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## Reduction of carbon dioxide and organic carbonyls by hydrosilanes catalysed by the perrhenate anion<sup>†</sup>

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The simple perrhenate salt [N(hexyl)<sub>4</sub>][ReO<sub>4</sub>] acts as a catalyst for the reduction of organic carbonyls and carbon dioxide by primary and secondary hydrosilanes. In the case of CO<sub>2</sub>, this results in the formation of methanol equivalents via silylformate and silylactal intermediates. Furthermore, the addition of alkylamines to the reaction mixture favours catalytic amine *N*-methylation over methanol production under certain conditions. DFT analysis of the mechanism of CO<sub>2</sub> reduction shows that the perrhenate anion activates the silylhydride forming a hypervalent silicate transition state such that the CO<sub>2</sub> can directly cleave a Si-H bond.

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### Introduction

The hydrosilylation of CO<sub>2</sub> to silylformates and silylethers is an attractive route for CO<sub>2</sub> utilisation as, unlike hydrogenation, this reaction is exergonic due to the comparative ease of Si-H bond activation and the strength of the Si-O bond, and the availability of relatively inexpensive and environmentally benign hydrosilanes.<sup>1–3</sup> A variety of catalysts have been discovered and developed for this reaction, including complexes of Co,<sup>4</sup> Cu,<sup>5–8</sup> Ir,<sup>9–14</sup> Ni,<sup>15–17</sup> Pd/Pt,<sup>18</sup> Rh,<sup>19</sup> Ru,<sup>20–25</sup> Sc,<sup>26</sup> Zn,<sup>27–29</sup> and Zr,<sup>30</sup> frustrated Lewis pairs,<sup>31–36</sup> organocatalysts,<sup>37–42</sup> alkali metal carbonates,<sup>43</sup> and even polar solvents such as DMF.<sup>38</sup> These reactions result in various reduction products including silyl-formates, -acetals, and -ethers, CO, and methane. This reactivity has been further exploited in a ‘diagonal’ approach to CO<sub>2</sub> reduction/functionalisation, for example in the use of CO<sub>2</sub> as the C1 source for amine *N*-methylation.<sup>3,37,44–54</sup> High oxidation state transition metal oxo complexes<sup>55</sup> have been studied in CO<sub>2</sub> reduction chemistry and have been shown to react to form metal carbonates where, in the case of molybdate, the resulting carbonate undergoes hydrosilylation to formate and silylated molybdate, albeit stoichiometrically.<sup>56–59</sup> Furthermore, rhenium(v) mono and dioxo complexes act as catalysts for the hydrosilylation of organic carbonyls,<sup>60–65</sup> but this activity is not generally seen for the parent perrhenate, ReO<sub>4</sub><sup>−</sup>. Sig-

nificantly, metal oxo complexes have the potential to overcome the oxygen and moisture sensitivity issues from which many current CO<sub>2</sub> hydrosilylation catalysts suffer.

It was shown previously that the pyridinium perrhenate salt [4,6-Bu<sub>2</sub>C<sub>5</sub>H<sub>2</sub>NH-2-CH{(CO)NH(C<sub>6</sub>H<sub>13</sub>)<sub>2</sub>}]<sub>2</sub>[ReO<sub>4</sub>] **1** acts as a catalyst for the epoxidation of alkenes by hydrogen peroxide under biphasic conditions.<sup>66</sup> Important to this chemistry is the formation of a lipophilic ion pair in which electrostatic and hydrogen-bonding interactions are enhanced in the hydrophobic phase. This catalyst has the advantages of ease of synthesis and manipulation and lack of air-sensitivity, and the recovery of the precious metal is driven by straightforward reverse phase-transfer. Given the efficacy of this new catalyst system and the use of high oxidation state rhenium oxo complexes in reduction catalysis, it is shown here that perrhenate can act as a catalyst for the reduction of carbon dioxide to methanol equivalents by hydrosilanes, that the methylation of alkylamines using CO<sub>2</sub> as the C<sub>1</sub> source occurs under similar catalytic conditions, and that this method can be used to reduce aldehydes and ketones to silylalcohols.

