and 1b); this echoes more recent results with main group FLPs where a truly "frustrated" system has been shown to be unnecessary so long as the Lewis acid and base can act in a cooperative fashion under the reaction conditions. The moniker "Cooperative Lewis Pairs" would seem to be increasingly appropriate. This also corroborates the DOSY study on 1a and 1b, which found that the Zr–P interaction is dynamic and a small amount of unbound Zr and PR₃ are present in solution. It is thought to be these species that react to form the desired products.

Crystalization under 1 bar CO₂ allows isolation of X-ray quality crystals of 6 in low (<5%) isolated yield. The molecular structure of 6 is shown in Figure 3. The solid state structure of 6 shows a slight lengthening of the Zr1–O2 bond length in comparison with 2 (1.962(2)Å vs 1.937(2)Å) indicative of a slight loss of the multiple bond character due to coordination of an additional ligand at the electron deficient Zr center. As expected, C30 appears to be tending toward sp² in character with values of 112.7(2)° and 117.8(2)° for the O3–C30–P1 and O2–C30–P1 angles, respectively, but a significantly larger angle (129.5(3)°) between O2–C30–O3 indicates that C30 retains some sp character. This is further reinforced by the only slightly longer C30–O2 bond length (1.278(4)Å) when compared to the C30–O3 double bond (1.220(4)Å).

2.5. Reactivity of Pairs with Tetrahydrofuran (THF). Treatment of chlorobenzene solutions of 1a–c with an excess of THF results in an immediate color change from orange to yellow and concomitant dissociation of the bound phosphine to yield what is proposed to be [(CpZr(THF)OMes)[B(C₆F₅)₄]]. 1a reacts further to give quantitative conversion to 7 within 16 h (31P NMR δ = 38.4 ppm), 1b undergoes a somewhat more rapid reaction to yield a species with a similar 31P NMR shift (31P NMR δ = 31.5 ppm) after 30 min assigned as 8. 1c shows no reaction at room temperature; however, upon heating to 80 °C for 6h full conversion to 9 is observed by 31P NMR spectroscopy (δ = 23.4 ppm). As with H₂ and CO₂, 1d and 1e show no further reaction with THF despite heating at 80 °C for 16 h (Scheme 6).

Scheme 6. Reactivity of FLP Systems 1a–e and 2a–e with THF

To further probe the mechanism of this reaction, postulated intermediate [(CpZr(THF)OMes)[B(C₆F₅)₄]] (1-THF) has been synthesized and isolated by reaction of 1 with THF, the molecular structure of which is shown in Figure 4. In comparison to 1, 1-THF shows a greater degree of bending of the Zr1–O1–Mes bond (139.8(8)° vs 153.2(2)°) in addition to a lengthening of the Zr1–O1 bond (1.972(1)Å vs 1.935(2)Å) due to the coordination of a more donating
ligand in the THF compared to chlorobenzene, thus further reducing the multiple bond character between the Zr and the aryloxide ligand.

Subsequent reaction of 1-THF with PR₃ (R = Cy, Et, Ph, Mes, C₆F₅) results in reactivity identical to that shown in Scheme 6. We can therefore propose that the reaction mechanism consists of an initial complexation of THF to the Lewis acidic zirconocene center, which activates the THF toward nucelophilic attack at the α-carbon by the phosphine. This mechanism fits well with the observed trend of the more nucleophilic phosphines giving more rapid reaction (PEt₃ > PCy₃ > PPh₃ > PMes > P(C₆F₅)₃). Reaction of 1-THF with PEt₃ being significantly more rapid than with PCy₃ is thought to be a purely steric effect, with PEt₃ having a cone angle of 132° compared to 170° for PCy₃. This mechanism is also consistent with works by Stephan et al. and Jordan et al., which describe the ring opening of Zr bound THF by phosphines and amines and independently conclude that the reaction proceeds by a Lewis acid activation of the C=O bond prior to nucleophilic attack at the α-carbon.

