H/D exchange under mild conditions in arenes and unactivated alkanes with $C_6D_6$ and $D_2O$ using rigid, electron-rich iridium PCP pincer complexes†

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The synthesis and characterization of an iridium polyhydride complex (Ir-H$_4$) supported by an electron-rich PCP framework is described. This complex readily loses molecular hydrogen allowing for rapid room temperature hydrogen isotope exchange (HIE) at the hydridic positions and the $\ Greek c -$H site of the ligand with deuterated solvents such as benzene-$d_6$, toluene-$d_6$ and THF-$d_8$. The removal of 1–2 equivalents of molecular H$_2$ forms unsaturated iridium carbene trihydride (Ir-H$_3$) or monohydride (Ir-H) compounds that are able to create further unsaturation by reversibly transferring a hydride to the ligand carbene carbon. These species are highly active hydrogen isotope exchange (HIE) catalysts using C$_6$D$_6$ or D$_2$O as deuterium sources for the deuteration of a variety of substrates. By modifying conditions to influence the Ir-H$_3$ speciation, deuteration levels can range from near exhaustive to selective only for sterically accessible sites. Preparative level deuterations of select substrates were performed allowing for procurement of $>95\%$ deuterated compounds in excellent isolated yields; the catalyst can be regenerated by treatment of residues with H$_2$ and is still active for further reactions.

Introduction

Metal-catalyzed hydrogen isotope exchange (HIE) is a key fundamental process for preparing isotopically labeled organic compounds necessary for studying reaction mechanisms and biological metabolic processes.1 Since the seminal work on homogeneous HIE by Garnett and Hodges2,3 as well as Shilov and co-workers,4,5 advances in analytical biology and chemistry have created an ever growing demand for deuterium4 and tritium7 labeled molecules.6 Consequently, an important body of research devoted to studying the homogeneously catalyzed exchange of hydrogen and deuterium atoms (H/D exchange) has developed over the last 25 years.7–12 Pioneering work by the Bergman group on pentamethylcyclopentadienyl supported iridium hydrides demonstrated low temperature C–H activation13–15 and it was subsequently shown that H/D exchange by cationic iridium complexes16–18 occurred under relatively mild conditions using C$_6$D$_6$ as a deuterium source. Most of these transformations involved selective deuterium incorporation into arene substrates10,11 with only a handful of catalysts capable of H/D exchange on unactivated alkane substrates under more forcing conditions.16,17,19–24

Tridentate “pincer” ligands have played a prominent role in homogeneous catalysis.25,26 For example, polyhydric pincer complexes of the late transition metals (particularly Ru and Ir) are known to be exemplary alkane dehydrogenation catalysts27 and CO$_2$ reduction catalysts,28,29 but recent work has demonstrated their promise as competent catalysts for promoting homogenous H/ D exchange. Leitner, Milstein and co-workers were the first to show that a Ru based PNP polyhydride, I Chart 1, could use C$_6$D$_6$ or D$_2$O for arene deuteration at 50 °C.30 Shortly after, Zhou and Hartwig revealed that the aliphatic iridium PCP polyhydride II could be used as a pre-catalyst to efficiently perform H/D exchange between olefins and C$_6$D$_6$ at room temperature without isomerization.31 Following these reports, PNP (III) and PCP (IV) complexes synthesized by Grubbs32 and Wendt,33 respectively, were also tested as HIE mediators. The iridium dihydride species III was able to selectively deuterate aromatic compounds and silanes, while the cyclohexyl-based PCP complex IV was found to successfully deuterate arenes and to a limited extent C$_{sp^3}$H sites, albeit at elevated temperatures of 150 °C.

Design elements of the ligands II/IV and III have been incorporated into a family of PCP ligands V introduced by our group34 and also explored by others.35–37 An attractive feature of these ligands is their ability to assume P$_{Csp^3}$P or P$_{C\text{carbene}}$P forms by the addition or removal of a hydrogen atom from the...
anchoring carbon of the pincer framework. This has precedence in the ligands II and IV but in the aryl-linked PCP systems V and VI (and the related benzothiophene-based system VI40) this is potentially more controllable because of the lack of β-hydrogens in the PCP linker framework, and through manipulation of the electronic properties of the ligand via substitution on the aryl linkers and/or rigidification of the ligand via additional linking of the backbone aryl groups.41,42 With these two structural levers, a library of PCP ligands has been prepared and a substantial range of rCO values has been found for [(PCcarbeneP)IrCO]+ species.43 We have shown that the rate of addition of N2O to the C=Ir bond can therefore be tuned substantially.44 Furthermore, the rigidification of the ligand tends to favor the PCcarbeneP form of the ligand vs. the PCsp3P form, while also providing greater protection against C–C bond cleavage reactions that destroy the integrity of the ligand in unlinked systems akin to V.45 The ability of the carbon atom in these PCP pincers to engage in ligand cooperativity has been exploited in hydrogen,35,46 silane47–49 and other small molecule50–53 activations in various systems. Furthermore, in polyhydridic systems, the dibenzyl hydrogen is actively involved in exchange with the metal hydrides,34,35 increasing the “hydrogen capacity” of these compounds relative to other (for example PNP) polyhydridic pincer complexes and potentially providing new pathways for catalysis.

In both the parent, unsubstituted, unlinked system V, and that incorporating the benzothiophene-based ligand VI, the Ir polyhydride is only stable under an atmosphere of dihydrogen but it was observed that slow H/D exchange in the sterically exposed C–H bonds of the ligand aryl backbone occurred in C6D6. We hypothesized that a more electron rich system would accelerate this process and indeed, the linked dihydroanthracene PCP ligand framework fitted with electron donating dimethylamino groups para to the PCP carbon (bottom of Chart 1) supports a stable, isolable polyhydride species Ir-H4 (Chart 1). While stable as a solid, this complex readily loses molecular hydrogen in solution to give species that are highly active HIE catalysts at low loadings and under mild conditions. Both C6D6 and D2O can be used as deuterium sources and selectivity can be modulated to some extent by controlling the catalyst speciation.

Results and discussion

Synthesis and characterization of iridium PCP pincer polyhydride Ir-H4

Using methodology established previously,46,54 the iridium PCP pincer complex Ir-OH was prepared by heating the previously...
reported Ir-Cl\textsuperscript{44} to 120 °C in toluene with an excess of CsOH for 24 hours (Scheme 1, red arrows). The resulting brown solution was filtered through Celite® and a brown powder was isolated in 80% yield after removing all volatiles in vacuo. The \textit{Ir-OH} complex displays a characteristic –OH triplet (\(J_{\text{HP}} = 3.8\) Hz) at 3.62 ppm in the \(^1\text{H}\) NMR spectrum as well as a singlet in the \(^{31}\text{P}\) \(^{1}\text{H}\) NMR at 51.9 ppm. The presence of an iridium-carbene moiety was indicated by a triplet signal at 196.0 ppm (\(J_{\text{CP}} = 2.2\) Hz) in the \(^{13}\text{C}\)\(^{1}\text{H}\) NMR spectrum.