### Results and discussion

#### Reduction of carbon dioxide

Initially, the reaction between CO<sub>2</sub> (2.5 bar), PhSiH<sub>3</sub> (2.0 mmol), and pyridinium perrhenate **1** (2.0 mol%) at 80 °C in C<sub>6</sub>D<sub>6</sub> in a Teflon-tapped NMR tube was monitored by <sup>1</sup>H NMR spectroscopy. However, only 15% consumption of hydrosilane is seen under these conditions after 16 h (Fig. S2 and S3, ESI<sup>†</sup>). In contrast, when the reaction is carried out using the simple lipophilic *quaternary* ammonium salt [N(hexyl)<sub>4</sub>][ReO<sub>4</sub>] **2** (2.5 mol%) with PhSiH<sub>3</sub> in C<sub>6</sub>D<sub>6</sub> at 1 bar of

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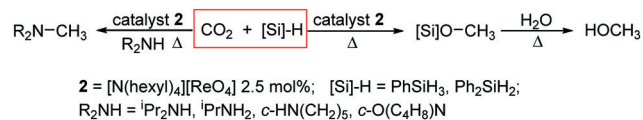
<sup>†</sup> Electronic supplementary information (ESI) available: Full experimental details and spectra. See DOI: 10.1039/c7cy00772h



CO<sub>2</sub> and at 80 °C, complete consumption of the hydrosilane is observed in the <sup>1</sup>H NMR spectrum after 12 h (Fig. S4†) along with the formation of silylformate (3), bis(silyl)acetal (4) and silylated methanol (5) products (Table 1, entry 1 and Scheme 1). Negligible depletion of hydrosilane (<3%) is seen when the reaction is carried out in the absence of CO<sub>2</sub> (under air) for 1 h (Fig. S5 and S6†), confirming that CO<sub>2</sub> is integral to the reaction. This air-stable perrhenate catalyst is also integral as no reaction is seen in the absence of 2 nor in the presence of 5 mol% of N(hexyl)<sub>4</sub>Br (Fig. S7†). Increased turnover and selectivity towards methanol equivalents is observed when switching solvents to the more polar CD<sub>3</sub>CN (Table 1, entries 3–6). In this case, complete consumption of hydrosilane is observed in 4 h after which the addition of water and heating at 80 °C for a further 2 h results in the formation of methanol, seen as a singlet at 3.28 ppm in the <sup>1</sup>H NMR spectrum (e.g. Fig. S9†). Prolonged reaction times to ensure full consumption of all Si–H containing species, i.e. no mono/oligomeric siloxyhydride resonances remaining at ca. 5.5 ppm, result in an increase in yield of 5. GC-MS analysis of the reduction with Ph<sub>2</sub>SiH<sub>2</sub> in CD<sub>3</sub>CN prior to quenching indicates the sole presence of Ph<sub>2</sub>Si(OMe)<sub>2</sub> confirming that all silylhydrides are consumed in the reaction (Table 1, entry 6, Fig. S60†).

The choice of hydrosilane is important to the reaction as Et<sub>3</sub>SiH (Table 1, entry 7) showed no conversion after 24 h at 80 °C, whereas (EtO)<sub>3</sub>SiH (Table 1, entry 8) is highly selective towards the formation of silylformate 3, and the use of more sterically demanding Ph<sub>3</sub>SiH (Table 1, entry 12) required elevated temperatures. This variation in reactivity is due to a combination of steric inaccessibility of the Si–H bond and the lower Lewis acidity of alkyl silanes.