Similar, but overall less rapid reactivity is observed with 2a–e. Upon addition of excess THF to chlorobenzene solutions of 2a–e an immediate color change from red to yellow is observed indicating formation of a Zr-THF adduct as observed for 1a–e. 2-THF was isolated and characterized by the addition of THF to a chlorobenzene solution of 2, and the molecular structure is shown in Figure 5. An interesting structural feature of 2-THF is that, unlike its Cp analogue, the aryloxide ligand appears to be locked in conformation, with free rotation about the Zr1–O1–Mes axis precluded by the additional steric bulk. This is evidenced by the ¹H NMR spectra for the two species, wherein 1-THF is seen to possess two equivalent ortho-CH₃ groups (δ = 1.84 ppm (broad)); however, in 2-THF, these become inequivalent (δ = 1.79 and 1.88 ppm).

The solid-state structure of 2-THF also shows a significant bending of the Zr1–O1–Mes bond when compared to 2 (153.6(1)° vs 176.7(2)°). This is accompanied by an extension of the Zr1–O1 bond upon binding of THF (1.937(2) Å to 1.984(1) Å) again indicating that binding of an additional donor ligand to the Zr center reduces the multiple bond character of the Zr1–O1 bond.

Analogous to compound 1a, species 2a reacts with an excess of THF at room temperature to yield 10 in 10 days (³¹P NMR δ = 36.8 ppm). Compound 2b again reacts significantly faster, proceeding to a >99% conversion to 11 in 3 days (³¹P NMR δ = 29.2 ppm). Compound 2c shows no reactivity with THF at room temperature, but upon heating to 80 °C complete conversion to 12 is observed within 12 h (³¹P NMR δ = 21.9 ppm). As with the system bearing the Cp ligand set, the analogous Cp* species 2d and 2e show no reaction with THF even at elevated temperature (80 °C, 24h). This general trend of the ring opening of cyclic ethers proceeding less rapidly with 2a–e than 1a–e is proposed to be a steric effect with Cp* hindering the attack of the incoming phosphine nucleophile.

2.6. Reactivity of Pairs with Alkynes. The reaction of FLPS with terminal alkynes can proceed via one of two mechanisms, with the majority of main-group FLP systems going via a 1,2-addition reaction with the nature of the resulting isomer generally controlled by steric factors. However, in previous work with Zr/P FLPS, it has been shown that a deprotonation reaction may also take place yielding a zirconium acetylidy and phosphonium species (Scheme 7).

In the case of 1a, upon addition of phenylacetylene (PhCCH), clean deprotonation is observed to yield [HPCys][B(C₆F₅)₄] (³¹P NMR δ = 33.1 ppm, Jₕₖ = 420 Hz) and a zirconium acetylide complex. Surprisingly, compound 1b shows a change in selectivity, and when treated with PhCCH it undergoes a slow reaction (20 °C, PhCl, 16 h) to yield a mixture of the two isomers of the 1,2-addition product (1:8). Separation of the isomers proved impossible due to their near identical solubility in a range of solvents. Reaction of 1c with PhCCH rapidly (20 °C, PhCl, <1 min) yields the 1,2-addition product 13 (³¹P [¹H] NMR δ = 20.1 ppm) with only the Zr/P trans isomer isolated. This was identified by comparison of the

Figure 4. Molecular structures of 1-THF as determined by single crystal X-ray diffraction. Thermal ellipsoids are drawn at the 50% probability level. Disorder around the THF ligand, hydrogens, and the [B(C₆F₅)₄]⁻ counterion are omitted for clarity. Selected bond lengths (Å) and angles (deg): Zr1−O1 1.972(1), Zr1−O2 2.206(1), O2−Zr1−O1 139.8(8), O1−Zr1−O2 96.82(5).

