Base-induced reversible H₂ addition to a single Sn(II) centre†

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A range of amines catalyse the oxidative addition (OA) of H₂ to [(Me₅Si)₂CH₂Sn(II)] (1), forming [(Me₅Si)₂CH₂SnH₂] (2). Experimental and computational studies point to ‘frustrated Lewis pair’ mechanisms in which 1 acts as a Lewis acid and involves unusual late transition states; this is supported by the observation of a kinetic isotope effect (KIE, k_H₂/k_D₂ = 1.51 ± 0.04) for Et₃N. When DBU is used the energetics of H₂ activation are altered, allowing an equilibrium between 1, 2 and adduct [1:DBU] to be established, thus demonstrating reversible oxidative addition/reductive elimination (RE) of H₂ at a single main group centre.

The ability of L₂Sn(n) compounds to undergo OA has been inversely correlated with the size of the singlet-triplet (HOMO-LUMO) gap, which may be diminished through the use of extremely strong σ-donor ligands. Aldridge et al. have employed a bis(boryl)tin(n) system to achieve the only example of direct OA of H₂ to a mononuclear Sn(n) centre, irreversibly forming the Sn(n) dihydride; boryl ligands are even stronger σ-donors than hydride or alkyl ligands, permitting a successful reaction outcome.†d

Introduction

In the past decade there has been significant interest in transition metal (TM) free systems which activate H₂.† Two main strategies have emerged to facilitate this reactivity: the use of low-valent main group (MG) compounds,‡ and so-called ‘frustrated Lewis pairs’ (FLPs).§ In both cases, reactivity arises from simultaneously having access to a high-lying HOMO and low-lying LUMO (Fig. 1). Various low-valent MG compounds containing multiple E–E bonds (E = Al, Si, Ga, Ge, Sn),☆ or single-site low-valent centres such as carbenes and heavier tetrylene analogues, have been shown to react with H₂.¶ The scope of Lewis bases (LBs) and, to a lesser extent, Lewis acids (LAs), which can be used in H₂-activating FLPs has expanded to include a number of elements from across the periodic table. This is principally due to the readily tuneable steric and electronic profiles of the individual LA and LB sites.¶•☆ Many FLP systems display reversible H₂ cleavage, which has facilitated their rapid expansion into the field of catalytic hydrogenation.¶•☆ The same is not true for low-valent MG compounds; examples of reversible H₂ activation are very rare and limited to antiaromatic boracycles,¶¶ a phosphorus-based singlet biradicaloid,¶∥ and only one low-valent group 14 compound: a dinuclear Sn(i) distannylene.¶‡ The design of single-site MG systems which are ergoneutral for H₂ activation requires fine-tuning of thermodynamic (e.g. weak E–H bond strengths promoting an accessible formal Eⁿ⁺/Eⁿ couple) and kinetic factors, both of which are constrained to a mononuclear species, and is hence especially challenging.

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Fig. 1. Representative orbital interactions between H₂ and main group compounds: (a) unsaturated E–E compounds e.g. distannynes (Ar = C₆H₆; 2.6–(C₆H₅)₂Sn₂; 2.6–(C₆H₅)₂: X = H, SiMe₃, F); for X = H, the reaction is reversible at 80 °C;¶¶ (b) single site low-valent centres e.g. carbenes;¶∥ (c) sterically hindered LAs and LBs (FLPs); (d) this work.
Conversely, the irreversible base-induced RE of H₂ from organostannanes is well-known.²⁴ Wesemann and others have studied RE from ArSnHₓ and [(Me₃Si)₂CH]SnHₓ compounds to yield various mononuclear Sn and Sn–Sn bound species (Ar = terphenyl).²⁵ Nevertheless, there has yet to be a report of reversible OA and RE occurring on a single Sn(n) scaffold. Lappert's stannylene [(Me₃Si)₂CH]₂Sn (1), which can act as both Lewis acid (LA) and base (LB), is a paradigmatic system for investigating OA to low-valent MG centres, yet to date its reactivity with H₂ has been unexplored.²⁶ Herein we report the use of FLP methodology to promote formal OA of H₂ to this simple dialkylstannylene. Furthermore, we document the first example of reversible H₂ addition to a single-site MG complex, which accesses an FLP via reversible dissociation of a classical 1-LB adduct; formation of the latter renders OA of H₂ to 1 energetically less favourable, enabling RE to occur from the Sn(n) dihydride and reform 1, which is in equilibrium with 1-LB.²⁷