Further solvent screening found the catalysis to be accelerated in pyridine, with complete hydrosilane consumption observed after 1.5 h (Table 1, entry 9). Using d<sub>7</sub>-DMF accelerates the reaction further, with complete hydrosilane depletion after 1 h and an increase in selectivity towards methanol formation compared to the reaction carried out in pyridine (Table 1, entries 10 and 11). Significantly however, reactions in DMF suffer from competitive solvent reduction by hydrosilane to (CD<sub>3</sub>)<sub>2</sub>NCDH<sub>2</sub>, consuming ca. 10% of the hydro-



**Scheme 1** Reactions of carbon dioxide with hydrosilanes catalysed by the lipophilic perrhenate salt 2.

silane in this process (Fig. S21†). The variation of solvent relates to an increase in turnover frequencies (TOFs), determined from the time taken to consume the hydrosilane at 80 °C, of 3.3 h<sup>-1</sup> for C<sub>6</sub>D<sub>6</sub> to 40 h<sup>-1</sup> for d<sub>7</sub>-DMF. The increase in rate is, in part, due to the increased solubility of CO<sub>2</sub> in the more polar solvent; indeed, a reaction carried out in CD<sub>3</sub>CN under 2 bar of CO<sub>2</sub> shows complete hydrosilane depletion within 1.5 h (Fig. S22†).

The longevity of the catalyst system was probed, and at the end of the first catalytic run with Ph<sub>2</sub>SiH<sub>2</sub> in d<sub>7</sub>-DMF (TOF 40 h<sup>-1</sup>), the NMR tube was recharged with Ph<sub>2</sub>SiH<sub>2</sub> and 1 bar of CO<sub>2</sub> and heated again at 80 °C; this procedure was repeated 3 times. A marginal drop in TOF is observed on the third recharge cycle (36 h<sup>-1</sup>) after which the reaction was quenched with water and selective formation of methanol is seen (ratio 1:0:99, Fig. S23†). Whilst only a small drop in activity was noted after subsequent recharges, the perrhenate catalyst has significant advantages over current reports due to the ease in which the perrhenate anion can be recovered through simple reverse-phase-transfer techniques.

The faster rate of the reaction in d<sub>7</sub>-DMF allowed its monitoring by <sup>1</sup>H and <sup>13</sup>C NMR spectroscopy at 300 K using <sup>13</sup>CO<sub>2</sub>. On depletion of hydrosilane, broad resonances are seen after 3 h consistent with the formation of silylformate 3 at 8.65 ppm (d, *J*<sub>HC</sub> 219 Hz) and 162.6 ppm (d, *J*<sub>CH</sub> 219 Hz) in the <sup>1</sup>H and <sup>13</sup>C NMR spectra, respectively (Fig. S24–S29†). After 5 h, quartet resonances at 51.9 ppm (*J*<sub>CH</sub> 144 Hz) appear in the <sup>13</sup>C NMR spectrum, consistent with [Si]OCH<sub>3</sub> 5 formation, along with some small triplet resonances at 82.8 ppm (*J*<sub>CH</sub> 171 Hz) for the bis(silyl)acetal [Si]OCH<sub>2</sub>O[Si] 4. In d<sub>7</sub>-DMF, the reaction between Ph<sub>2</sub>SiH<sub>2</sub> and CO<sub>2</sub> in the absence of perrhenate catalyst also leads to silylformate and silyl ether reduction products in a ratio of 38:62.<sup>38</sup> However, this

**Table 1** Hydrosilylation of carbon dioxide catalysed by 2.5 mol% [N(hexyl)<sub>4</sub>][ReO<sub>4</sub>] (2) under 1 bar of CO<sub>2</sub>, 0.2 mmol of hydrosilane at 80 °C