Dissolving \textit{Ir-OH} in benzene and exposing it to 4 atm of \(\text{H}_2\) gas for 3 hours resulted in a color change from dark reddish-brown to yellow-orange. The \(^{31}\text{P}\) \(^{1}\text{H}\) NMR spectrum for the solution exhibited a single peak at 56.5 ppm and upon solvent removal a brown residue was formed. This residue was then stirred in \textit{n}-pentane under 1 atm of \(\text{H}_2\) for 20 minutes producing a beige powder, \textit{Ir-H}_4, that was isolated in 98% yield following the removal of all volatiles in vacuo. Alternatively, we found stirring the precursor \textit{Ir-Cl} with NaOH (10 eq.) under \(\text{H}_2\) gas (4 atm) in benzene for 3 hours produced a similar red-orange solution (Scheme 1, blue arrow). Following filtration through a pad of Celite®, the solution was stirred for 20 minutes under \(\text{H}_2\) gas. All volatiles were removed in vacuo and a beige powder was again isolated in 94% yield. Compared to the parent \textit{ortho}-phenylenediyne \textit{V}\textsuperscript{20} (Chart 1) and the benzophenoxobased ligand congener \textit{VI}\textsuperscript{46} which both could not be isolated due to decomposition via loss of \(\text{H}_2\) under vacuum, complex \textit{Ir-H}_4 is remarkably stable as a solid and can be stored indefinitely at room temperature under an inert atmosphere. Furthermore, it can be interrogated via NMR spectroscopy in cyclohexane-\textit{d}_{12} or methylyclohexane-\textit{d}_{14} under argon, whereas the others (\textit{V} and \textit{VI}) must be kept under an atmosphere of \(\text{H}_2\) in order to avoid decomposition. The \(^{1}\text{H}\) NMR spectrum for \textit{Ir-H}_4 in cyclohexane-\textit{d}_{12} displays a single triplet (average \(J_{\text{HP}} = 10.2\) Hz) resonance at \(-10.32\) ppm and a singlet at 5.52 ppm, integration indicating the presence four rapidly exchanging hydridic nuclei and one \(\alpha\)-C–H–Ir proton, respectively. Moreover, the ligand resonances in the spectrum clearly point to \(C_5\) symmetry with 12 equivalent dimethyl amino protons represented by a singlet at 2.85 ppm and two signals at 1.23 ppm and 1.84 ppm for the inequivalent –CH\textsubscript{3} groups attached to the carbon backbone linker. The \(^{13}\text{C}\) \(^{1}\text{H}\) NMR also corroborates the formation of \textit{Ir-H}_4 showing the loss of the carbene triplet associated with the \textit{Ir-Cl} and \textit{Ir-OH} complexes and the presence of a signal at 21.0 ppm coupling to the \(\alpha\)-C–H proton at 5.52 ppm in the \(^{3}\text{H}\)–\(^{13}\text{C}\) HSQC spectrum.

Crystals of the compound were obtained by slowly cooling a supersaturated solution of \textit{Ir-H}_4 in isopropanol from 50 °C to 20 °C and crystallographic analysis corroborates the solution data described above. The hydrogen atoms associated with the metal center were located from the Fourier map but low accuracy in \textit{Ir-H} and \(\text{H–H}\) distances preclude conclusions concerning their arrangement as hydride or dihydrogen ligands. The data show the anchoring \(\alpha\)-carbon (C1 in Fig. 1) is pyrimalized (the sum of the angles about C1 is 335.2(8)°), and the central 6 membered ring containing this \(\alpha\)-carbon has also adopted a “boat”–like conformation. The sp\textsuperscript{3} hybridization of the \(\alpha\)-carbon is also indicated by the central Ir1–C1 bond length of 2.184(5) Å which is similar to other sp\textsuperscript{3} hybridized ir–C bonds we have observed with these \(\text{NR}_2\) substituted ligand frameworks\textsuperscript{45} and lengthened significantly from the values of \(1.93\) Å seen in \textit{PC}_{\text{carbene}}P complexes of this ligand.\textsuperscript{44}

Given the lack of crystallographic data concerning the iridium hydrides in \textit{Ir-H}_4, we turned to criteria developed by Halpern and co-workers\textsuperscript{35} to shed some light on what form dominates the speciation of \textit{Ir-H}_4 in solution. While it is typically difficult to unambiguously categorize the hydride bonding mode for these complexes, we were able to rule out the possibility that \textit{Ir-H}_4 is a “non-classical” Ir(r) bis-dihydrogen species (Scheme 2). The \(T_1\) (min) relaxation for the hydridic nuclei of \textit{Ir-H}_4 in methylyclohexane-\textit{d}_{14} (Fig. S1†) was found to be 186 ms at 600 MHz (\(10^\circ\text{C} \pm 0.5\) °C), approximately 155 ms when adjusting to a 500 MHz field strength. These values are comparable to those obtained for other PCP pincer polyhydride complexes reported by Heinekey,\textsuperscript{56} Wendt\textsuperscript{44} and us;\textsuperscript{35} a collection of comparative data is given in Table S1.† These data point to an equilibrating mixture of a “classical” Ir(n) tetrahydride and the “intermediate” Ir(m) dihydrogen–dihydride isomers shown

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**Scheme 1** Synthetic routes to iridium PCP polyhydride \textit{Ir-H}_4.
methylcyclohexane-d_{14}. All four iridium hydrides and the benzyllic position on the ligand are rapidly deuterated under these conditions, but by halting before full deuteration, the residual hydride resonance in methylcyclohexane-d_{14} at −10.25 ppm displays a resolved seven line pattern with an average observed J_{H-D} of 3.1 ± 0.1 Hz when collected at 50 °C (Fig. S2†). This coupling correlates to an approximated average H–H atomic distance of 1.02–1.25 Å (Table S1†)36 and is consistent with equilibrating classical tetrahydride (H–H > 1.2 Å) and intermediate dihydrogen (H–H ≈ 1.0–1.2 Å) dihydride structures (Table S1†)37−39 as depicted in Scheme 2. Unlike the related system based on IV, attempts to freeze out the exchange between the isomers of the latter and/or the equilibrium between the two via variable temperature ^1H[^31P] NMR spectroscopy on Ir-H₄ in methylcyclohexane-d_{14} showed no decoalescence of the hydride resonance to temperatures as low as −110 °C (Fig. S3†). This indicates a lower energetic barrier for hydride exchange than the estimated 8.7 kcal mol⁻¹ at −70 °C reported by Wendt et al. in the system IV.34 Indeed, M06 DFT calculations indicate the potential energy surface for postulated tetrahydride and dihydrogen–dihydride complexes is relatively flat with an energy range of <2 kcal mol⁻¹ between these four structures (Scheme 2). For the full covalent Ir-H bonds, the distances are ∼1.6 Å. In the dihydrogen–dihydride structures, the dihydrogen distances are ∼0.85 Å, which are only slightly stretched compared to free H₂.29 The barriers are <12 kcal mol⁻¹ for conversion between tetrahydride and dihydrogen–dihydride complexes. The transition-state distances for the forming H–H bonds are ∼1.1 Å. Overall, these DFT thermodynamic and kinetic estimates suggest that all intermediate and classical structures are accessible in solution and the spectroscopically observed complex lies within the dihydrogen–dihydride tetrahydride continuum. While the benzyl hydrogen on the Csp³ carbon of the ligand certainly exchanges with the four iridium hydrgens rapidly on the chemical timescale, the two peaks only undergo slight broadening at temperatures below which loss of H₂ and conversion to other species ensues (vide infra). Line shape analysis to probe the barrier was not possible with the spectral data available in this system, unlike that of IV.34

