

CrossMark  
click for updatesCite this: *Chem. Sci.*, 2016, 7, 921

# Mechanistic studies on the addition of hydrogen to iridaepoxide complexes with subsequent elimination of water†

Lauren E. Doyle, Warren E. Piers,\* Javier Borau-Garcia, Michael J. Sgro and Denis M. Spasyuk

Iridium complexes of the PC<sub>sp</sub><sup>2</sup>P ligand in which the donors are linked by 2,3-benzo[*b*]thiophene groups engage in the cooperative activation of N<sub>2</sub>O and the resulting iridaepoxides can be treated with dihydrogen to effect elimination of water and regeneration of the starting iridium complex. The mechanism of the steps in this reaction have been investigated using low temperature NMR investigations that reveal H/D exchange processes that point to a highly reactive kinetic product of hydrogen addition to the iridaepoxide. This intermediate is also involved in the water elimination pathway, and model compounds have been synthesized to provide further evidence for the mechanistic proposals for water elimination. The adaptable donor properties of the PC<sub>sp</sub><sup>2</sup>P ligand framework, particularly the anchoring carbene donor, plays a significant role in the ability of these compounds to mediate the transformation of N<sub>2</sub>O in this way.

Received 21st September 2015

Accepted 26th October 2015

DOI: 10.1039/c5sc03575a

www.rsc.org/chemicalscience

## Introduction

Tridentate “pincer” ligand frameworks continue to attract interest for supporting complexes of metals from across the periodic table that serve as platforms for small molecule activation and selective bond cleavage.<sup>1,2</sup> Their appeal stems from the large number of permutations for EE'E'' donor arrays, the extensive electronic and steric tunability possible *via* modification of the linking features of the ligand framework, and the opportunity to design ligands of formal charge ranging from 0 to -3. Furthermore, through incorporation of functions capable of accepting and delivering hydrogen atoms or protons, ligand cooperativity can feature prominently in catalytic cycles mediated by pincer ligand metal complexes.<sup>3</sup>

Recently, we introduced a variation on the venerable PCP<sup>4</sup> motif that incorporates a diaryl carbene donor in the central anchoring position of the pincer framework,<sup>5</sup> exemplified by ligands **I** and **II**.<sup>6,7</sup> These ligands are not only quite electron rich, but allow for the possibility of cooperativity through engagement of the M=C unit of the chelate array.<sup>8</sup> Furthermore, depending on the metal employed, the carbene unit in the ligand assumes either Schrock-like<sup>9-13</sup> or Fischer-like behavior.<sup>14,15</sup> Thus, in nickel complexes, the carbene is highly

nucleophilic, whereas in iridium manifolds, it is less so. This “continuum of reactivity” can be exploited in the deployment of such ligands for small molecule activation.<sup>16</sup>



Despite the lowered nucleophilicity of the carbene carbon, the iridium complexes of ligands of type **I** or **II** react with N<sub>2</sub>O to afford “iridaepoxide” complexes in which the oxygen atom has added to the Ir=C units.<sup>7</sup> In compounds of ligand type **I**, C-C bond activations<sup>17</sup> ensue from the initially observed iridaepoxide, which eventually leads to the dismantling of the pincer ligand framework.<sup>18</sup> However, the rigidity imparted by the 2,3-benzo[*b*]thiophene linker in ligand **II** raises the barrier to this undesired ruination of the ligand framework, allowing for other chemistry to be explored. For instance, reaction of iridaepoxide **1·Cl** with H<sub>2</sub> gives an iridium dihydride complex, **2·Cl<sub>trans</sub>**, where the two hydride ligands reside in positions *trans* to the iridaepoxide moiety (Scheme 1). Upon heating to ≈ 100 °C, this compound eliminates water to return to the iridium carbene complex **3·Cl**. Attempts to garner information concerning the mechanism of water loss were to some degree hampered by the fact that dihydride **2·Cl<sub>trans</sub>** is a thermodynamic isomer that we proposed must convert to a kinetic isomer **2·Cl<sub>cis</sub>** in which the hydride ligands are *cis*-disposed with respect to the iridium oxygen bond of the iridaepoxide unit. The reasons for this

University of Calgary, Department of Chemistry, 2500 University Drive N.W., Calgary, Alberta, T2N 1N4, Canada. E-mail: wpiers@ucalgary.ca

† Electronic supplementary information (ESI) available: Description of general methods, Fig. S1-3, NMR spectra, crystallographic data for **3·OH**, **1·OH**, **1·Br**, **7·Cl<sub>3</sub>**, **7·(OTf)<sub>2</sub>Cl** and **8**. CCDC 1423154-1423159. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c5sc03575a





Scheme 1 Hydrogen addition and water elimination to iridaepoxide 1-Cl.

kinetic preference are not clear, but they may be related to the seminal studies of Eisenberg on the preferred paths for addition of H<sub>2</sub> to Vaska's complex along a P–Ir–CO vector instead of P–Ir–Cl.<sup>19</sup> Regardless, the kinetic data we obtained by following the loss of water from 2·Cl<sub>trans</sub> provided information on the loss of H<sub>2</sub> from this compound, which is the rate limiting step in the loss of water, and the process by which H<sub>2</sub>O was eliminated was essentially opaque to these kinetic investigations.<sup>7</sup> Here, we provide more evidence for this proposal through a combination of labeling studies, low temperature NMR investigations and model synthesis of likely intermediates in the water elimination path.

## Results and discussion

The structure and properties of compound 2·Cl<sub>trans</sub> were reported in our preliminary communication.<sup>7</sup> To summarize, it is stable in toluene solution at temperatures ranging from 25–80 °C, but EXSY spectroscopy indicates that the two diastereotopic hydride ligands (characterized as triplets in the <sup>1</sup>H NMR spectrum at –7.58 and –15.22 ppm) undergo exchange even at room temperature on the timescale of these experiments. However, there is no exchange with free H<sub>2</sub> at room temperature as measured by exposure of a solution of 2·Cl<sub>trans</sub> to D<sub>2</sub>; only when temperatures reach 80–90 °C does label begin to appear in these positions, but as indicated above, water elimination also occurs in this temperature regime.