	Silane	Time (h)	Solvent	Conversion <sup>a</sup> (%)	Ratio 3:4:5 <sup>b</sup>	TOF <sup>c</sup> (h <sup>-1</sup> )
1	PhSiH <sub>3</sub>	12	C <sub>6</sub> D <sub>6</sub>	95	51:18:22	3
2	Ph <sub>2</sub> SiH <sub>2</sub>	12	C <sub>6</sub> D <sub>6</sub>	100	73:16:11	3
3	PhSiH <sub>3</sub>	3	CD <sub>3</sub> CN	96	15:0:85	10
4	Ph <sub>2</sub> SiH <sub>2</sub>	4	CD <sub>3</sub> CN	100	16:0:84	13
5	PhSiH <sub>3</sub> <sup>d</sup>	16	CD <sub>3</sub> CN	100	2:0:98	3
6	Ph <sub>2</sub> SiH <sub>2</sub> <sup>d</sup>	16	CD <sub>3</sub> CN	100	7:0:93	3
7	Et <sub>3</sub> SiH	24	CD <sub>3</sub> CN	0	0	0
8	(EtO) <sub>3</sub> SiH	24	CD <sub>3</sub> CN	15	99:0:1	0.3
9	Ph <sub>2</sub> SiH <sub>2</sub>	1.5	d <sub>5</sub> -py	100	38:0:62	27
10	PhSiH <sub>3</sub>	1	d <sub>7</sub> -DMF	100	10:0:90	40
11	Ph <sub>2</sub> SiH <sub>2</sub>	1	d <sub>7</sub> -DMF	100	16:0:84	40
12	Ph <sub>3</sub> SiH <sup>e</sup>	8	d <sub>7</sub> -DMF	25	21:0:79	1

<sup>a</sup> Consumption of hydrosilane. <sup>b</sup> NMR ratio vs. Me<sub>3</sub>SiPh after water quench. <sup>c</sup> TOF = (conversion/catalyst loading)/time. <sup>d</sup> Prolonged reaction times to consume all Si–H containing species. <sup>e</sup> 150 °C.



background reaction is very slow, taking 144 h to consume the  $\text{Ph}_2\text{SiH}_2$  at 300 K compared with 10 h when using perrhenate (Fig. S30<sup>†</sup>).

A series of stoichiometric reactions were undertaken to probe the catalytic mechanism. It was first thought that the mechanism may follow a similar route to the molybdate dianion which reacts initially with  $\text{CO}_2$  to form the carbonate complex  $\text{Mo}(\text{CO}_3)\text{O}_3^{2-}$ ; <sup>56</sup> however, no reaction of 2 with  $^{13}\text{CO}_2$  in the absence of hydrosilane is seen, even after prolonged heating at 80 °C (Fig. S31<sup>†</sup>), thus ruling out initial carbonate formation in this catalytic cycle. Also, no intermediates are seen in stoichiometric reactions of 2 with varying equivalents of hydrosilane in the absence of  $\text{CO}_2$ . It is of note, however, that upon extended heating (>16 h at 80 °C) decomposition of 2 occurs to form trihexylamine, siloxanes and a brown precipitate containing rhenium oxide clusters (Fig. S32–S34<sup>†</sup>). In light of these experiments, the mechanism of this reaction was evaluated computationally.