Figure 5. Molecular structure of 2-THF as determined by single crystal X-ray diffraction. Hydrogens, [B(C₆F₅)₄]⁻ counterion, and solvent of crystallization are omitted for clarity. Selected bond lengths (Å) and angles (deg): Zr1−O1 1.984(1), Zr1−O2 2.282(1), O2−Zr1−O1 95.49(5), Zr1−O1−Mes 153.6(1).
1H NMR spectra to the crystallographically characterized analogue 14 vide infra. In both cases, the alkenyl proton exhibits a $^{3}$J$_{PH}$ coupling of 45 Hz indicating an identical geometry around the double bond. A further change in selectivity is observed with 1d with the favored reaction pathway reverting back to deprotonation, such that upon treatment of 1d with PhCCH immediate formation of [H-PMes$_3$][B(C$_6$F$_5$)$_4$] is detected by $^{31}$P NMR spectroscopy ($^{31}$P NMR $\delta$ = -27.5 ppm, $^{1}$J$_{PH}$ = 478 Hz). Complex 1e exhibits no reaction with PhCCH. The pairs 2a–e exhibit a broadly similar trend in reactivity; however, both 2a and 2b yield a mixture of deprotonation and 1,2-addition products in ratios of 2:1 and 3:2, respectively. Again the system containing PPh$_3$, 2c, reacts cleanly and rapidly with PCCH to generate only the 1,2-addition product 14 ($^{31}$P[${^1}$H] NMR $\delta$ = 17.4 ppm), the molecular structure of which is shown in Figure 6. As is observed with 1d and 1e, 2d forms only the deprotonation product, [H-PMes$_3$][B(C$_6$F$_5$)$_4$], and 2e does not react upon addition of PhCCH.

The molecular structure of 14 (Figure 6) reveals the trans-Zr/P conformation with the Ph moiety of PhCCH geminal to PPh$_3$. This conformer is assumed to be preferred as it reduces steric clashes between the bulky Cp$^*$ ligands with the phenyl rings of both PPh$_3$ and PhCCH. C31 appears to possess a greater degree of sp$^2$ character when compared to C30 as evidenced by the large Zr1–C30–C31 angle (143.3(2)$^\circ$), this could again be attributed to the steric strain enforced by the interaction between the Ph group and the bulk aryloxide ligand on Zr.

These two competing reaction pathways have previously been observed by Erker et al., the main group FLP system P$^\text{III}$Bu$_3$/B(C$_6$F$_5$)$_3$ reacting with a terminal alkyn to give the deprotonation product, whereas PAr$_3$/B(C$_6$F$_5$)$_3$ (PAR$_3$ = P(0-tolyl)$_3$ or P(Ph)$_2$[2,5-bis(trifluoromethyl)phenyl]) cleanly yields the 1,2-addition product. This reactivity can generally be attributed to electronic factors with the more basic phosphines favoring deprotonation; however, there may also be a steric effect as P(0-tolyl)$_3$ and PMes$_3$ are considered to be electronically similar, but have significantly different steric parameters, with cone angles of 194$^\circ$ and 212$^\circ$, respectively. This could be responsible for the switch in reactivity from 1,2-addition (P(0-tolyl)$_3$), Erker et al.) to deprotonation (P(Mes)$_3$, vide supra).

3. CONCLUSION

We have synthesized a range of intermolecular zirconium/phosphine FLPs derived from zirconocene cations and tertiary phosphines of varying steric and electronic properties. A DOSY NMR spectroscopic study on these systems has shown the nature of the Lewis acid/Lewis base interactions present in all cases. These pairs show for the first time the ability of intermolecular FLPs containing a transition metal fragment as the Lewis acid to react in an analogous fashion to their intramolecular counterparts. These new systems are shown to mediate the activation of a range of small molecules (D$_2$, CO$_2$, THF, phenylacetylene) with the reactivity toward these substrates highly dependent on the steric and electronic nature of the phosphine employed, a factor which had remained previously unexplored with transition metal FLPs. It has been found that the phosphine must be of sufficient basicity to promote such reactions; in all cases, systems using the weakly basic P(C$_6$F$_5$)$_3$ (1e and 2e) show no reactivity toward the small molecules studied. Given sufficient Lewis basicity, high steric bulk in the phosphine used is surprisingly unimportant; indeed, the least bulky phosphine used here, PE$_3$, gives the cleanest results. In addition, the base used has a dramatic effect on selectivity, as evidenced by the switch in reaction mode with phenylacetylene from 1,2-addition to deprotonation when the less bulky PPh$_3$ (1c and 2c) is replaced with the significantly more bulky P(Mes)$_3$ (1d and 2d). These results show that the use of intramolecular systems is not a prerequisite for transition metal FLPs and open many other possibilities for the design of intermolecular transition metal frustrated or cooperative Lewis pairs.