In order to assess the results of this exchange experiment, the spectroscopic signature of 2·Cl<sub>trans</sub>-d<sub>2</sub> was obtained on a separately prepared sample obtained by treatment of 1·Cl with D<sub>2</sub>. The <sup>31</sup>P{<sup>1</sup>H} NMR chemical shift of 24.1 ppm for the dideuterated isotopologue is almost baseline resolved from that of undeuterated 2·Cl<sub>trans</sub> at 24.0 ppm in the 162 MHz spectrum. We therefore sought to assess the kinetic isotope effect for the oxidative addition of H<sub>2</sub> to 1·Cl by reacting it with 1 atm of

a 1 : 1 mixture of H<sub>2</sub>/D<sub>2</sub> at room temperature. Surprisingly, rather than the expected mixture of 2·Cl<sub>trans</sub> and 2·Cl<sub>trans</sub>-d<sub>2</sub>, both the <sup>31</sup>P{<sup>1</sup>H} and the <sup>1</sup>H{<sup>31</sup>P} NMR spectra revealed that a near-statistical mixture of the four possible isotopologues (Fig. 1a) was observed (Fig. 1b, middle trace); the presence of the d<sub>2</sub> isotopologue was confirmed by <sup>2</sup>H and <sup>31</sup>P{<sup>1</sup>H} NMR spectroscopy. Furthermore, the presence of significant quantities of HD was detected in the <sup>1</sup>H NMR spectrum (4.47 ppm, <sup>1</sup>J<sub>HD</sub> = 43 Hz, 1 : 1 : 1 triplet). This puzzling result was



Fig. 1 (a) Reaction of compound 1·Cl with either a 1 : 1 mixture of H<sub>2</sub>/D<sub>2</sub> or pure HD leads to the mixture of isotopologues of compound 2·Cl<sub>trans</sub>. (b) Hydride region of the <sup>1</sup>H{<sup>31</sup>P} NMR spectrum of pure 2·Cl<sub>trans</sub> (top), the reaction mixture when 1·Cl is treated with H<sub>2</sub>/D<sub>2</sub> (middle) and the reaction mixture when 1·Cl is treated with HD (bottom). Note that, while the outer resonances in each pair are due to the 2·Cl<sub>trans</sub>-d<sub>1</sub> isotopomers, we have not made absolute assignments for these resonances.



corroborated by a complementary experiment in which the reaction of  $1\cdot\text{Cl}$  with pure HD gave the same mixture of four isotopologues (Fig. 1b, bottom trace), along with detectable amounts of  $\text{H}_2$  in the  $^1\text{H}$  NMR spectrum (singlet, 4.51 ppm).<sup>‡</sup> It was further observed in a separate experiment that equimolar mixtures of  $2\cdot\text{Cl}_{\text{trans}}$  and  $2\cdot\text{Cl}_{\text{trans}}\text{-d}_2$  exhibited *no scrambling* of the label to form either of the two  $2\cdot\text{Cl}_{\text{trans}}\text{-d}_1$  isotopomers unless heated to above  $80^\circ\text{C}$ —the temperature regime in which hydrogen/deuterium elimination from  $2\cdot\text{Cl}_{\text{trans}}\text{-d}_n$  begins. Since the scrambling to form the four isotopologues of  $2\cdot\text{Cl}_{\text{trans}}$  occurred in reactions done at *room temperature*, this requires that the isotope redistribution in the  $\text{H}_2/\text{D}_2$  mixture and the pure HD must be occurring *prior* to final formation of the thermodynamic isomer  $2\cdot\text{Cl}_{\text{trans}}$ .

Given this conclusion, we postulated that these rapid H/D scrambling processes are mediated by the proposed kinetic isomer  $2\cdot\text{Cl}_{\text{cis}}$ . As briefly discussed previously,<sup>7</sup> this kinetic isomer can be observed spectroscopically at low temperatures ( $-38$  to  $-78^\circ\text{C}$ ) *via* both  $^1\text{H}$  and  $^{31}\text{P}$  NMR experiments by treatment of  $1\cdot\text{Cl}$  with  $\approx 2\text{--}3$  atm of  $\text{H}_2$  at  $-78^\circ\text{C}$ . As long as the temperature remains below  $-38^\circ\text{C}$ , only  $1\cdot\text{Cl}$  and  $2\cdot\text{Cl}_{\text{cis}}$  are observed; the thermodynamic isomer only begins to appear if the sample is warmed above this temperature. In the low temperature regime,  $1\cdot\text{Cl}$ ,  $\text{H}_2$  and  $2\cdot\text{Cl}_{\text{cis}}$  are in rapid equilibrium, as indicated by a changing ratio of  $1\cdot\text{Cl}$  and  $2\cdot\text{Cl}_{\text{cis}}$  and the broadening of the resonances for  $2\cdot\text{Cl}_{\text{cis}}$  in the  $^{31}\text{P}$  NMR spectrum as the temperature is raised. At  $-78^\circ\text{C}$ , the equilibrium constant for the addition of  $\text{H}_2$  to  $1\cdot\text{Cl}$  to form  $2\cdot\text{Cl}_{\text{cis}}$  is  $145\text{ M}^{-1}$  and the equilibrium constant can be evaluated at various temperatures and treated using a van't Hoff analysis (Fig. S1†). The thermodynamic parameters thus derived indicate that, while enthalpically favored ( $\Delta H^\circ = -4.9(3)\text{ kcal mol}^{-1}$ ), the entropic penalty ( $\Delta S^\circ = -15(2)\text{ cal mol}^{-1}\text{ K}^{-1}$ ) renders the formation of  $2\cdot\text{Cl}_{\text{cis}}$  at room temperature essentially thermoneutral ( $\Delta G_{298}^\circ = -0.4(5)\text{ kcal mol}^{-1}$ ), completely consistent with our qualitative observations. The equilibrium constant measured at  $-78^\circ\text{C}$  when substituting deuterium for hydrogen, is larger ( $K_{\text{eq}}(\text{D}_2) = 590\text{ M}^{-1}$ ), and a substantial inverse equilibrium isotope effect (EIE) of 0.37 is obtained. The observation of an inverse EIE in oxidative addition of  $\text{H}_2/\text{D}_2$  is expected in this low temperature regimen, and the reasons for this phenomenon have been lucidly rationalized by Parkin and co-workers.<sup>20–22</sup>