in Scheme 2. The experimental T₁ of 155 ms is somewhat lower than the calculated T₁ value of 203 ms calculated from the parameters obtained from the M06 density functional theory (DFT) optimized structure of Ir-H₄, suggesting a significant contribution from any or all of the dihydrogen–dihydride isomers, which also can be optimized using DFT and give rise to T₁ values of ∼15 ms using the Halpern methodologies. Significantly, the non-classical bis-dihydride structure could not be optimized by DFT, suggesting that this formally Ir(0) isomer is not favored with this electron rich pincer ligand and does not contribute to the speciation of Ir-H₄ in solution.

Another key metric in this family of compounds is the averaged J_{H-D} coupling constant in the Ir-HD₃ isotopologue. For Ir-H₄, this species is trivially prepared by dissolving Ir-H₄ in C₆D₆ and stirring for 15 minutes at room temperature (vide infra), removing the solvent and redissolving the sample in methylycyclohexane-d_{14}. All four iridium hydrides and the benzyllic position on the ligand are rapidly deuterated under these conditions, but by halting before full deuteration, the residual hydride resonance in methylcyclohexane-d_{14} at −10.25 ppm displays a resolved seven line pattern with an average observed J_{H-D} of 3.1 ± 0.1 Hz when collected at 50 °C (Fig. S2†). This coupling correlates to an approximated average H–H atomic distance of 1.02–1.25 Å (Table S1†)36 and is consistent with equilibrating classical tetrahydride (H–H > 1.2 Å) and intermediate dihydrogen (H–H ≈ 1.0–1.2 Å) dihydride structures (Table S1†)37−39 as depicted in Scheme 2. Unlike the related system based on IV, attempts to freeze out the exchange between the isomers of the latter and/or the equilibrium between the two via variable temperature ^1H[^31P] NMR spectroscopy on Ir-H₄ in methylcyclohexane-d_{14} showed no decoalescence of the hydride resonance to temperatures as low as −110 °C (Fig. S3†). This indicates a lower energetic barrier for hydride exchange than the estimated 8.7 kcal mol⁻¹ at −70 °C reported by Wendt et al. in the system IV.34 Indeed, M06 DFT calculations indicate the potential energy surface for postulated tetrahydride and dihydrogen–dihydride complexes is relatively flat with an energy range of <2 kcal mol⁻¹ between these four structures (Scheme 2). For the full covalent Ir-H bonds, the distances are ∼1.6 Å. In the dihydrogen–dihydride structures, the dihydrogen distances are ∼0.85 Å, which are only slightly stretched compared to free H₂.29 The barriers are <12 kcal mol⁻¹ for conversion between tetrahydride and dihydrogen–dihydride complexes. The transition-state distances for the forming H–H bonds are ∼1.1 Å. Overall, these DFT thermodynamic and kinetic estimates suggest that all intermediate and classical structures are accessible in solution and the spectroscopically observed complex lies within the dihydrogen–dihydride tetrahydride continuum. While the benzyl hydrogen on the Csp³ carbon of the ligand certainly exchanges with the four iridium hydrgens rapidly on the chemical timescale, the two peaks only undergo slight broadening at temperatures below which loss of H₂ and conversion to other species ensues (vide infra). Line shape analysis to probe the barrier was not possible with the spectral data available in this system, unlike that of IV.34

Solution behavior of Ir-H₄

While Ir-H₄ is quite stable in the solid state, evidence exists for reversible loss of H₂ in solution (Scheme 3). When dilute solutions in benzene were exposed to vacuum, the pale-yellow color darkened slowly to a maroon-tinged hue. Readmission of H₂ gas to the vessel gave back the yellow solution; monitoring this transformation by NMR spectroscopy indicated that the maroon solution was essentially spectroscopically identical to the yellow solutions of Ir-H₄. However, when heating solutions of Ir-H₄ to 80 °C under a static vacuum and monitoring by NMR spectroscopy, other resonances became apparent. In the ^1H NMR spectrum, a new resonance at 2.71 ppm in the NMMe₃ region and a faint peak at −10.30 ppm were observed in about a 12 : 3 ratio. In the ^31P[^1H] NMR spectrum, resonance at 66.4 ppm grew in relative to that of Ir-H₄ at 56.5 ppm (Scheme 3 and Fig. S4 and S5†). Cooling the sample resulted in loss of this

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**Scheme 2** (a) Outline of isomerism in Ir-H₄ with benzene-solvated enthalpies (Gibbs energies) in kcal mol⁻¹ (b) Ir-H and H–H distances (Å) found in M06 DFT structures.
resonance and a return to solutions that were >98% Ir-H4. We hypothesized that this maroon species was either the PCsp3P supported Ir(III) dihydride or the PCarbenep ligated Ir(III) trihydride shown in Scheme 3. These should be distinguishable by the 13C resonance of the PCP ligand, but because it is produced in such small amounts via this method, we sought ways to prepare or at least enrich the speciation of this compound in order to obtain more concentrated samples.

 Heating Ir-H4 in benzene under dynamic vacuum did lead to a much darker colored solution, but the 31P{1H} NMR spectra indicated that a third species with a resonance at 53.6 ppm was dominant in these samples. Separate synthesis, NMR spectroscopy and X-ray crystallography confirmed this product to be the PCarbenep ligated Ir(III) trihydride shown in Scheme 3. In attempts to generate Ir-H3/Ir-H4 by treating Ir-H3 with the dehydrogenation agent tert-butyl ethylene, varying mixtures of the three components were obtained depending on the equivalency of olefin employed; when >3 equivalents were used, Ir-Ph was cleanly produced and isolated as a green solid in 72% yield. The 13C{1H} NMR spectrum of Ir-Ph in benzene-d6 shows a distinct carbene triplet at 221.1 ppm (JCP = 4.2 Hz) while the 1H NMR spectrum shows three separate aryl proton signals for the new phenyl group as well as characteristic ligand resonances. Crystallographic data (Fig. 2) show a central Ir1–C1 distance of 1.965(4) Å. This is shorter than that found in Ir-H4 and indicative of an iridium carbene bond similar to analogous structures. Accordingly, unlike the boat-like shape found in the Ir-H4 the central 6-membered ring of the ligand framework adopts a planar conformation, as a result of the sp3 hybridization found at the carbene center. The rigidity of the ligand employed here stabilizes this phenyl complex with respect to the intriguing C–C bond formation reaction observed by Wendt in a the more flexible IV.40 Taken together, the above observations suggest that Ir-H4 rather easily loses one equivalent of H2, even thermally, to give Ir-H3/Ir-H4, but when exposed to vacuum and heat, a second equivalent of H2 is eliminated to form the (unobserved) compounds Ir /[Ir-H] depicted in Scheme 3. Consistent with the relative ease for H2 loss, the DFT calculated energy of Ir-H4 is only endergonic by ~5 kcal mol−1 relative to Ir-H4. Calculations suggest that formation of the dihydrogen–dihydride complex and dihydrogen dissociation is required before 1,2-hydrogen shift from the ligand to Ir to form Ir-H3, which only requires a barrier of ~15 kcal mol−1. The Ir-H/[Ir] structures are calculated to be 6.6 and 12.2 endergonic relative to Ir-H4. Overall, the thermodynamic and kinetic DFT calculations suggest that Ir-H3, Ir-H3, Ir-H, and Ir-[Ir] structures are all viable. These compounds rapidly C–H activate benzene and, after elimination of another equivalent of H2, Ir-Ph is produced; this compound rapidly converts back to Ir-H4 upon treatment with H2, with loss of C9H8. The speciation of the Ir-H3/Ir-H4 manifold was solidified as being mainly Ir-H3 by mixing isolated Ir-H4 and Ir-Ph in