### Results and discussion

1 is in a rapid solution-phase equilibrium with its dimer [1]₂, which has been crystallographically characterised and contains a formal Sn=Sn double bond.²⁸ When a d₆-toluene solution of 1/[1]₂ was placed under an atmosphere of H₂ (4 bar) in a sealed NMR tube, no change was observed in the ¹H NMR spectrum, even after prolonged periods (>48 h), confirming that neither 1 nor [1]₂ can react with H₂ alone. Separately, addition of Et₃N (20 mol%) to a solution of 1 resulted in no perturbation of their ¹H NMR resonances, suggesting no interaction between the components; i.e. the formation of an FLP.²⁹ Placing this new mixture under H₂ (4 bar, RT) resulted in the solution turning from deep red to colourless over the course of 24 h, with the ¹H NMR spectrum revealing complete consumption of 1 and a new Sn–H triplet resonance at δ = 5.11 ppm [J(¹H–¹H) = 2.2 Hz] with attendant satellites [J(¹H–¹¹Sn–¹H) = 1704 Hz; J(¹¹Sn–¹¹Sn–¹H) = 1784 Hz], in addition to signals for the Si(CH₃)₃ and methine protons [δ/ppm = 0.17 (s) and -0.42 (t, J(¹H–¹H) = 2.2 Hz), respectively] (see Fig. 2). ¹¹Sn NMR spectroscopy showed only a triplet of triplets at −196 ppm [J(¹¹Sn–¹¹Sn–¹H) = 1784 Hz, J(¹¹Sn–¹¹Sn–¹H) = 87 Hz] which collapsed to a singlet upon ¹H decoupling. Collectively these data correspond to the previously unreported dihydride [(Me₃Si)₂CH]₂SnH₂ (2), which was confirmed by comparison with an authentic sample prepared by the reaction of LiAlH₄ and [(Me₃Si)₂CH]₂SnCl₂ (see ESIF† for details).

#### Isotopic investigation

When D₂ was used in place of H₂, the methine peak present in the ¹H NMR spectrum of the product mixture resolved as a singlet, while the Sn–H signal was absent and replaced by a Sn–D signal at δ = 5.11 ppm [J(¹¹Sn–¹H–¹H) = 262 Hz, J(¹¹Sn–¹¹Sn–¹H) = 274 Hz] in the ²H NMR spectrum. These results demonstrate the formation of dideuteride 2-D₂,²⁸ and that the Sn-bound protons in 2 must originate from the hydrogen atmosphere.

In order to probe the mechanism further, a d₆-toluene solution of 1/[1]₂ and Et₃N was reacted with a 1 : 1 mixture of H₂/D₂. The resultant ¹H NMR spectrum was very similar in appearance to that of 2, with two exceptions: the relative integration of the Sn–H peak did not match that of the methine signal (1.2 : 2; consistent with the faster rate of reaction with H₂ vs. D₂ – vide infra), and the C–H resonance was composed of overlapping peaks commensurate with a mixture of 2 and 2-D₂.

No spectroscopic evidence was seen for the formation of 2-HD, which was independently and selectively obtained by analogous reaction of 1/[1]₂ under an HD atmosphere. These observations provide strong evidence that delivery of both atoms from H₂/D₂/HD to a single Sn centre occurs either simultaneously, or in a near-concerted fashion.

#### Kinetic analysis

By analogy with established FLP systems, and the microscopic reverse of the polar mechanism by which dehydrogenation of ArSnHₓ species is proposed to occur,³⁰ we envisaged a reaction mechanism in which 1 and Et₃N form a weakly associated ‘encounter complex’ which subsequently reacts with H₂ (Scheme 1).³¹ Assuming that encounter complex formation is a rapid pre-equilibrium prior to rate-limiting H₂ activation gives the expected rate law: rate = k′[Et₃N][H₂], where k′ = (k₁k₃)/k₋₁. Calorimetric studies on H₂ activation by the FLP Mes₃P/B(C₆F₅)₃ (Mes = 2,4,6-C₆Me₃) found the rate to be very accurately modelled as a single, termolecular step, which formally gives the same rate law.²²

To confirm the order of catalytic Et₃N, the method of time (t) scale normalisation was used;²³ normalisation to the scale of t·[Et₃N]ₓ resulted in the superposition of all reactant traces only when x = 1, confirming the rate to be first order with respect to the amine (Fig. 3a). Determination of reaction order with respect to 1 requires its concentration to be known accurately at any given time in a reaction mixture. However, since the
observed $^1$H NMR resonances are a weighted average of the signals from 1 and $[1]_2$ ($\Delta G_{293K} = 3.1 \text{ kcal mol}^{-1}$), with both species present at significant concentrations under reaction conditions, simple observation of the concentration of 1 is not directly possible by $^1$H NMR spectroscopy.$^{16}$ The concentration of 1 can, however, be calculated from the total concentration of “$R_2Sn$” species in solution, [Tot], present as either monomer or dimer, which are related to the concentrations of 1 and $[1]_2$ by:

$$\text{[Tot]} = [1] + 2([1]_2) \tag{1}$$

The dimerisation equilibrium of 1 can be expressed as:

$$K_0 = \frac{[1]^2}{([1]_2)} \tag{2}$$

Combining eqn (1) and (2) and solving for [1] yields:

$$[1] = \frac{1}{4} \left( \sqrt{K_0} \sqrt{8\text{[Tot]}} + K_0 - K_0 \right) \tag{3}$$