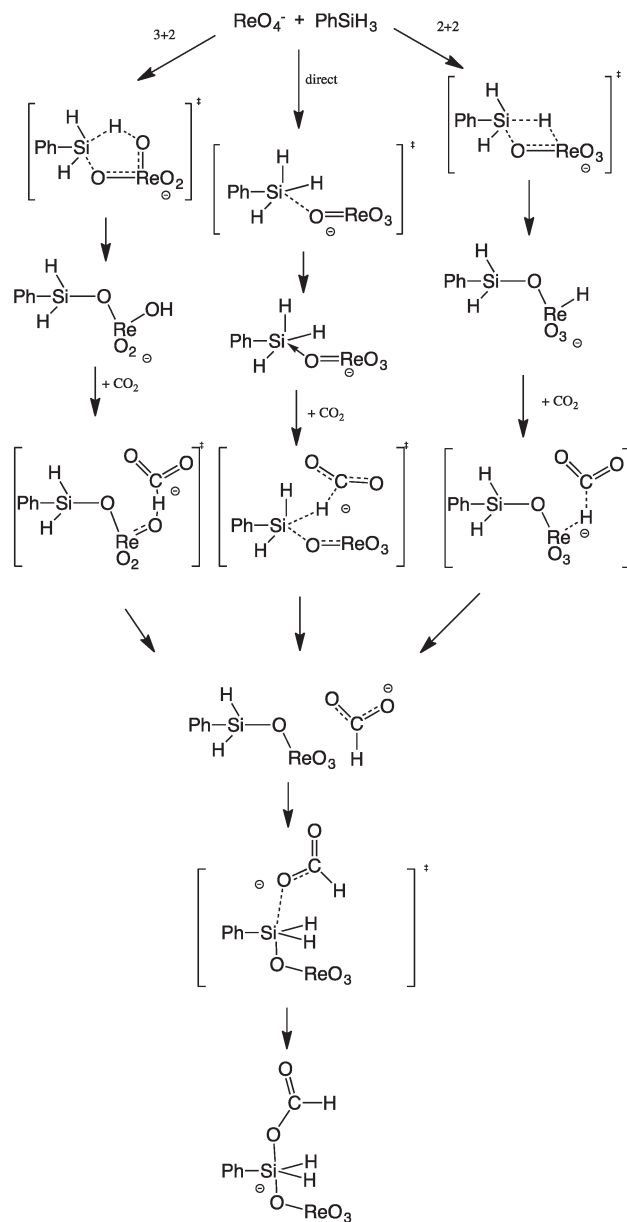
### DFT modelling of the catalytic mechanism

To explain how the carbon dioxide reacts with the hydrosilane and the perrhenate catalyst, three different principle pathways have been elucidated (Scheme 2).

Two pathways include activation of the H–Si bond of the relevant hydrosilane by either a [3 + 2] or [2 + 2] cycloaddition to the Re=O or O=Re=O fragment of the perrhenate to form a siloxyhydride, with subsequent reaction of this intermediate with  $\text{CO}_2$  by hydride migratory insertion.<sup>63,65</sup> If Si–H activation would occur to form a hydride, then the most probable mode would be [2 + 2] addition. However, the DFT results favour an alternative  $\Delta G$  energy profile in the gas phase (Fig. 1) which includes the formation of a simple complex of the hydrosilane with perrhenate without preliminary Si–H activation, forming a five-coordinate hypervalent silicon centre.<sup>2,60,67–70</sup> The  $\text{CO}_2$  activation occurs then directly at a Si–H bond to form a  $\text{HCO}_2^-$  species that adds to the silicon to form the observed silylformate (here as adduct of  $\text{ReO}_4^-$ ). This adduct can be seen as a starting point of activating the next Si–H bond by a second  $\text{CO}_2$ . In principle,  $\text{CO}_2$  can also undergo a cycloaddition to silyl perrhenate intermediates, but the barriers are higher in all pathways. Calculations incorporating a solvent continuum were carried out, with neither benzene nor DMF models significantly altering the preferred mechanism or its energies (see ESI<sup>†</sup>). Furthermore, alternative mechanisms involving the cycloaddition of  $\text{CO}_2$  to a rhenium silicate intermediate were explored, but were found to have higher energy barriers than the route described above (see ESI<sup>†</sup>).

### N-Methylation of amines using carbon dioxide

The observation that silylformates are formed during the catalytic reaction and their implication as intermediates in the N-methylation of amines using  $\text{CO}_2$  led us to assess the use of 2 to catalyse this latter reaction. Accordingly, the reaction between  $\text{CO}_2$ ,  $\text{HN}^i\text{Pr}_2$ , 4 eq. of  $\text{Ph}_2\text{SiH}_2$ , and 2.5 mol% of 2 at 80 °C in  $\text{CD}_3\text{CN}$  was monitored by  $^1\text{H}$  NMR spectroscopy. Af-



**Scheme 2** The pathways considered for the DFT investigation of the reduction of  $\text{CO}_2$  by hydrosilanes catalysed by the perrhenate salt 2.