Fig. 2 Synthesis and thermal ellipsoid diagram of Ir-Ph (ellipsoids shown in 50% probability; hydrogen atoms have been omitted for clarity). Selected metrical data for Ir-Ph: bond distances (Å): Ir1–C1, 1.965 (4); Ir1–C33, 2.118(4); Ir1–P1, 2.2581(13); Ir1–P2, 2.2671(17); C1–C2, 1.467(6); C1–C14 1.469(6); bond angles (°): P1–Ir1–P2, 166.46(4); C1–Ir1–C33, 179.53(19); P1–Ir1–C1, 83.46(13); P1–Ir1–C33, 96.38(13); Ir1–C1–C2, 122.6(3); Ir1–C1–C14, 122.9(4); C2–C1–C14, 114.5(4).
a $\approx 2 : 3$ ratio in benzene; this experiment resulted in an equilibrium mixture of these two compounds along with about 40% \textbf{Ir-H$_3$} which allowed us to acquire meaningful $^1$H and $^{13}$C NMR data on the latter. Both are consistent with $C_{2v}$ symmetry and the presence of a PC$_{\text{carbene}}$P ligand. In the $^1$H NMR spectrum, no resonance in the region between 3 and 6 ppm consistent with the presence of an $\alpha$-C(H)-Ir proton was observed, while a characteristic resonance at 245.0 ppm in the $^{13}$C($^1$H) NMR spectrum is diagnostic of a carbene carbon. This signal correlated with the triplet integrating to 3 hydrogens at $-10.12$ ppm in the 2D $^2$H$_2$-$^{13}$C HMBC spectrum. At room temperature, the single resonance for the three hydrides in \textbf{Ir-H$_3$} indicate they are in rapid exchange, presumably via a 1,2-hydride shift from the iridium to the carbene carbon, i.e., via \textbf{Ir-H$_2$}. The apparent thermodynamic preference for \textbf{Ir-H$_3$} over \textbf{Ir-H$_2$} in this system contrasts with observations reported by Wendt and co-workers,\textsuperscript{14,15} but these structures are very close in energy and in rapid exchange. DFT calculations using a variety of functionals (M06, wb97X-D, M11, and others) all indicate that the \textbf{Ir-H$_2$} structure is slightly lower in energy than \textbf{Ir-H$_3$}; the methodology has limitations for distinguishing these types of isomers accurately.

**HIE with iridium polyhydrides \textbf{Ir-H$_n$}**

As briefly alluded to above, dissolution of \textbf{Ir-H$_4$} into C$_6$D$_6$ (and indeed toluene-$d_8$ or THF-$d_8$) leads to the rapid deuteration of the $\alpha$-C(H)-Ir and \textbf{Ir-H} positions at 20 $^\circ$C to form \textbf{Ir-D$_4$} (Scheme 4a) and so it had to be characterized in deuterated cyclohexane or methylycyclohexane solvents which do not surrender deuterium at room temperature. Heating \textbf{Ir-H$_4$} in C$_6$D$_6$ to 65 $^\circ$C for 24 hours lead to trace deuteration of the ligand NME$_2$ and iso-propyl methyl groups (Fig. S6†). Given the coordinative saturation of \textbf{Ir-H$_4$}, we hypothesized that hydrogen loss to form \textbf{Ir-H$_3$} was required to initiate the HIE process. Indeed, mixing \textbf{Ir-H$_4$} and \textbf{Ir-Ph} as described above to generate solutions enriched in \textbf{Ir-H$_3$} leads to much higher deuteration activity. Heating solutions of \textbf{Ir-H$_4$/Ir-H$_3$/Ir-Ph} (3 : 2 : 2) in C$_6$D$_6$ to 65 $^\circ$C for 24 hours gave the three species essentially exhaustively deuterated in the positions shown in Scheme 4b (Fig. S7†). Separate experiments showed that heating pure \textbf{Ir-Ph} under these conditions gave only 70% conversion to \textbf{Ir-Ph$_{\text{dir}}$} after 24 hours and no deuteration of the ligand NME$_2$ or iso-propyl groups was observed (Fig. S8†), suggesting that \textbf{Ir-Ph} is not an effective HIE promoter in this system.

While the $^1$Pr groups are likely deuterated via an intramolecular C-H activation, deuteration of the more remote NME$_2$ groups suggest that intermolecular catalytic deuteration of other substrates under mild conditions might be possible. We chose \textit{meta}-xylene as a test substrate to evaluate the various iridium species as HIE catalysts due to the sterically varied aryl C-H bonds present, as well as the benzylic sp$^3$ hybridized C-H bonds. The results of these experiments are summarized in Fig. 3. Solutions containing \textit{meta}-xylene and 1 mol% of either \textbf{Ir-H$_4$} (argon, 1 atm), \textbf{Ir-H$_4$} (static vacuum), \textbf{Ir-Ph} (argon, 1 atm) or \textbf{Ir-Ph} activated with D$_2$ gas ($\approx 1$ equivalent) were heated to 50 $^\circ$C in benzene-$d_8$ and monitored by $^1$H NMR spectroscopy. Each panel in Fig. 3 indicates the rate of loss of substrate protons as time progresses for a different C-H position in the substrate, as a function of the catalyst precursor employed (colored shapes). Cyclooctane was employed as an internal standard. As can be seen in panel a, none of the catalyst formulations deuterate the sterically encumbered 2 position of \textit{meta}-xylene at 50 $^\circ$C; the other positions, however, are deuterated to varying degrees. In general, \textbf{Ir-H$_4$} under argon and \textbf{Ir-Ph} behave similarly for deuteration of the C-H bonds in the benzylic, and aryl 4/5 positions, with moderate activity observed only for the sterically accessible aryl C-H bond in the 5 position. The catalyst systems which would be expected to generate \textbf{Ir$/[\text{Ir-H$_4$}]}$ (Scheme 3) or \textbf{Ir-H$_3$/Ir-H$_4$}, namely \textbf{Ir-Ph} plus D$_2$ (blue triangles) or \textbf{Ir-H$_4$} under vacuum (green inverted triangles) fared much better, with the former able to almost completely deuterate \textit{meta}-xylene at all three positions under these mild conditions. Notably, when \textbf{Ir-H$_4$} is simply exposed to vacuum (green triangles), there is an acceleration in the rate of deuteration consistent with the slow formation of \textbf{Ir-H$_3$} through loss of H$_2$ to the headspace, whereas when the active species is generated through activation of \textbf{Ir-Ph} with D$_2$, a more typical first order time evolution is observed, further implicating \textbf{Ir-H$_3$} as the active catalyst or catalyst precursor.