Inserting eqn (3) into the expected rate law (vide supra) gives:

$$\frac{-d[\text{Tot}]}{dt} = \frac{k^*}{4} \left( \sqrt{K_0} \sqrt{8\text{[Tot]}} + K_0 - K_0 \right) \tag{4}$$

where, if the amount of $H_2$ is sufficiently high that its concentration remains approximately constant:

$$k^* = \frac{k_1k_2}{k_{-1}} [B][H_2] \tag{5}$$

Rearrangement and integration by substitution of eqn (4) (see ESI†) gives:

$$\sqrt{\frac{8\text{[Tot]}}{K_0}} + 1 + \ln \left( \frac{\sqrt{K_0} + \sqrt{8\text{[Tot]}} - \sqrt{K_0}}{\sqrt{K_0} + \sqrt{8\text{[Tot]}} - \sqrt{K_0}} \right) = \frac{\sqrt{8\text{[Tot]}}}{K_0} + 1$$

$$= -k^* t \tag{6}$$

Therefore, plotting the variable portion of the LHS of this expression against $t$ gives a straight line of gradient $-k^*$, confirming the proposed first-order dependence on 1 (Fig. 3b).

Using the known value of $[H_2]$ in toluene at 4 bar (293 K)$^4$ provides a value of $k'_{[H_2]} = 0.47 \pm 0.03 \text{ M}^{-1} \text{s}^{-1}$. As well as $Et_3N$, 2-tert-butyl-1,1,3,3-tetramethylguanidine (Barton’s base, TBTMG) and 1,2,2,6,6-pentamethyldiisopropylidine (PMP), were also found to form FLPs with 1/[1]$_2$, with corresponding rates of $H_2$ cleavage: $k'_{[TBTMG]} = 5.0 \pm 0.3 \text{ M}^{-1} \text{s}^{-1}$, $k'_{[PMP]} = 0.0266 \pm 0.0018 \text{ M}^{-1} \text{s}^{-1}$. Despite the similar basicity to $Et_3N$, the bulkier Hünig’s base ([P(H$_2$)$_2$]$^+$/[P$_8$H$_8$]$^-$) was ineffective for $H_2$ heterolysis, as was the weaker base 2,4,6-collidine ([P$_8$H$_8$]$^-$) $^{14}$ Clearly $H_2$ activation requires that the LB be sufficiently basic and not too sterically encumbered, in line with observations of other FLP systems.$^{29}$

A kinetic analysis of the isotopic systems permitted quantification of the KIE: $k'_{[H_2]}/k'_{[D_2]} = 1.51 \pm 0.04$ when $Et_3N$ was used as the base. In addition, the acceleration in rate from a more polar solvent could also be quantified: $k'_{[Tol]}^{[THF]}/k'_{[toluene]} = 1.97 \pm 0.04$ (when $Et_3N$ was used).
Coordinating bases

When the less sterically bulky 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) is used, an interaction with 1 can be clearly seen in the $^{13}$C($^1$H) NMR spectrum: upon gradual addition of DBU to 1/\([1]_2\), the methine resonance undergoes a substantial upfield shift, reaching a limiting value of $\delta = 18.5$ ppm (10-fold excess of DBU). Using the established $^{13}$C NMR chemical shift values for 1 and \([1]_2\) (60.0 ppm and 28.7 ppm, respectively),\textsuperscript{38e} this is consistent with a fast equilibrium between 1-DBU, 1 and \([1]_2\) (Scheme 2; see ESI\textsuperscript{f} for full details). A value of $\Delta G = -3.7 \pm 0.2$ kcal mol$^{-1}$ for the formation of 1-DBU from \([1]_2\) was obtained from a van’t Hoff analysis of variable temperature UV-Vis spectra.

While the reaction of 1/DBU mixtures (containing 0.1–10 equivalents of DBU) with H$_2$ proceed rapidly, they do not reach completion, indicative of a reversible process (see Fig. S7 in ESI\textsuperscript{f}).