With this behavior in mind, in order to probe for hydrogen/deuterium scrambling in the kinetic dihydrido compound,  $1\cdot\text{Cl}$  was treated with a mixture of  $\text{H}_2$  and  $\text{D}_2$  ( $\approx 1.4 : 1$ ) at  $-78^\circ\text{C}$ . The 243 MHz  $^{31}\text{P}\{^1\text{H}\}$  spectrum at the top of Fig. 2 is observed initially; here, the broad resonance for iridaepoxide  $1\cdot\text{Cl}$  at approximately 25.1 ppm is accompanied by two sharper, baseline resolved signals at 23.25 and 23.09 ppm for the  $2\cdot\text{Cl}_{\text{cis}}\text{-d}_2$  and  $2\cdot\text{Cl}_{\text{cis}}$  isotopologues, respectively.<sup>§</sup> This spectrum remains constant at this temperature, but as the sample is warmed, the chemical shifts for the isotopologues of  $2\cdot\text{Cl}_{\text{cis}}$  drift downfield and the signals broaden slightly. Furthermore, a new resonance of intermediate chemical shift begins to appear (Fig. 2,  $-58^\circ\text{C}$  spectrum). At  $-38^\circ\text{C}$ , the amount of  $2\cdot\text{Cl}_{\text{cis}}\text{-d}_n$  is significantly diminished as the equilibrium shifts to the left and the pattern of the signal is difficult to discern; however, if the sample is re-



Fig. 2 Reaction of compound  $1\cdot\text{Cl}$  with a mixture of  $\text{H}_2/\text{D}_2$  at  $-78^\circ\text{C}$ , followed by warming to  $-38^\circ\text{C}$  and then re-cooling to  $-78^\circ\text{C}$  as monitored by  $^{31}\text{P}\{^1\text{H}\}$  NMR spectroscopy at 243 MHz in  $\text{CD}_2\text{Cl}_2$ . The equilibrium between  $1\cdot\text{Cl}$  and the  $(\text{H})_2$  and  $(\text{D})_2$  isotopologues of  $2\cdot\text{Cl}_{\text{cis}}$  is apparent in the  $t=0$  spectrum; warming results in production of the  $(\text{H})(\text{D})$  isotopomers, which are resolved in the  $-78^\circ\text{C}$  spectrum obtained upon re-cooling to low temperature.

cooled to  $-78^\circ\text{C}$ , and the speciation of the equilibrium shifts towards the addition product, the  $^{31}\text{P}\{^1\text{H}\}$  NMR spectrum clearly displays signals for all four possible isotopologues of the kinetic isomer  $2\cdot\text{Cl}_{\text{cis}}\text{-d}_n$ . Also of significance is the observation that the  $^{31}\text{P}$  resonances for the separate  $2\cdot\text{Cl}_{\text{cis}}\text{-d}_1$  isotopomers coalesce at temperatures above  $-78^\circ\text{C}$  into one averaged signal, indicating that these two species are in rapid exchange, likely *via* an  $\eta^2\text{-HD}$  complex (III, Fig. 2, inset). Furthermore, in the  $^1\text{H}$  NMR spectrum obtained for this sample ( $-78^\circ\text{C}$ ), in addition to the singlet for  $\text{H}_2$  at 4.55 ppm, the distinct, isotopically shifted 1 : 1 : 1 triplet for HD is observed at 4.52 ppm.<sup>¶</sup> This experiment clearly establishes  $2\cdot\text{Cl}_{\text{cis}}$  as the agent of the facile isotope scrambling in the iridium dihydrides and the redistribution of label in the headspace gas as observed in the experiments depicted in Fig. 1.



Given the clear implication of the kinetic isomer  $2 \cdot \text{Cl}_{\text{cis}}$  in the observed H/D scrambling, and its potential involvement in the process by which water is eliminated,<sup>7</sup> we sought to probe the effect on its formation as a function of the X type ligand present in  $1 \cdot \text{Cl}$ . It is well-established that the exothermicity of hydrogen addition to Vaska's complex is strongly influenced by the nature of the X ligand,<sup>23</sup> and that the addition product is favored as the halide is changed from  $\text{F} < \text{Cl} < \text{Br} < \text{I}$ .<sup>24,25</sup> Since compound  $1 \cdot \text{Cl}$  is phenomenologically similar to Vaska's complex, we postulated that the position of the equilibrium for formation of the kinetic  $\text{H}_2$  addition product might be influenced by changing this group. We therefore synthesized the hydroxo and bromo derivatives shown in Scheme 2 and measured the equilibrium constants for formation of the dihydrido derivatives  $2 \cdot \text{OH}_{\text{cis}}$  and  $2 \cdot \text{Br}_{\text{cis}}$ . The carbene hydroxo complex  $3 \cdot \text{OH}$  was prepared using cesium hydroxide as a reagent.<sup>15,26</sup> This species was then used to prepare the bromo derivative  $3 \cdot \text{Br}$  *via* treatment with LiBr. Both compounds exhibit characteristic triplet resonances ( $^2J_{\text{CP}} = \approx 2.5 \text{ Hz}$ ) in the  $^{13}\text{C}$  NMR spectra at 172.2 ( $3 \cdot \text{OH}$ ) and 161.4 ( $3 \cdot \text{Br}$ ) ppm for the carbene carbon of the  $\text{PC}_{\text{sp}^2}\text{P}$  ligand. Compound  $3 \cdot \text{OH}$  was further characterized by X-ray crystallography, the first example of this family of monomeric hydroxo compounds to be structurally characterized; its molecular structure is shown in Fig. 3 along with selected metrical parameters. The ligand adopts a nearly planar geometry, and features an Ir=C bond distance of 1.943(5) Å. The Ir–O distance of 2.004(4) Å is between those of 1.978(12) Å and 2.020(6) Å found in other four coordinate, monomeric iridium hydroxo compounds.<sup>27–29</sup> Both the carbene



Fig. 3 Molecular structure of  $3 \cdot \text{OH}$ . Hydrogen atoms have been omitted for clarity. Displacement ellipsoids are shown at the 50% probability level. Selected bond lengths (Å): Ir(1)–P(1), 2.2904(13); Ir(1)–P(2), 2.2817(13); Ir(1)–C(1), 1.943(5); Ir(1)–O(1), 2.004(4). Selected bond angles (°): C(1)–Ir(1)–O(1), 175.5(2); P(1)–Ir(1)–P(2), 166.08(5).

hydroxo and bromo compounds react smoothly with  $\text{N}_2\text{O}$  to yield the corresponding iridaepoxides  $1 \cdot \text{OH}$  and  $1 \cdot \text{Br}$ , as shown in Scheme 2 and each of these compounds has also been characterized crystallographically. Fig. 4 shows the molecular structure of one of the two molecules found in the unit cell for the hydroxo complex; metrical parameters are essentially identical for both molecules. The molecular structure of  $1 \cdot \text{Br}$  can be found in Fig. S2.† Metrical parameters within the series of compounds  $1 \cdot \text{X}$  are broadly similar, with the exception of those distances and angles pertaining to the X group. Notably, the C(1)–O(1) distance within the iridaepoxide unit is virtually



Scheme 2 Synthesis of compounds  $1 \cdot \text{X}$  and addition of  $\text{H}_2$ .