Since it was clear from the above experiments that \textbf{Ir-Ph} activated with D$_2$ gave the most active HIE catalyst, further optimization of conditions for this reaction were undertaken with this as a starting point (Table 1). Entries 1 and 2 reaffirm
that a catalyst is required, and that \textbf{Ir-Ph} itself is a poor catalyst with low activity. Entries 3–6 show that there is an optimal amount of added D$_2$ required to activate the catalyst of 1–3 equivalents. With excess D$_2$ (entry 7), activity drops off markedly, presumably because the iridium speciation is now essentially 100\% Ir-D$_4$, which (as shown in Fig. 3) is not active under these conditions. Raising the temperature 15 degrees to 65 °C (entries 8–12) leads to greater HIE activity; comparison of entry 4 with entry 10 underscores this observation. Even the difficult 2-position sees some deuteration at this stage. As entries 11 and 12 show, near exhaustive deuteration of the benzylic, 4/6 and 5 positions can be achieved at 65 °C, with 1 mol\% of \textbf{Ir-Ph} activated with 1.4 equivalents of D$_2$; entry 12 uses double the amount of C$_6$D$_6$ in order to drive the deuteration in these positions to near completion.

In comparison to other PCP (and PNP) pincer catalysts$^{31-34}$ for HIE reactions (Chart 1), the iridium systems described here appear to be more active under milder conditions. Furthermore, through manipulation of the speciation of the active \textbf{Ir-H$_n$} compounds, the selectivity of deuteration can also be to some extent controlled. Finally, given the stability of these systems to water, D$_2$O can also be used as a deuterium source rather than

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**Table 1** Optimization of conditions for exhaustive deuteration of \textit{meta}-xylene with C$_6$D$_6$\textsuperscript{a}

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<th>Entry</th>
<th>Temp. (°C)</th>
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<th>4/6</th>
<th>5</th>
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$^a$ Optimization performed with \textit{meta}-xylene (0.38 mmol) and \textbf{Ir-Ph} (1 mol\%) in C$_6$D$_6$ (0.5 mL). D$_2$ gas was injected into 0.015 M solutions of \textbf{Ir-Ph} in C$_6$D$_6$ and stirred for 10 minutes prior to its addition to \textit{m}-xylene in C$_6$D$_6$ (0.25 mL). Conversion was measured via $^1$H NMR using cyclohexane as an internal standard. $^b$ Volume of C$_6$D$_6$ increased to 1.0 mL.
C₆D₆. The results of HIE with a number of substrates, under three sets of conditions are summarized in Fig. 4. Using the optimized conditions of entry 11 in Table 1 developed for meta-xylene, HIE for several substrates were examined (Fig. 4, black data). Deuterium incorporation was found to be quite high for several arene based substrates with unhindered Csp²–H positions in fluorobenzene, naphthalene, aniline and 2,6-lutidine being deuterated almost quantitatively. In the case of naphthalene even the less reactive bay positions were fully deuterated in contrast to III and IV, which required longer reaction times and higher temperatures to achieve even partial deuteration of these positions in naphthalene. More interestingly, typically less active Csp³–H positions were significantly deuterated in substrates such as THF, triethylamine, 2,6-lutidine and meta-xylene under very mild conditions with no sign of catalyst decomposition. In the case of naphthalene even the less reactive bay positions were fully deuterated in contrast to III and IV, which required longer reaction times and higher temperatures to achieve even partial deuteration of these positions in naphthalene. More interestingly, typically less active Csp³–H positions were significantly deuterated in substrates such as THF, triethylamine, 2,6-lutidine and meta-xylene under very mild conditions with no sign of catalyst decomposition. This level of deuteration for the Csp³–H sites, along with precedents from previous systems, prompted further investigation into the deuteration of aliphatic hydrocarbons. Despite no detectable H/D exchange occurring between benzene-d₆ and cyclooctane at 50 °C the increased temperature of 65 °C resulted in mild deuteration of 9% after 24 hours which increased to 36% after 5 days. For n-heptane, deuteration of the terminal methyl groups was found to be 58% while internal methylene positions were deuterated to 37% after 5 days. The $^{13}$C{¹H} NMR revealed that the majority of deuteration performed on the interior methylene groups occurred the 2 and 6 positions. This is intriguing considering reactivity of cyclopentadienyl Ru complex reported by Nikonov was limited to the terminal methyl C–H sites. Reduced activity was also observed with phenol while catalysis was completely halted in the presence of ethyl benzoate and pyridine. The $^{31}$P{¹H} NMR spectra of these latter reactions showed that complexes Ir-Ph-d₅, Ir-D₃ and Ir-D₄ were no longer present in solution. While exhaustive deuteration is desirable in some situations, mild, selective deuteration has some interest for drug discovery and evaluation. The data in Table 1 indicates that, under conditions where Ir-D₄ dominates, deuteration is emphasized at the more sterically accessible sites (see entry 9). We therefore examined HIE with the substrates in Fig. 4 under these conditions (blue data). Using these conditions, good selectivity is possible with compounds containing directing groups and less accessible Csp³–H sites. While fluorobenzene still showed almost quantitative deuterium incorporation, naphthalene, 2,6-lutidine and N,N-dimethylbenzamide showed higher levels of deuteration at accessible sites relative to less accessible ones. Catalytic HIE of aniline had a slightly greater preference for...
and ortho-sites. The increased propensity of the iridium catalyst to bind at the more strongly donating ortho site likely resulted in this mild enhancement. While selectivity was observed in phenol, results were almost identical to the exhaustive conditions with the -OH and ortho-positions being deuterated to 42% and 37%, respectively. More surprisingly, deuteriation for ethyl benzoate and pyridine became possible. Ethyl benzoate was significantly deuterated at both the meta- and para-positions while pyridine showed low level deuteriation at all sites. For both pyridine and ethyl benzoate, the greater amount of Ir-D4 may stave off immediate catalyst decomposition by providing increased levels of molecular D2 present in solution which help prevent these substrates from strongly binding to the iridium center. Under these conditions deuteration at Csp3 ortho-positions were obtained. The experiments were repeated with 1 mol% of Ir-H4 dissolved in neat THF (8.3 mmol) under static vacuum and the red data in Fig. 4 shows the results HIE using D2O with Ir-H4 under static vacuum as the catalyst precursor. To solubilize the iridium species, reactions were performed in a biphasic mixture of D2O and cyclohexane. For meta-xylene, H/D exchange occurred in trace amounts at the benzyl position, 20% at the 4 and 6 positions and quantitatively at the 5 position. Deuteration of the other substrates in Fig. 4 using D2O gave broadly similar results to those obtained under the “selective” conditions giving rise to the blue data, with a few notable differences. Fluorobenzene was again exhaustedly deuterated while naphthalene and 2,6-lutidine had deuterium unevenly distributed throughout the aren systems. Aniline was deuterated almost quantitatively in all positions while deuteriation in phenol was greatly enhanced to 95%, although limited to the ortho and -OH sites. Deuterium levels in pyridine were found to be slightly higher than in the previous attempts but again catalysis was hampered by catalyst degradation (detected via 31P{1H} NMR). Aliphatic hydrocarbons were found to be mostly unreactive though trace H/D exchange had occurred at the terminal methyl positions of n-heptane. Unlike the results produced using Ir-Ph and 9 mol% D2 with benzene-d6, H/D exchange was found to be possible in both triethylamine and THF, albeit in low levels.