4. EXPERIMENTAL SECTION

4.1. General Considerations. Unless otherwise stated, all manipulations were undertaken under an atmosphere of argon or nitrogen using standard glovebox (M-Braun O$_2$ < 0.1 ppm, H$_2$O < 0.1 ppm) and Schlenk line techniques and all glassware were oven and vacuum-dried prior to use. Cp$_2$ZrCl$_2$, Cp$_2$ZrCl$_2$MeLi (1.6 M in Et$_2$O), PCy$_3$, PE$_3$, PPh$_3$, P(Mes)$_3$, and P(C$_6$F$_5$)$_3$ were purchased from Sigma-Aldrich and used as received. [P(C$_6$F$_5$)$_3$]$_2$ was purchased from Acros Organics and used as received. PCy$_3$ was purchased from Sigma-Aldrich and dried prior to use by stirring a hexane solution over CaH$_2$ before removal of the solvent in vacuo and sublimation (25 $^\circ$C, 2 × 10$^{-2}$ Torr). Phenylacetylene was purchased from Sigma-Aldrich and purified by distillation before use. Reagent gases (D$_2$ and CO$_2$) were dried prior to use by passing through a ~78 $^\circ$C trap. Cp$_2$ZrMe$_2$ and Cp$_2$ZrMe$_2$ were synthesized according to literature protocols. Common laboratory solvents (Et$_2$O, DCM, hexane, THF) were purified using a Grubbs type purification system. Nonstandard solvents (chlorobenzene, pentane) were purchased from Sigma-Aldrich and distilled from CaH$_2$ prior to use.

NMR spectra were recorded using JEOL ECP-300 (300 MHz), Varian-400 (400 MHz), and Varian NMR-500 (500 MHz) spectrometers. Deuterated solvents were obtained from Sigma-Aldrich (d$_{6}$-benzene, d$_{6}$-THF, and d$_{2}$-DCM) or Apollo Scientific (d$_{5}$-PhBr) and distilled from CaH$_2$ prior to use. Spectra of air sensitive compounds were recorded using NMR tubes fitted with J. Young valves.
2.4. Synthesis of Zr Lewis acids. Cp₂Zr(Me)-OMes was prepared by a modified literature procedure. In a glovebox, a chlorobenzene (0.5 mL) solution of 1 (30 mg, 0.028 mmol) and an equimolar amount of the corresponding phosphine (0.028 mmol, a = PCy₃ (8.1 mg), b = PEt₃ (3.4 mg), c = PPh₃ (7.6 mg), d = PMes (11.3 mg), e = P(C₆F₅)₃ (14.5 mg)) were weighed out and dissolved in PhCl (0.7 mL) before transferring to an NMR tube fitted with a J. Young valve. Following removal from the glovebox, the sample was subjected to a freeze–pump–thaw degassing cycle prior to refilling with 1 bar D₂ in all cases, no change in the ³¹P NMR spectra was observed following addition of D₂.

3.0. Reactions of FLPs with C₂H₄. In a glovebox, 2 (30 mg, 0.026 mmol) and an equimolar amount of the corresponding phosphine (0.026 mmol, a = PCy₃ (7.1 mg), b = PEt₃ (3.0 mg), c = PPh₃ (6.7 mg), d = PMes (9.9 mg), e = P(C₆F₅)₃ (14.0 mg)) were weighed out and dissolved in PhCl (0.7 mL) before transferring to an NMR tube fitted with a J. Young valve. Following removal from the glovebox, the sample was subjected to a freeze–pump–thaw degassing cycle prior to refilling with 1 bar D₂ in all cases, no change in the ³¹P NMR spectra was observed following addition of D₂.