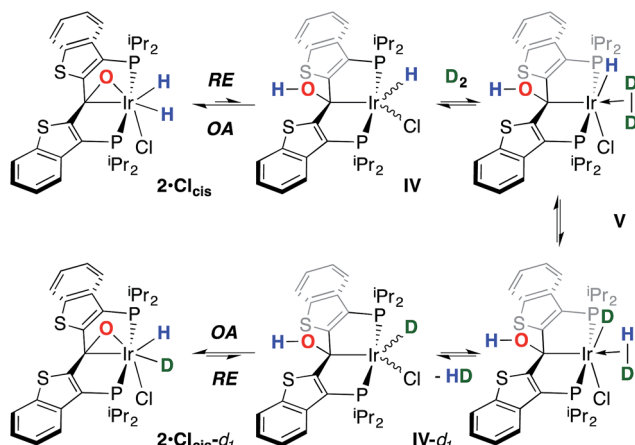
Fig. 4 Molecular structure of  $1 \cdot \text{OH}$ . Hydrogen atoms have been omitted for clarity. Displacement ellipsoids are shown at the 50% probability level. Selected bond lengths (Å): Ir(1)–P(1), 2.3110(10); Ir(1)–P(2), 2.3135(10); Ir(1)–C(1), 2.078(4); Ir(1)–O(1), 2.084(3); Ir(1)–O(2), 1.972(4); C(1)–O(1), 1.359(5). Selected bond angles (°): C(1)–Ir(1)–O(1), 38.12(13); C(1)–O(1)–Ir(1), 70.7(2); Ir(1)–C(1)–O(1), 71.2(2); P(1)–Ir(1)–P(2), 166.23(4).



unchanged within these derivatives, indicating little effect on the C–O bond order as X is changed.

With these compounds in hand, reactions with H<sub>2</sub> were studied at both –78 °C and at ambient temperatures. The hydroxo complex **1**·OH exhibits no reaction with H<sub>2</sub> at –78 °C, indicating that the equilibrium constant for formation of a hydrogen oxidative addition product, **2**·OH<sub>cis</sub>, analogous to the chloro case discussed above, is very small. When the temperature of this sample is warmed to >45 °C, a reaction ensues that produces two equivalents of water and a polyhydride PC<sub>sp</sub><sup>3</sup>P complex, **4** Scheme 2, that is analogous to several such compounds reported in the literature.<sup>30</sup> Compound **4** was only stable under an atmosphere of dihydrogen and so was characterized *in situ* by multinuclear NMR spectroscopy. This was further complicated by the observation of partial deuteration (*via* the solvent, toluene-*d*<sub>8</sub>) in the ligand aromatic and benzylic positions, as well as the Ir–H moieties,<sup>30–32</sup> under the high temperature conditions under which it forms. Nevertheless, the observed generation of **4** and water suggests that H<sub>2</sub> oxidative addition to **1**·OH does occur at higher temperatures, but facile O–H reductive eliminations<sup>33,34</sup> ensue that eventually lead to the observed products. In contrast, compound **1**·Br exhibits behavior more akin to that of **1**·Cl, forming a kinetic isomer at low temperatures and an isolable thermodynamic isomer at higher temperatures (Scheme 2). The equilibrium constant measured for the formation of **2**·Br<sub>cis</sub> is about 4 times larger than that found for the chlorido derivative, indicating that the exothermicity of hydrogen oxidative addition to these compounds follows a similar trend to that observed in the Vaska's complex series.<sup>22</sup> The greater stability of **2**·Br<sub>cis</sub> is reflected in the much slower release of water from this derivative. At 110 °C, elimination of H<sub>2</sub>O is extremely slow, and complete conversion is only observed when heating in *o*-xylene at 150 °C for 2–3 hours.

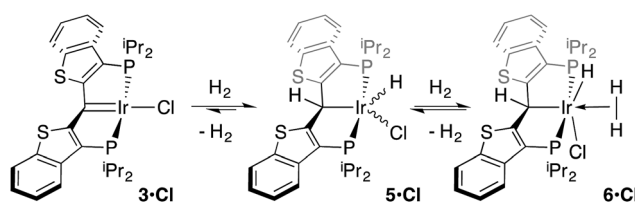
While the kinetic isomer **2**·Cl<sub>cis</sub> is thus strongly implicated in the observed H/D scrambling, the mechanism by which it mediates this facile process is a matter of some speculation. A further question regarding the relationship, if any, between the label redistribution process and the path by which water elimination from **2**·Cl (*trans* or *cis*) occurs is also significant. In our initial communication, we postulated that, because the Ir–O bond of the iridaepoxide moiety and an Ir–H bond are now *cis* to one another in the kinetic isomer, it is plausible that reductive elimination of O–H<sup>33,34</sup> could be a first step towards overall release of water from **2**·Cl<sub>cis</sub> to form the carbene chloride **3**·Cl (Scheme 1).<sup>7</sup> We believe that the H/D scrambling observed is consistent with this proposal (Scheme 3). Reductive elimination of O–H from **2**·Cl<sub>cis</sub> leads to the hydrido chloride Ir(III) species **IV** bearing an alcohol function on the anchoring carbon of a PC<sub>sp</sub><sup>3</sup>P pincer ligand. This compound, while not observed here, is directly analogous to the product formed *via* a single C–H bond activation in the ligand attachment chemistry to iridium for these ligands—lacking the OH function, of course. These five coordinate Ir(III) compounds are highly fluxional,<sup>6,30,35,36</sup> and have been shown to bind H<sub>2</sub> to form η<sup>2</sup>-dihydrogen hydrido complexes in which all three hydrogens are in rapid exchange; the specific case involving ligand **II** is no



Scheme 3 Proposed path for H/D scrambling mediated by **2**·Cl<sub>cis</sub>.