On a preparative scale, these biphasic conditions can be used with neat substrates as summarized in Fig. 5. Both benzene and fluorobenzene (8.3 mmol each) were deuterated at all positions to 90% and 94%, respectively, with D2O (305 mmol) at 80 °C using 1 mol% of Ir-H4. Following extraction of the organic layers and distillation of the liquids, Ir-H4 was recovered in excess of 90% and deuterated benzene and fluorobenzene were obtained. The experiments were repeated with the recovered Ir-H4 and deuterium with minimal loss of catalyst performance, as shown in Fig. 5.

Using a slightly modified methodology, both THF and n-heptane were partially deuterated and isolated in good yields. Ir-H8 was not soluble in THF/D2O mixtures, so a 6 M solution of NaOH in D2O was employed to render the medium biphasic. Heating Ir-H4 (1 mol%) dissolved in neat THF (8.3 mmol) under static vacuum to 100 °C in the presence of a D2O with NaOH (6 M) solution resulted in both the 2 and 3 positions were deuterated to 74% and 65%, respectively. Lower temperatures produced significantly lower yields as did performing the reaction under 1 atmosphere of argon. Deuterated THF was isolated via distillation and the catalyst was again recovered in 90% yield. NMR analysis of the recovered iridium material in THF revealed the mixture was composed of Ir-H4, Ir-H3 and Ir-OD (76%, 3% and 21%, respectively). All material was easily converted back to Ir-H4 by placing the solution under 4 atmospheres of H2 gas. In order to partially deuterate n-heptane slightly higher temperature (120 °C) was required. After 24 hours the material was isolated and NMR analysis revealed n-heptane had been substantially deuterated to 63% at the terminal methyl positions and 28% among all methylene positions.
Conclusions

Since our initial report in 2012 (ref. 35) on iridium complexes of the PCarbeneP framework, buttressed by robust aryl groups linking the arms of the pincer to the central carbene, the ligand design has evolved into high rigidity and high electron-donating ligand employed here to prepare and isolate Ir-H$_4$. Three features of this advanced ligand design make it a highly effective HIE catalyst system (Scheme 5). The electron richness of the ligand allows for the stabilization of the Ir-H$_4$ form of the catalyst such that it is isolable, storable, and able to fully regenerate if the system is exposed to excess dihydrogen. However, loss of one equivalent of H$_2$ is facile, allowing for access to the Ir-H$_2$/Ir-H$_3$ manifold, which is moderately active as an HIE catalyst and able to effect selective deuteration of sterically accessible positions on some substrates. We propose that further removal of H$_2$, formally speaking, allows access to an even more active Ir[II]Ir-H catalyst; this species, while not observed experimentally, is implied by the stoichiometry of the reactions that lead to its generation. This species is highly effective at activating a wide variety of C=H and C-D bonds via oxidative addition, including bonds normally reticent towards activation. While highly reactive, this species is robust under HIE conditions by virtue of the electron richness of the ligand.

A second useful feature in this context, is the ability of these compounds to cooperatively move hydrogen atoms from metal to pincer carbene via a low barrier 1,2-shift. This increases their hydrogen capacity in each HIE active manifold while also providing a mechanism for opening coordination space for hydrogen capacity in each HIE active manifold while also cooperatively move hydrogen atoms from metal to pincer carbene via a low barrier 1,2-shift. This increases their hydrogen capacity in each HIE active manifold while also providing a mechanism for opening coordination space for hydrogen capacity in each HIE active manifold while also facilitating the electron richness of the ligand. This has the effect of “flattening the curve” on the energy surface joining the PCarbeneP (as in Ir-H$_3$ and Ir-[]]) and PCarbeneP (e.g. Ir-H$_3$ and Ir-H) forms of the active species by making the C-H bond in the PCarbeneP form weaker and rendering the overall HIE process more exhaustive deuteration of a number of substrates, including on a preparative scale, under mild conditions and using Cd$_6$ or D$_2$O as a deuterium source.

Experimental section

General considerations

All compounds were stored, handled and manipulated, unless otherwise stated, using standard techniques in a VAC glovebox or on a double manifold high vacuum line under an atmosphere of argon purified via passage through an OxisorBW scrubber (Model 641-02 Matheson Gas Products). An M-Braun solvent purification system was used to dry n-pentane which was subsequently stored over sodium/benzophenone ketal in a 500 mL thick-walled glass pressure flask. Benzene, benzene-d$_6$, toluene, toluene-d$_6$, THF-d$_8$ and cyclohexane were dried over sodium/benzophenone ketal and stored under vacuum in a 100 mL thick-walled glass pressure flask. All dried solvents were degassed and vacuum distilled prior to use. Triethylamine, aniline, pyridine and 2,6-lutidine were dried over CaH$_2$ and distilled prior to use. Fluorobenzene and ethyl benzoate were dried over 3 Å molecular sieves and filtered off. Methylcyclohexane-d$_{14}$, cyclohexane-d$_{12}$, cyclooctane, and n-heptane were purchased from Sigma-Aldrich and degassed via freeze-pump-thaw prior to use. Phenol, N,N-dimethylaniline, and naphthalene were placed under 1 atmosphere of argon prior to use. H$_2$ (99.5%) and D$_2$ (99.98%) were purchased from Praxair and used without further purification. Iridium(III) chloride hydrate was purchased from Pressure Chemicals Inc. and used as received. [Ir(COE)$_2$Cl]$_2$ and [Ir-Cl] were prepared as previously reported. $^1$H and $^{13}$C(H) NMR chemical shifts were referenced to residual solvent protons and naturally abundant $^{13}$C resonances for all deuterated solvents. $^{65}$ Chemical shift assignments are based on $^1$H, $^{13}$P[$^1$H], $^{11}$C[$^1$H], $^1$H-1H-COSY, $^{1}$H-13C-HSQC and $^{1}$H-11C-HMBD NMR experiments performed on Bruker Ultrashield 400, Ascend-500, or Avance-600 MHz spectrometers using TopSpin version 3.2 software. X-ray crystallographic analyses were performed on a Nonius system equipped with a Bruker Apex-II CCD using samples coated in Paratone 8277 oil (Exxon) and mounted on a glass fibre. Elemental analyses were performed by staff of the Instrumentation Facility in the Department of Chemistry, University of Calgary.