³¹P NMR (121 MHz, PhCl) δ 33.6 (1:1:1 triplet, Jₚ₋ₚ = 68 Hz, [DCPy*]⁺), ³¹P NMR (46 MHz, PhCl) δ 4.22 (d, Jₚ₋ₚ = 68 Hz, [DCPy*]⁺) 5.98 (s, Zr-D).
Compound 5. H NMR (500 MHz, PhCl) δ 1.02 (9H, m, CH3 (PEt3)), 1.69 (30H, s, CPt), 1.87 (6H, m, CH2 (PEt3)), 1.94 (3H, s, ortho-CH3), 2.00 (3H, s, ortho-CH3), 2.19 (3H, s, para-CH3), 6.67 (2H, s, Ar-CH). 13C NMR (125 MHz, PhCl) δ 11.7 (s, C73.3), 129.7 (s, C), 161.6 (d, δPC (THF)). Compound 6. δ 1.8 (9H, s, C), 3.0 (2H, m, ortho-CH3), 7.6 mg, 3. δ 1.02 (9H, m, CH3 (PEt3)), 1.69 (30H, s, CPt), 1.87 (6H, m, CH2 (PEt3)), 1.94 (3H, s, ortho-CH3), 2.00 (3H, s, ortho-CH3), 2.19 (3H, s, para-CH3), 6.67 (2H, s, Ar-CH). 13C NMR (125 MHz, PhCl) δ 5.4 (d, δPC (THF)) = 4 Hz, 11.8 (d, δPC = 42 Hz), 18.1 and 19.6 (s, ortho-C). 156.2 (s, δPC (Cyclo)), 161.6 (d, δPC = 108 Hz, C(O) O). 31P NMR (121 MHz, PhCl) δ 22.5 (s).

Reactivity of [Cp2ZrOMes][B(C6F5)4]/PhCl (2a-ε). In a glovebox, 2 (30 mg, 0.026 mmol) and an equimolar amount of the corresponding phosphine (0.026 mmol, a = PCy3 (7.1 mg), b = PEt3 (3.0 mg), c = PPh3 (6.7 mg), d = PMes (9.9 mg), e = P(C6P3) (14 mg)) were weighed out and dissolved in PhCl (0.7 mL) before transferring to an NMR tube fitted with a J. Young valve. Following removal from the glovebox, the sample was subjected to a freeze-pump-thaw degassing cycle prior to refilling with 1 bar CO2 via a ~78 °C trap. In the cases of 2a, 2b, and 2c, an immediate color change from red to yellow was observed. In all cases, isolation under 1 bar CO2 was attempted, but was only possible in the case of 6 and in <5% yield. As such all spectral data was obtained in situ.

Compound 6. δ 1.8 (9H, m, CH3 (PEt3)), 1.69 (30H, s, CPt), 1.87 (6H, m, CH2 (PEt3)), 1.94 (3H, s, ortho-CH3), 2.00 (3H, s, ortho-CH3), 2.19 (3H, s, para-CH3), 6.67 (2H, s, Ar-CH). 13C NMR (125 MHz, PhCl) δ 11.7 (s, C73.3), 129.7 (s, C), 161.6 (d, δPC (THF)). Compound 7. Yield = 24 mg (60%). H NMR (500 MHz, PhCl) δ 1.80-2.04 (3H, m, PCy3-β-CH2 and γ-CH2), 2.10 (6H, s, ortho-CH3), 2.21 (3H, s, para-CH3), 2.28 (2H, m, δ-CH2), 4.12 (2H, t, 6JCH = 6 Hz, α-CH2). 6.23 (10H, s, Cp), 6.69 (2H, s, Ar-H). 13C NMR (125 MHz, PhCl) δ 16.3 (d, δPC = 43 Hz, CH2), 12.8 (s, ortho-CH2), 20.7 (α-CH2 = 17.8 Hz, γ-CH2), 20.8 (s, para-CH3), 23.3 (s, δPC (Cyclo)), 27.5 (d, δPC = 12 Hz, meta-CH2 (Cyclo)), 27.8 (d, δPC = 4 Hz, ortho-CH (Cyclo)), 30.8 (d, δPC = 42 Hz, ipso-CH (Cyclo)), 36.7 (d, δPC = 14 Hz, β-CH2), 73.3 (s, α-CH3), 113.7 (s, C), 125.6 (s, para-CH3), 127.6 (s, ortho-C), 129.7 (s, meta-C), 162.0 (s, ipso-C). 31P NMR (121 MHz, PhCl) δ 31.5 (s). ESI-MS (+ve detection) 707.3522 m/z [M]+, 353.2962 m/z [HO(C6H4)PEt3]+.