exception (Scheme 4). Treatment of the carbene chloride complex **3**·Cl with an atmosphere of H<sub>2</sub> gives an equilibrium mixture of compounds **5**·Cl and **6**·Cl; in the presence of H<sub>2</sub>, the equilibrium lies far to the side of dihydrogen complex **6**·Cl. This species is characterized by a broad resonance at –5.35 ppm for the η<sup>2</sup>-H<sub>2</sub> ligand and a hydride signal at –21.7 ppm, also broad. When the H<sub>2</sub> atmosphere is removed, compound **5**·Cl is isolable as an analytically pure orange solid. Treatment of **5**·Cl with D<sub>2</sub> regenerates **6**·Cl as its *d*<sub>1</sub>-isotopologue; exchange of D into the benzylic position of the PC<sub>sp</sub><sup>3</sup>P ligand is also observed as would be expected. The rapid exchange process for the hydrogen/deuterium atoms in **6**·Cl is also likely to be present in **V**, providing a valid mechanism for the scrambling observed when **IV** comes into contact with D<sub>2</sub>; exchange leads not only to **IV**-*d*<sub>1</sub>, but the HD that is also observed in these reactions (Scheme 3). Oxidative addition of the PC<sub>sp</sub><sup>3</sup>P ligand O–H bond regenerates **2**·Cl<sub>cis</sub> as its *d*<sub>1</sub> isotopologue; as the spectra in Fig. 2 demonstrate, the diastereotopic positions in this compound are in rapid exchange even at –68 °C.

Overall, this proposal accounts for the statistical redistribution of isotopes observed (as opposed to the pairwise exchange of H<sub>2</sub> and D<sub>2</sub>, for example), but necessitates that reductive elimination/oxidative addition of O–H is rapidly reversible under these conditions. Furthermore, experimental observations suggest that if **2**·Cl<sub>cis</sub> is in equilibrium with **IV**, the equilibrium favors the observed dihydrido species. Close examination of the spectra shown Fig. 2 indicate that, as the sample containing **2**·Cl<sub>cis</sub> and **2**·Cl<sub>cis</sub>-*d*<sub>2</sub> is warmed from –78 °C to –38 °C, the diproteo isotopologue disappears faster than the dideutero isotopologue, providing qualitative evidence for



Scheme 4 Addition of H<sub>2</sub> to **3**·Cl.



a normal kinetic isotope effect in the rate limiting step of the process leading to isotope scrambling. This is consistent with expectations for an O–H(D) reductive elimination initial step. It is also consistent with an alternative view of  $2 \cdot \text{Cl}_{\text{cis}}$  as an  $\eta^2$ -ketone complex (Scheme 5); here, the reversible conversion to **IV** would be viewed as a 1,2-insertion of the C=O double bond into the *cis* disposed hydrido ligand and the reverse process,  $\beta$ -elimination. The metrical data obtained for the structure of  $2 \cdot \text{Cl}_{\text{trans}}$ , specifically the C–O distance of 1.303(12) Å, and the downfield resonance for the iridaepoxide carbon (99.5 ppm) is consistent with the  $\eta^2$  ketone form contributing significantly to the structural description of this compound. Our experimental data do not distinguish between these two views of such a process, but both, we believe, are reasonable suggestions to explain the observed H/D scrambling.

In this mechanistic model, the H/D scrambling observed proceeds *via* the key species **IV** and we proposed that the path to water elimination also originates from this species.<sup>7</sup> While the H/D scrambling occurs with a very low barrier, since elimination of water only occurs at significantly higher temperatures, the rate limiting step for that process must have a much higher barrier. Cleavage of the C–O bond in **IV** *via* migration of the OH group in **IV** to give the putative  $\text{PC}_{\text{sp}^2}\text{P}$  iridium(III) hydroxo hydrido chloride complex **VI** (Scheme 6) would be expected to have a higher barrier.<sup>37,38</sup> Once **VI** is formed, the facility with which O–H bond reductive elimination occurs in these systems renders this species unstable towards rapid water elimination, giving  $3 \cdot \text{Cl}$ . Such a scenario is supported by the observation and characterization of model compounds for the six coordinate Ir(III) intermediate **VI**. For example, the  $\text{PC}_{\text{sp}^2}\text{P}$  carbene chloride  $3 \cdot \text{Cl}$  can be treated with an excess of hexachloroethane ( $\text{C}_2\text{Cl}_6$ ) at 50 °C to yield the highly insoluble trichloride complex  $7 \cdot \text{Cl}_3$  in excellent isolated yield (Scheme 6). The insolubility of this species in most common solvents made full spectroscopic characterization difficult, but the solid obtained is pure by elemental analysis and an X-ray crystal structure analysis confirms its identity (Fig. 5). The iridium center exhibits octahedral geometry and the Ir(1)–C(1) distance of 1.938(8) Å is slightly longer than the distances we have observed for these



Scheme 6 Proposed mechanism for water elimination from  $2 \cdot \text{Cl}_{\text{cis}}$ .

ligands in most square planar, Ir(I) derivatives. The Ir(1)–Cl(2) bond length (*trans* to the carbene donor) is significantly longer at 2.428(2) Å than the lengths recorded for the *cis* chlorides (2.348(2) and 2.356(2) Å).

A second, more soluble member of this family of compounds can be made by treatment of the iridaepoxide complex  $1 \cdot \text{Cl}$  with two equivalents trimethylsilyl triflate. This reaction proceeds smoothly at room temperature to form the  $\text{PC}_{\text{sp}^2}\text{P}$  iridium(III) bis triflate chloride compound  $7 \cdot (\text{OTf})_2\text{Cl}$  (Scheme 7) along with one equivalent of hexamethyldisiloxane. Interestingly, use of only one equivalent of TMSOTf yields a 1 : 1 mixture of  $1 \cdot \text{Cl}$  and  $7 \cdot (\text{OTf})_2\text{Cl}$ , indicating that reaction of the second equivalent of TMSOTf with the product of the reaction of  $1 \cdot \text{Cl}$  with the first equivalent is very rapid. In any case, because of its greater solubility, compound  $7 \cdot (\text{OTf})_2\text{Cl}$  was fully characterized spectroscopically and features a highly downfield shifted signal at



Scheme 5 Two views of the conversion of  $2 \cdot \text{Cl}_{\text{cis}}$  to **IV**.