Synthesis of Ir-0H

To a 100 mL thick walled round bottomed pressure flask charged with CsOH·H$_2$O (0.866 g, 5.16 mmol) and a Teflon stir bar, Ir-Cl (0.388 g, 0.516 mmol) dissolved in toluene (25 mL) was added. The flask was sealed and heated to 120 °C. After stirring for 24 hours the green-brown solution had turned dark brown. This solution was cooled to room temperature and filtered through a pad of Celite® into a 50 mL round bottom flask. All volatiles were removed from the filtrate in vacuo and n-pentane (20 mL) was vacuum transferred into the flask at −78 °C. The flask was warmed to room temperature and the solution was stirred for 2 hours. All volatiles were removed in vacuo. A brown powder (0.301 g, 4.101 mmol) was isolated in 80% yield. Elemental analysis calc (%): C, 52.37; H, 7.00; N, 3.82. Found (average of 4 trials): C, 53.41; H, 7.25; N, 3.59 (the carbon value is high due to small amounts of unremovable toluene). APCI-MS (M$^+$) for C$_{33}$H$_{31}$IrN$_2$OP$_5$: calc’d: 734.3100. Found: 734.3070.

1H NMR (500 MHz, Cd$_6$) δ 6.70 (dvt, $^1$J$_{HH}$ = 2.3 Hz, $^1$J$_{HP}$ = 3.7 Hz, 2H, ArH), 6.34 (d, $^1$J$_{HH}$ = 2.3 Hz, 2H, ArH), 3.64 (t, $^1$J$_{HP}$ = 3.8 Hz, 1H, Ir-0H), 2.95 (m, 4H, PCH(CH$_3$)$_2$), 2.36 (s, 12H, N(CH$_3$)$_2$), 1.65 (s, 6H (18H with PCH(CH$_3$)$_2$), -CH$_2$(CH$_3$)$_2$-), 1.63 (d, $^2$J$_{HH}$ = 7.5 Hz, $^2$J$_{HP}$ = 7.3 Hz, 12H (18H with -C(CH$_3$)$_2$-), PCH(CH$_3$)$_2$), 1.40 (dvt, $^2$J$_{HH}$ = 7.1 Hz, $^2$J$_{HP}$ = 7.1 Hz, 12H, PCH(CH$_3$)$_2$). $^{13}$C([H])
NMR (126 MHz, C6D6) δ 197.0 (t, JCP = 2.2 Hz, Ir=–C), 158.7 (vt, JCP = 18.8 Hz, ArC), 149.9 (vt, JCP = 3.9 Hz, ArC), 142.5 (vt, JCP = 20.0 Hz, ArC) 130.3 (vt, JCP = 7.8 Hz, ArC), 116.3 (s, ArCH), 114.4 (s, ArCH), 127.4 (s, –C(CH3)=), 120.2 (s, N(CH3)2), 28.2 (s, –C(CH3)=), 25.3 (vt, JCP = 12.8 Hz, PCH(CH3)2), 20.0 (vt, JCP = 2.7 Hz, PCH(CH3)2) 19.7 (s, PCH(CH3)2). 31P{1H} NMR (203 MHz, C6D6) δ 51.9 (s).

Synthesis of Ir-H4

Method 1: to a 100 mL thick walled glass pressure vessel equipped with a Teflon stirbar, Ir-OH (0.196 g, 0.267 mmol) dissolved in benzene (15 mL) was loaded. The mixture was degassed via the freeze–pump–thaw method and placed under 4 atm of H2 gas. After stirring for 3 hours all volatiles were removed in vacuo resulting in a dark red-brown residue. To this 0.78 atm of PCH(CH3)2 were added via vacuum transfer at −78 °C. The solution was warmed to room temperature and stirred for 20 minutes under 1 atm of H2. The n-pentane was removed under high vacuum and a beige powder (0.188 g, 0.260 mmol) was isolated in 57.8% yield. Elemental analysis calcd (%): C, 53.24; H, 6.78; N, 3.88. Found: C, 53.48; H, 8.11; N, 3.73. Method 2: to a 100 mL thick walled glass pressure vessel charged with NaOH (0.266 g, 6.65 mmol) and a Teflon stirbar, Ir-Cl (0.500 g, 0.665 mmol) dissolved in benzene (35 mL) was loaded. The mixture was degassed via the freeze–pump–thaw method and then placed under 4 atm of H2 gas. After stirring for 3 hours the now orange-yellow solution was degassed and filtered through a pad of Celite® in a glovebox. The filtrate was placed in a 50 mL round bottom flask and all volatiles were removed in vacuo. To the dark red residue n-pentane (25 mL) was vacuum transferred at −78 °C. The mixture was warmed to room temperature, placed under 1 atmosphere of H2 gas and stirred for 20 minutes. All volatiles were removed in vacuo and a beige powder (0.454 g, 0.627 mmol) was isolated in 94% yield. X-ray quality crystals were obtained by slowly cooling a supersaturated solution of Ir-H4 in isopropanol from 50 °C to 20 °C. Elemental analysis calcd (%): C, 53.42; H, 7.68; N, 3.88. Found: C, 53.43; H, 7.88; N, 3.80.1H NMR (500 MHz, C6D6) δ 6.84 (d, JHH = 1.9 Hz, 2H, ArH), 6.59 (d, JHH = 1.9 Hz, 2H, ArH), 6.48 (d, JHH = 1.9 Hz, 2H, ArH), 6.52 (s, 1H, Ir-CH), 2.85 (s, 12H, N(CH3)2), 2.10 (m, 2H, PCH(CH3)2), 1.91 (m, 2H, PCH(CH3)2), 1.84 (s, 3H, –C(CH3)=), 1.23 (s, 3H, –C(CH3)=), 1.16 (d, JHH = 7.9 Hz, JHP = 7.4 Hz, 6H, PCH(CH3)2), 1.07 (m, 12H, PCH(CH3)2), 0.78 (d, JHH = 8.0 Hz, JHP = 7.3 Hz, 6H, PCH(CH3)2), −10.32 (t, JHH = 10.7 Hz, 4H). 13C{1H} NMR (126 MHz, C6D6) δ 245.6 (s, Ir-C), 141.2 (s, ArCH), 135.4 (s, ArCH), 126.6 (s, PhCH), 120.8 (s, PhCH), 117.1 (s, ArCH), 41.3 (s, –C(CH3)=), 27.5 (d, JHH = 7.8 Hz, 1H, PCH(CH3)2), 20.1 (vt, JCP = 2.4 Hz, PCH(CH3)2) 19.3 (br s, PCH(CH3)2). 31P{1H} NMR (203 MHz, C6D6) δ 53.6 (s).