Compound 8. δ 1.8 (9H, m, CH3 (PEt3)), 1.70 (2H, m, β-CH2) and γ-CH2). 2.09 (6H, s, ortho-CH3), 2.21 (3H, s, para-CH3), 2.24 (2H, m, δ-CH2), 2.27 (6H, m, CH2 (PEt3)), 2.31 (2H, m, δ-CH2), 4.10 (2H, t, 6JCH = 6 Hz, α-CH2). 6.23 (10H, s, Cp), 6.69 (2H, s, Ar-H). 13C NMR (125 MHz, PhCl) δ 5.62 (d, δPC = 5 Hz, CH2 (PEt3)), 12.1 (d, δPC = 49 Hz, CH2 (PEt3)), 18.2 (s, ortho-CH3), 18.9 (d, δPC = 45 Hz, CH2), 19.3 (d, δPC = 5 Hz, γ-CH2), 20.8 (s, para-CH3), 36.2 (d, δPC = 14 Hz, β-CH2), 73.3 (s, α-CH3), 113.7 (s, C), 125.6 (s, para-CH3), 127.6 (s, ortho-C), 129.7 (s, meta-C), 162.0 (s, ipso-C). 31P NMR (121 MHz, PhCl) δ 38.0 (s). ESI-MS (+ve detection) 545.2118 m/z [M]+, 191.1536 m/z [HO(C6H4)PEt3]+.
13C NMR (125 MHz, d5-PhBr) δ 11.7 (s, Cp’), 8.3 (Jp = 50 Hz, 5-CH2), 8.7 (s, para-CH3), 19.9 (s, ortho-CH3), 20.8 (s, ortho-CH3), 22.9 (d, Jp = 4 Hz, 4-CH2), 37.6 (d, Jp = 15 Hz, β-CH2), 70.7 (s, α-CH2), 121.3 (s, Cp’), 128.5 (s, meta-C), 129.6 (d, JpH = 38 Hz, ipso-C (PPh3)), 131.5 (d, Jp = 11 Hz, meta-C (PPh3)), 134.7 (d, Jp = 4 Hz, para-C (PPh3)), 157.9 (s, ipso-C). Ortho-C peak for the triphenylphosphine is obscured by the solvent.

31P NMR (121 MHz, PhCl) δ 22.2 (s, ESI-MS (+ve detection) 829.3684 m/z [M]+, 335.1563 m/z [HO(C6H3)2PPh3]⁺).

Synthesis of [Cp’2Zr(THF)OMes][B(C6F5)4] (2-THF). In a glovebox, THF (0.25 mL) was added dropwise to a stirred chlorobenzene (1 mL) solution of 2 (119 mg, 0.1 mmol), resulting in a yellow solution. The product was isolated via precipitation into a large volume (25 mL) of rapidly stirred hexane. The resulting pale yellow powder was washed with pentane (3 × 5 mL) and dried in vacuo (90 mg, 71%). Crystals of 2-THF suitable for analysis by single crystal X-ray diffraction were obtained by layering a chlorobenzene solution with pentane (7 days).