Fig. 5 Molecular structure of  $7 \cdot \text{Cl}_3$ . Hydrogen atoms have been omitted for clarity. Displacement ellipsoids are shown at the 50% probability level. Selected bond lengths (Å): Ir(1)–P(1), 2.343(2); Ir(1)–P(2), 2.346(2); Ir(1)–C(1), 1.938(8); Ir(1)–Cl(1), 2.348(2); Ir(1)–Cl(2), 2.428(2); Ir(1)–Cl(3), 2.356(2). Selected bond angles (°): C(1)–Ir(1)–Cl(1), 92.0(2); C(1)–Ir(1)–Cl(2), 178.6(2); C(1)–Ir(1)–Cl(3), 90.3(2); P(1)–Ir(1)–P(2), 170.47(8).





Scheme 7 Model reactions for water elimination.

219.1 ppm (the  $^2J_{\text{CP}}$  is unresolved) for the carbene carbon, and two distinct resonances at  $-78.8$  and  $-79.1$  ppm in the  $^{19}\text{F}$  NMR spectrum, indicating that the two triflate ions are diastereotopic. A solid state structural analysis on single crystals of this species confirms the geometry of the complex as drawn in Scheme 7; the molecular structure is given in Fig. 6. The structure is similar to that of trichloride  $7\cdot\text{Cl}_3$  but the ligand backbone exhibits some torsion between the planes of the benzothiophene linkers, possibly due to the more bulky triflate ligands interacting with the phosphine iso-propyl groups. The Ir(1)–C(1) distance in  $7\cdot(\text{OTf})_2\text{Cl}$  is slightly longer at  $1.948(7)$  Å and the differences in the Ir(1)–O(1) ( $2.197(5)$ ) and O(4) ( $2.131(5)$ ) distances are less pronounced than the analogous Ir–Cl distances in  $7\cdot\text{Cl}_3$ .

Compound  $7\cdot(\text{OTf})_2\text{Cl}$  is significant because it is constitutionally related to the compound we obtained upon double

protonation of  $1\cdot\text{Cl}$  with triflic acid.<sup>7</sup> This product, **8** in Scheme 7, although obtained very cleanly upon treatment of  $1\cdot\text{Cl}$  with HOTf at  $-20$  °C in methylene chloride, was not previously isolable and so a precise structural assignment was not possible at the time of the original report.<sup>7</sup> Since  $7\cdot(\text{OTf})_2\text{Cl}$  is related to this species by the addition or elimination of one equivalent of water, we attempted to generate **8** through treatment of  $7\cdot(\text{OTf})_2\text{Cl}$  with stoichiometric amounts of water. These experiments met with mixed success, but we found that stirring  $7\cdot(\text{OTf})_2\text{Cl}$  in  $\text{CD}_2\text{Cl}_2$  that had not undergone our normal rigorous drying procedure, eventually generated clean samples of compound **8**. Water free  $7\cdot(\text{OTf})_2\text{Cl}$  was slowly regenerated by adding a few activated molecular sieves to this sample; this interconversion is demonstrated by the series of  $^{31}\text{P}$  NMR spectra shown in Fig. S3.† We were also able to generate **8** by doubly protonating  $1\cdot\text{Cl}$  with HOTf in wet  $\text{CD}_2\text{Cl}_2$  and this procedure led to the growth and isolation of single crystals; the X-ray structural determination confirmed the structure of this species as that shown in Scheme 7. Fig. 7 depicts the molecular structure of **8**, along with an outer sphere triflate anion from an adjacent unit cell, which exhibits hydrogen bonding with one of the aquo ligand protons as shown. Here, it is the triflate anion located *cis* to the carbene carbon in  $7\cdot(\text{OTf})_2\text{Cl}$  that has been formally displaced by the aquo ligand. Presumably the mechanism of this substitution is dissociative in nature, and this geometry for **8** represents the thermodynamically most stable product, with the weakly donating triflate anion occupying the position *trans* to the carbene carbon.

As we reported previously<sup>7</sup> (and also shown in Scheme 7), compound **8** can be reduced with two equivalents of  $\text{Cp}^*_2\text{Co}$  to



Fig. 6 Molecular structure of  $7\cdot(\text{OTf})_2\text{Cl}$ . Hydrogen atoms and the iso-propyl groups on phosphorus have been omitted for clarity. Displacement ellipsoids are shown at the 50% probability level. Selected bond lengths (Å): Ir(1)–P(1), 2.3797(17); Ir(1)–P(2), 2.3772(17); Ir(1)–C(1), 1.948(7); Ir(1)–Cl(1), 2.3035(17); Ir(1)–O(1), 2.197(5); Ir(1)–O(4), 2.131(5). Selected bond angles (°): C(1)–Ir(1)–Cl(1), 96.5(2); C(1)–Ir(1)–O(1), 177.7(3); C(1)–Ir(1)–O(4), 93.9(2); P(1)–Ir(1)–P(2), 164.76(6).



Fig. 7 Molecular structure of **8**. Hydrogen atoms, except for those on O(1), and the iso-propyl groups on phosphorus have been omitted for clarity. Displacement ellipsoids are shown at the 50% probability level. Selected bond lengths (Å): Ir(1)–P(1), 2.3642(13); Ir(1)–P(2), 2.3648(12); Ir(1)–C(1), 1.960(4); Ir(1)–O(1), 2.098(3); Ir(1)–O(2), 2.196(3); Ir(1)–Cl(1), 2.3200(12). Selected bond angles (°): C(1)–Ir(1)–O(1), 93.25(15); C(1)–Ir(1)–O(2), 175.46(16); O(1)–Ir(1)–O(2), 84.85(13); P(1)–Ir(1)–P(2), 171.34(4).







Elemental anal. calcd (%) for  $C_{38}H_{44}ClF_6IrO_6P_2S_4$  ( $7 \cdot (\text{OTf})_2\text{Cl}$  + toluene): C 40.44; H 3.93. Found: C 40.14; H 4.13.

### Low temperature addition of $H_2$ , $D_2$ or HD to $1 \cdot \text{Cl}$

All reactions involving the addition of 1 atm of  $H_2$ ,  $D_2$  or HD gas were performed by degassing the reaction mixture in a J-Young NMR tube, followed by addition of 1 atm of the gas at room temperature. Reactions carried out under 2–3 atm of a gas were performed by degassing the reaction mixture in a J-Young NMR tube and submerging the entire length of the tube in liquid nitrogen ( $-196^\circ\text{C}$ ) while open to the gas. Once sealed, the sample was warmed to the desired temperature ( $-78$  to  $-38^\circ\text{C}$ ).