The reversibility can be explicitly demonstrated by the (CH$_3$)$_3$Si region of the $^1$H NMR spectrum, whereby addition of DBU to a solution of 2 led to the appearance of a signal corresponding to the dehydrogenated mixture 1-DBU $\leftrightarrow$ 1 $\leftrightarrow$ [1]$_2$; this increased in intensity at the expense of the (CH$_3$)$_3$Si peak of 2 (Fig. 4a–c). No H$_2$ is observed in the $^1$H NMR spectrum as the solution was degassed multiple times in order to accelerate the reaction – however, the very small amount of H$_2$ generated (approx. 0.3 bar) would likely hamper detection. Furthermore, the methine resonance of the 1-DBU $\leftrightarrow$ 1 $\leftrightarrow$ [1]$_2$ mixture is subject to a significant upfield shift compared to [1][1]$_2$ (dependent upon the DBU concentration), and so is obscured beneath the relatively intense (CH$_3$)$_3$Si region. Upon charging this reaction with H$_2$, restoration of 2 was rapidly observed (Fig. 4d). For the equilibrium involving H$_2$ (Scheme 2), an equilibrium constant, $K_{eq} = 164 \pm 5$, in favour of 2 can be calculated from the relative intensities of the (CH$_3$)$_3$Si resonances, providing $\Delta G = -3.0$ kcal mol$^{-1}$ (1 bar H$_2$).

Using the similarly unhindered but less basic 4-(dimethyl amino)pyridine (DMAP) also gave an adduct 1-DBU, but no reaction with H$_2$ at room temperature. However, heating a solution of 1 with excess DMAP (4 bar H$_2$, 2 h, 100 °C) yielded 2 in 31% conversion.

Computational investigation

To gain further insight into the mechanism of H$_2$ activation, DFT calculations were performed for various 1/LB pairs;\textsuperscript{39} the computed reaction profiles for both the Et$_3$N- and DBU-mediated reactions are depicted in Fig. 5. When LB = Et$_3$N, the reaction was found to proceed via initial H$_2$ heterolysis leading to a tight ion pair intermediate [H][Et$_3$NH]$^+$ (int$_1$). Facile rearrangement to int$_2$ and subsequent delivery of the H$^+$ to the lone pair on the [H]$^+$ moiety furnishes 2 (Fig. 6a); a very similar mechanism was found when LB = DBU. In support of this polar mechanism, the rate using Et$_3$N as the LB was found to be faster in THF ($k_1$/[THF]/$k_{(toluene)}$ = 1.97 ± 0.04). The low

![Graphical representation of reaction profiles](image-url)
step for reactions with di
PA activation free energy.

with 1 free \([\text{bases DBU (Fig. 6b) and DMAP, adducts favourable relative to tent with experimental reaction rates. For the coordinating D}

reduces the absolute value of \(\text{D}

and int1. All \(\text{bases are compiled in Table 1}

barriers to rearrangement of the intermediates also offer an explanation as to why H/D exchange is not observed upon reaction with an H\(_2\)/D\(_2\) mixture or HD: collapse of the ion pairs is likely much faster than solvent cage escape.

Although to the located transition states (TSs) are energetically close-lying, the overall reaction barrier appears to be determined by the H\(_2\) splitting step, which is in line with kinetic measurements. Free energy data computed for the H\(_2\) splitting step for reactions with different bases are compiled in Table 1 alongside other properties. For Et\(_3\)N, TBTMG and PMP, no favourable adduct formation was found with 1, and the \(\Delta G^1\) values follow the order TBTMG < Et\(_3\)N < PMP, which is consistent with experimental reaction rates. For the coordinating bases DBU (Fig. 6b) and DMAP, addsucts favourable relative to free \([1]_2\) and base were computationally determined. This reduces the absolute value of \(\Delta G_{\text{reaction}}\) such that an

equilibrium is experimentally observed in the case of DBU. For DMAP, the activation barrier is found to be much higher, paralleling results seen by experiment where elevated temperatures are required to obtain product 2.

The energies of all intermediates int\(_1\) are computed to be well above the reference state, which follows from the weak Lewis acidity of 1. The stabilities of int1 species correlate very well with the general trend in PA and pK\(_a\), but this is not strictly true for the TSs, where steric factors are more important. Unstable int1 intermediates imply late TSs for the H\(_2\) activation step, which is shown by significantly elongated H–H distances in the TS structures. The experimentally observed KIE (1.51 \pm 0.04) supports this finding, which is commensurate with rate-limiting H\(_2\)/D\(_2\) activation involving considerable H–H/D–D bond breaking.\(^{32}\)

Conclusions

In conclusion, we have demonstrated the ability of FLP-mediated reactivity to enable the formal oxidative addition of H\(_2\) to an otherwise inert MG centre, and in doing so have also observed the first example of reversible H\(_2\) addition to a single-site MG complex. We have utilised experimental and computational means to comprehensively explore the mechanism of this transformation and found that H\(_2\) activation in this system differs from those based on more typical FLPs, due to the high-energy nature of the immediate H\(_2\) splitting products, resulting in rare examples of late TSs. The development of methods to harness this FLP-promoted OA/RE H\(_2\) reactivity for hydrogenation catalysis is currently underway.

Conflicts of interest

There are no conflicts to declare.

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