1H NMR (500 MHz, d5-PhBr) δ 1.59 (4H, s, THF (C3-C4)), 1.61 (30H, s, Cp’), 1.79 (3H, s, ortho-CH3), 1.88 (3H, s, ortho-CH3), 2.19 (3H, s, para-CH3), 3.55 (4H, s, THF (C2-C5)), 6.03 (1H, s, Ar-H), 6.77 (1H, s, Ar-H). 13C NMR (125 MHz, d5-PhBr) δ 10.2 (s, Cp’-Me), 17.0 (s, ortho-CH3), 18.3 (s, ortho-CH3), 20.1 (s, para-CH3), 25.0 (s, THF (C3-C4)), 67.9 (s, THF (C2-C5)), 122.0 (s, Cp’), 129.4 (s, meta-C), 154.8 (s, ipso-C). NB: Remaining peaks in 13C NMR are obscured by the PhBr solvent.

4.8. Reaction of Pairs with Phenylacetylene (PhCCH). Reactivity of [Cp’ZrOMes][B(C6F5)4]/PhP3 (1a-e). In a glovebox, 1 (30 mg, 0.028 mmol) and an equimolar amount of the corresponding phosphine (0.028 mmol, a = PCy3 (8.1 mg), b = PEt3 (3.4 mg), c = PPh3 (7.6 mg), d = PMes3 (11.3 mg), e = PCF3(5) (15.4 mg)) were weighed out and dissolved in PhCl (0.7 mL) before transferring an NMR tube fitted with a J. Young valve. Excess phenylacetylene (5 drops) was subsequently added, and in the case of 1a-d an instantaneous lightening of the yellow color was observed. The progress of the reactions was monitored by 31P NMR spectroscopy. Collected spectral data is detailed below.

1a. Reaction complete in <1 min. Mixture of products could not be sufficiently separated to allow further characterization.

1b. Reaction complete in 16 h. Mixture of products could not be sufficiently separated to allow further characterization.

1c. Reaction complete in <1 min, and compound 13 was isolated in a glovebox by precipitation into rapidly stirred hexane (20 mL) and washed with pentane (3 × 5 mL) before drying in vacuo (24.0 mg, 92%). Crystals of 14 suitable for analysis by single crystal X-ray diffraction were obtained by layering a PhCl solution of 14 with pentane (5 days).

1H NMR (500 MHz, d5-PhBr) δ 1.59 (30H, s, Cp’), 1.17 (3H, s, para-CH3), 2.10 (3H, s, ortho-CH3), 2.12 (3H, s, ortho-CH3), 6.40 (2H, s, Ar-H), 7.50–7.76 (15H, m, PPh3), 8.36 (1H, d, JpH = 45 Hz, α-H). 13C NMR (125 MHz, d5-PhBr) δ 15.4 (s, Cp’), 17.5 (s, para-CH3), 22.9 and 23.6 (s, ortho-CH3), 124.8 (Cp’), 128.1 (s, para-CH3), 131.7 (s, ortho-CH3), 132.5 (s, meta-C), 136.5 (d, JpH = 14 Hz, meta-C (PPh3)), 137.6 (d, JpH = 4 Hz, para-C (PPh3)), 158.9 (ipso-C), 233.1 (d, JpH = 10 Hz, Zr-CH(C)). 31P NMR (121 MHz, d5-PhBr) δ 17.4 (d, JpH = 48 Hz, 14). NB: Remaining peaks in 31P NMR are obscured by the PhBr solvent.

1d. Reaction complete in <1 min. In situ analysis of the reaction mixture by 31P NMR spectroscopy showed clean conversion to deprotonation products; however, the zirconium acetylide complex could not be isolated cleanly.

1e. No reaction was evident by 31P NMR spectroscopy.


(10) Dehydrocoupling of amine-boranes has been achieved stoichiometrically with main group systems, but never in a catalytic sense. (a) Miller, A. J. M.; Bercaw, J. E. Chem. Commun. 2010, 46, 1709.