### Low temperature addition of 1 : 1 $H_2/D_2$ to $1 \cdot \text{Cl}$

A 1 : 1 mixture of  $H_2$  :  $D_2$  was formed by first fully evacuating a double manifold vacuum line and then introducing  $H_2$  (300 mmHg) followed by  $D_2$  (300 mmHg). This mixture was allowed to equilibrate for a minimum of 6 hours after which a previously degassed sample in a J-Young NMR tube was opened to allow the mixture in. The same procedure was followed using 420 mmHg  $H_2$  for a 1.4 : 1 mixture of  $H_2/D_2$ .

### Measurement of equilibrium experiments

In order to dissolve a sufficient amount of  $H_2$  in solution without forming any of the thermodynamic isomer  $2 \cdot X_{trans}$ ,  $H_2$  was added to a solution of  $1 \cdot X$  (15 mM) in 0.5 mL  $CD_2Cl_2$  at  $-196^\circ\text{C}$  and warmed to  $-78^\circ\text{C}$  in a dry ice/acetone bath. The sample was then quickly transferred to the NMR spectrometer, which had previously been cooled to  $-78^\circ\text{C}$ . A  $^{31}\text{P}\{^1\text{H}\}$  experiment (243 MHz,  $d_1 = 2$  seconds, 16 scans) was then performed at each temperature and integrations were used to determine relative concentrations of the iridium complexes. To measure the concentration of  $H_2$  dissolved in solution during the thermodynamic van't Hoff equilibrium experiments, the reaction conditions were duplicated using an internal standard. Pentachlorobenzene (1.3 mg, 0.005 mmol, 10 mM) was dissolved in 0.5 mL  $CD_2Cl_2$  in a J-Young NMR tube which was subsequently degassed, charged with  $H_2$  gas and introduced into the NMR spectrometer at  $-78^\circ\text{C}$  as described above. A  $^1\text{H}$  NMR experiment (600 MHz,  $d_1 = 20$  seconds, 8 scans) was performed at each temperature and integrations were used to determine the concentration of  $H_2$  in solution; a correction of 1.33 was used to account for the  $\approx 25\%$  parahydrogen that is undetectable by  $^1\text{H}$  NMR spectroscopy.<sup>39,40</sup> As the concentration of  $D_2$  gas in solution could not be easily measured using NMR spectroscopy, it was estimated based on reported isotope fractionation factors.<sup>41</sup> Thus, the concentration of  $D_2$  in solution used for these calculations is  $1.1 \times [H_2]$  at a given temperature.

## Conclusions

The  $PC_{sp^2}P$  ligand systems exemplified by structures **I** and **II** are a unique class of PCP pincer ligands in that the anchoring  $C=M$  donor is capable of entering into cooperative reactivity with small molecule substrates. The reason for this partly stems from the generally reactive nature of metal carbenes, but here it

is also due to the adaptable nature of the character of the  $C=M$  unit in diaryl carbenes. This class of carbenes is somewhat special in that they lie somewhere between the extremes of Fischer type and Schrock type carbenes.<sup>16,42</sup> Although not strongly so, aryl substituents are  $\pi$ -donating and, in the case of ligands **I** and **II**, this results in an ability to function as both “Schrock-like” and “Fischer-like”, depending on the nature of the metal center.

This adaptability is clearly manifested in the comparative behavior of the iridium<sup>6</sup> and nickel<sup>9</sup> complexes of ligand **I**, but is also potentially a factor within the iridium manifold in the chemistry discussed herein. For example, the proposed transformation of **IV** into **VI** as shown in Scheme 6 is illustrative of this notion. Although **IV** can be clearly formulated as an Ir(III) derivative, transfer of the OH group to the iridium center is in some sense an oxidative addition reaction. But because the  $PC_{sp^2}P$  ligand that forms in this step can take on Fischer-like character, the iridium center is still formally Ir(III) or, put another way, the ligand enables the stabilization of highly reactive Ir(V) compounds ( $7 \cdot X_3$ ) or intermediates (**VI**). In so doing, the iridium complexes of ligand **II** provide a pathway for the overall hydrogenation of  $N_2O$  to water and dinitrogen.

The mechanistic details of this process have been probed by a combination of deuterium labeling experiments and low temperature NMR spectroscopic investigations. While developing this chemistry from an essentially stoichiometric process to a catalytic one would be desirable,<sup>43</sup> the reactions of the iridium-based products and catalytic intermediates with hydrogen have precluded us from achieving this goal as of yet. The results of these investigations, however, do provide the basis for further ligand modifications that are currently being pursued.

## Acknowledgements

Funding for this work was provided by NSERC of Canada in the form of a Discovery Grant and an Accelerator Supplement to W. E. P. W. E. P. also thanks the Canada Research Chair secretariat for a Tier I CRC (2013–2020). L. E. D. thanks NSERC for a CGSD scholarship and Alberta Innovates Technology Futures for a Graduate Student Scholarship.

## Notes and references

‡ The HD employed was 96 mol%, 98 atom% D, so small amounts of  $H_2$  are present; however more  $H_2$  is present in these samples than would be expected from the source gas.

§ The isotopologues are found in a nearly 1 : 1 ratio due to the excess of  $H_2$  in the  $H_2/D_2$  mixture used.

¶ While patterns reminiscent of those shown in Fig. 1 for the hydride ligands appear in the upfield region, they are not as well resolved as those found for the *trans* isomer. For  $2 \cdot Cl_{cis}$ , the  $^{31}\text{P}$  NMR spectrum is more suited for resolving the isotopologues.

1 *Topics in Organometallic Chemistry: Organometallic Pincer Chemistry*, ed. G. van Koten and D. Milstein, Springer, Heidelberg, 2013.