Generation of Ir-H3

To a J-Young tube charged with Ir-Ph (0.010 g, 0.025 mmol) dissolved in C6H6 (0.25 mL), Ir-H4 (0.013 g, 0.037 mmol) in C6H6 (0.35 mL) was added dropwise over 5 minutes. The solution changed from dark green to a dark reddish brown color. Ir-H3 is generated in situ and is isolable. 1H NMR (500 MHz, C6H6) spliced with C6D6 δ 6.78 (m, Ar-H), 6.73 (d, JHH = 2.5 Hz Ar-H), 2.58 (s, 12H, N(CH3)2), 2.25 (m, 4H, PCH(CH3)2), 1.63 (s, 6H, –C(CH3)=), 1.46 (d, JHP = 7.5 Hz, PCH(CH3)2), 1.41 (m, 12H, PCH(CH3)2), 1.37 (m, 2H, PCH(CH3)2), 1.07 (m, 12H, PCH(CH3)2), 0.78 (d, JHP = 8.0 Hz, JHP = 7.3 Hz, 6H, PCH(CH3)2), −10.32 (t, JHP = 10.7 Hz, 4H). 13C{1H} NMR (126 MHz, C6D6) δ 151.6 (vt, JCP = 15.4 Hz, ArC) 149.2 (vt, JCP = 4.3 Hz, ArC), 145.2 (vt, JCP = 7.4 Hz, ArC) 139.6 (vt, JCP = 24.4 Hz, ArC), 112.1 (s, ArCH), 111.6 (s, ArCH), 41.9 (s, N(CH3)2), 40.8 (s, –C(CH3)=), 30.3 (s, –C(CH3)=), 30.1 (vt, JCP = 14.8 Hz, PCH(CH3)2), 25.0 (s, –C(CH3)=), 23.1 (vt, JCP = 16.7 Hz, PCH(CH3)2), 21.0 (s, Ir-CH), 20.2 (s, PCH(CH3)2), 20.2 (vt, JCP = 2.2 Hz, PCH(CH3)2) 19.4 (vt, JCP = 3.7 Hz, PCH(CH3)2), 18.0 (s, PCH(CH3)2). 31P{1H} NMR (203 MHz, C6D6) δ 66.4 (s).

Preparation of stock solutions

Solution A: complex Ir-Ph (0.060 g, 0.076 mmol, 0.015 M) was dissolved with benzene-d6 in a volumetric flask (5 mL). Solution B: cyclohexane was added to a volumetric flask (5 mL) containing complex Ir-H4 (0.068 g, 0.094 mmol, 0.019 M) and cyclohexane-d12 (0.550 g, 5.07 mmol, 1.15 M). Solution C: cyclohexane was added to a volumetric flask (1 mL) containing Solution B (0.450 mL) and substrate (0.85 mmol).

General procedure for deuteration of substrates in C6D6

Stock Solution A (0.80 mL) was added to a glass reaction tube (length: 8.5 cm length, inner diameter: 1.5 cm) equipped with a Teflon stirbar. The flask was sealed with a rubber stopper and placed in a glovebox and tert-butylthylene (0.105 g, 1.069 mmol) was added. After stirring for 10 minutes all volatiles were removed in vacuo. The dark green-brown residue was then sonicated in n-pentane (5 mL). The solid was then filtered through a fritted Büchner funnel and washed with n-pentane (3 × 2 mL). A dark green powder (0.143 g, 0.180 mmol) was isolated in 73% yield. X-ray quality crystals were obtained from a saturated solution of Ir-Ph in n-pentane that was sealed at room temperature in a 20 mL scintillation vial for 1 week. Elemental analysis calcd (%): C, 57.48; H, 6.98; N, 3.53. Found: C, 57.59; H, 6.99; N, 3.52.
1.4 mol% (0.405 mL, 0.017 mmol at 20 °C) or 9 mol% (2.6 mL, 0.11 mmol) D₂ gas was injected into the flask. After stirring for 10 minutes at 20 °C, the solution (0.250 mL, 0.0038 mmol of [Ir]) was injected into a J.-Young tube containing substrate (0.38 mmol), cyclohexane (0.046 mmol) and benzene-d₆ (0.75 mL). The solution was heated to 65 °C for 24 hours and conversion was measured using cyclohexane as an internal standard. Cyclohexane was monitored without the use of an internal standard; no detectable traces of deuterium incorporation were visible using ¹H, ³H or ¹³C[¹H] NMR.

General procedure for deuteration of neat benzene, fluoroacetic acid and n-heptane with D₂O

Stock Solution C (0.450 mL) was added to a pressure tube (10 mL) charged with cyclohexane (0.300 mL), argon sparged D₂O (0.4 mL, 0.022 mmol) and a Teflon stirbar. Another sample was prepared by placing stock Solution C (0.450 mL) and cyclohexane (0.300 mL) in an NMR tube; this sample was used as a starting material reference. The flask was sealed under argon (1 atm) with a Teflon valve and heated to the appropriate temperature. After 24 hours the organic phase was carefully transferred to an NMR tube under argon without exposure to air. Conversion was measured via ¹H NMR using the cyclohexane signal as the internal standard. The reference samples containing water miscible/soluble compounds (THF, triethylamine, 2, 6-lutidine, pyridine, aniline and phenol) were extracted with D₂O prior to running NMR. The ¹H NMR spectra of the organic phases for these samples were compared to the organic phases of the reaction mixtures to measure conversion. Aliquots of the D₂O fractions (0.2 mL) for both the reference and reaction mixtures were also taken and added to separate NMR samples containing D₂O (0.4 mL) and DMSO (10 µL). The ¹H NMR spectra for these samples indicated comparable levels of deuteration were present in the D₂O phase.

General procedure for deuteration of neat benzene, fluoroacetic acid and n-heptane with D₂O

Substrate (8.31 mmol) was loaded into a pressure tube (10 mL) charged with Ir[H₄] (0.060 g, 0.083 mmol) and a Teflon stirbar. Argon sparged D₂O (6.0 mL, 330 mmol) was carefully added to the reaction flask under a steady flow of argon. The vessel was sealed with a Teflon valve and heated to 80 °C for 24 hours. The flask containing n-heptane was freeze–pump–thawed three-fold and heated to 120 °C under static vacuum for 24 hours. Organic phases were extracted under a flow of argon via syringe and liquids were distilled under reduced pressure from the iridium catalyst. Benzene was recovered in 81% yield (0.561 g, 6.72 mmol) with 90% deuterium incorporation. Fluoroacetic acid was recovered in 82% yield (0.725 g, 6.81 mmol) with 94% deuteration incorporation at all positions. n-Heptane was recovered in 88% yield (0.791 g, 7.40 mmol) with 63% deuterium at the terminal methyl positions and 28% deuteration incorporation throughout the methylene position.

Deuteration of THF with D₂O

THF (0.600 g, 8.32 mmol) was added to a pressure tube (10 mL) charged with Ir[H₄] (0.060 g, 0.083 mmol) and a Teflon stirbar. Argon sparged D₂O (6 mL) containing NaOH (1.45 g, 36.3 mmol, 6 M) was added to the flask under a flow of argon. The mixture was freeze–pump–thawed three-fold and sealed under static vacuum. After heating at 100 °C for 24 hours the organic phase was extracted under a flow of argon via syringe and distilled under reduced pressure from the iridium catalyst. Deuterated THF was recovered in 84% yield (0.547 g, 7.03 mmol) with deuterium incorporated in 74% and 65% at the 2 and 3 positions, respectively.

Conflicts of interest

There are no conflicts to declare.

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Notes and references


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