- 2 Pincer and Pincer-Type Complexes: Applications in Organic Synthesis and Catalysis, ed. Kalman J. Szabo and Ola F. Wendt, Wiley, Weinheim, Germany, 2014.
- 3 J. R. Khusnutdinova and D. Milstein, *Angew. Chem., Int. Ed.*, 2015, **54**, 12236–12273.
- 4 D. Gelman and R. Romm, in *Organometallic Pincer Chemistry*, ed. G. van Koten and D. Milstein, Springer, Berlin Heidelberg, 2013, vol. 40, ch. 10, pp. 289–317.
- 5 C. C. Roberts, D. M. Matías, M. J. Goldfogel and S. J. Meek, *J. Am. Chem. Soc.*, 2015, **137**, 6488–6491.
- 6 R. J. Burford, W. E. Piers and M. Parvez, *Organometallics*, 2012, **31**, 2949–2952.
- 7 L. E. Doyle, W. E. Piers and J. Borau-Garcia, *J. Am. Chem. Soc.*, 2015, **137**, 2187–2190.
- 8 H. D. Empsall, E. M. Hyde, R. Markham, W. S. McDonald, M. C. Norton, B. L. Shaw and B. Weeks, *J. Chem. Soc., Chem. Commun.*, 1977, 589–590.
- 9 D. V. Gutsulyak, W. E. Piers, J. Borau-Garcia and M. Parvez, *J. Am. Chem. Soc.*, 2013, **135**, 11776–11779.
- 10 P. Cui, C. C. Comanescu and V. M. Iluc, *Chem. Commun.*, 2015, **51**, 6206–6209.
- 11 C. C. Comanescu and V. M. Iluc, *Organometallics*, 2014, **33**, 6059–6064.
- 12 C. C. Comanescu, M. Vyushkova and V. M. Iluc, *Chem. Sci.*, 2015, **6**, 4570–4579.
- 13 J. Borau-Garcia, D. V. Gutsulyak, R. J. Burford and W. E. Piers, *Dalton Trans.*, 2015, **44**, 12082–12085.
- 14 R. J. Burford, W. E. Piers and M. Parvez, *Eur. J. Inorg. Chem.*, 2013, 3826–3830.
- 15 R. J. Burford, W. E. Piers, D. H. Ess and M. Parvez, *J. Am. Chem. Soc.*, 2014, **136**, 3256–3263.
- 16 M. T. Whited and R. H. Grubbs, *Acc. Chem. Res.*, 2009, **42**, 1607–1616.
- 17 M. Gozin, A. Weisman, Y. Bendavid and D. Milstein, *Nature*, 1993, **364**, 699–701.
- 18 J. D. Smith, J. Borau-Garcia, W. E. Piers and D. Spasyuk, *Can. J. Chem.*, 2015, 1–4, DOI: 10.1139/cjc-2015-0251.
- 19 P. P. Deutsch and R. Eisenberg, *Chem. Rev.*, 1988, **88**, 1147–1161.
- 20 T. Hascall, D. Rabinovich, V. J. Murphy, M. D. Beachy, R. A. Friesner and G. Parkin, *J. Am. Chem. Soc.*, 1999, **121**, 11402–11417.
- 21 G. Parkin, *Acc. Chem. Res.*, 2009, **42**, 315–325.
- 22 G. Parkin, *J. Labelled Compd. Radiopharm.*, 2007, **50**, 1088–1114.
- 23 L. Vaska, *Acc. Chem. Res.*, 1968, **1**, 335–344.
- 24 F. Abu-Hasanayn, K. Krogh-Jespersen and A. S. Goldman, *Inorg. Chem.*, 1993, **32**, 495–496.
- 25 F. Abu-Hasanayn, A. S. Goldman and K. Krogh-Jespersen, *Inorg. Chem.*, 1994, **33**, 5122–5130.
- 26 K. A. Woerpel and R. G. Bergman, *J. Am. Chem. Soc.*, 1993, **115**, 7888–7889.
- 27 C. A. Miller, T. S. Janik, C. H. Lake, L. M. Toomey, M. R. Churchill and J. D. Atwood, *Organometallics*, 1994, **13**, 5080–5087.
- 28 D. Sieh, M. Schlimm, L. Andernach, F. Angersbach, S. Nüchel, J. Schöffel, N. Šušnjar and P. Burger, *Eur. J. Inorg. Chem.*, 2012, **2012**, 444–462.
- 29 B. J. Truscott, D. J. Nelson, C. Lujan, A. M. Z. Slawin and S. P. Nolan, *Chem.–Eur. J.*, 2013, **19**, 7904–7916.
- 30 T. J. Hebden, K. I. Goldberg, D. M. Heinekey, X. Zhang, T. J. Emge, A. S. Goldman and K. Krogh-Jespersen, *Inorg. Chem.*, 2010, **49**, 1733–1742.
- 31 J. Zhou and J. F. Hartwig, *Angew. Chem., Int. Ed.*, 2008, **47**, 5783–5787.
- 32 V. M. Iluc, A. Fedorov and R. H. Grubbs, *Organometallics*, 2011, **31**, 39–41.
- 33 O. Blum and D. Milstein, *Angew. Chem., Int. Ed. Engl.*, 1995, **34**, 229–231.
- 34 D. Morales-Morales, D. W. Lee, Z. Wang and C. M. Jensen, *Organometallics*, 2001, **20**, 1144–1147.
- 35 M. Findlater, K. M. Schultz, W. H. Bernskoetter, A. Cartwright-Sykes, D. M. Heinekey and M. Brookhart, *Inorg. Chem.*, 2012, **51**, 4672–4678.
- 36 M. R. Kita and A. J. M. Miller, *J. Am. Chem. Soc.*, 2014, **136**, 14519–14529.
- 37 J. Choi, Y. Choliy, X. Zhang, T. J. Emge, K. Krogh-Jespersen and A. S. Goldman, *J. Am. Chem. Soc.*, 2009, **131**, 15627–15629.
- 38 S. Kundu, J. Choi, D. Y. Wang, Y. Choliy, T. J. Emge, K. Krogh-Jespersen and A. S. Goldman, *J. Am. Chem. Soc.*, 2013, **135**, 5127–5143.
- 39 Y. A. Ustynyuk and I. P. Gloriovov, *Russ. Chem. Bull.*, 2009, **58**, 1841–1846.
- 40 R. Eisenberg, *Acc. Chem. Res.*, 1991, **24**, 110–116.
- 41 J. Muccitelli and W.-Y. Wen, *J. Solution Chem.*, 1978, **7**, 257–267.
- 42 T. M. Trnka and R. H. Grubbs, *Acc. Chem. Res.*, 2000, **34**, 18–29.
- 43 W. B. Tolman, *Angew. Chem., Int. Ed.*, 2010, **49**, 1018–1024.