ter 2 h, complete loss of  $\text{HN}^i\text{Pr}_2$  is observed with the concomitant formation of  $\text{MeN}^i\text{Pr}_2$  in quantitative yields (Table 2, entry 1; Fig. S35<sup>†</sup>). The excess hydrosilane is then consumed in the formation of  $[\text{Si}]\text{OCH}_3$  (5) as noted by the characteristic resonances in the  $^1\text{H}$  NMR spectrum between 3.4–3.6 ppm. The scope of the amine substrate was evaluated, revealing that several aliphatic amines are N-methylated, including isopropylamine (to form  $\text{Me}_2\text{N}^i\text{Pr}$ ), piperidine, pyrrolidine and morpholine (Table 2, entries 2–5). The N-methylation of aromatic primary or secondary amines at this temperature does not occur (Table 2, entries 6 and 7), and instead the hydrosilane is consumed in competitive  $\text{CO}_2$  reduction. However, lowering the temperature of reaction negates the further reduction of silylformate by hydrosilane to favour reaction with



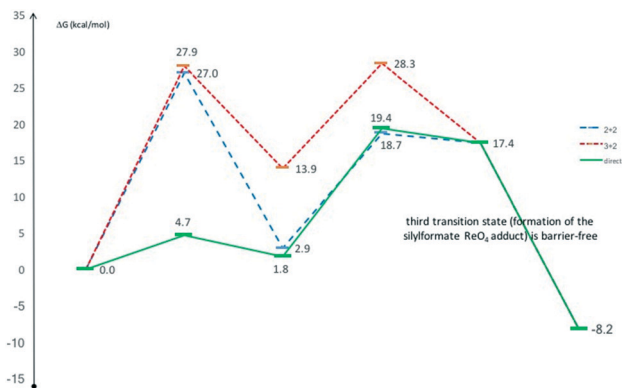


Fig. 1 Calculated Gibbs free-energy profile of the three different pathways.

the poorly nucleophilic *N*-methylaniline and, after 100 h at 20 °C, its conversion to *N,N*-dimethylaniline was seen in 40% yield. Interestingly, the major product of this reaction is the aminal  $\text{CH}_2(\text{NMePh})_2$  in 60% yield, seen by its characteristic resonances at 2.86 and 4.78 ppm in the  $^1\text{H}$  NMR spectrum (Fig. S41†). Similar aminal formation from  $\text{CO}_2$  has been seen previously in organocatalysed or Ru-catalysed reactions.<sup>47,54</sup> As the aminal can be prepared by the condensation reaction of *N*-methylaniline and formaldehyde, its formation in this catalysed reaction is likely a result of transient formaldehyde production which can arise from the decomposition of  $\text{CH}_2(\text{OSiH}_2\text{Ph})_2$ .<sup>71</sup>

It was reported recently that selective *formylation* of amines is possible using hydrosilanes and  $\text{CO}_2$  in polar solvents such as DMF and DMSO.<sup>72</sup> In contrast, it was found that the simple aliphatic amine  $\text{HN}^i\text{Pr}_2$  underwent *N*-methylation in DMF at room temperature in the absence of 2, and subsequent *in situ* monitoring of this reaction by  $^1\text{H}$  NMR spectroscopy found the initial rates essentially identical for the catalysed and uncatalysed reaction (Fig. S40†). Reactions in DMF for the other amine substrates are also selective towards *N*-formylation, which was possible in the absence of 2; however, to achieve *N*-methylation catalyst 2 was required.

### Reduction of aldehydes and ketones

Following on from the success of the  $\text{CO}_2$  reduction, attention was turned to  $\text{C}=\text{O}$  reduction for a variety of carbonyl

containing substrates.<sup>73–75</sup> An initial trial NMR reaction with benzaldehyde was carried out using the standard reaction conditions of 1.2 eq. of  $\text{PhSiH}_3$ , 0.5 mL of  $\text{CD}_3\text{CN}$  and 2.5 mol% of 2 at 80 °C. After 1 h, the  $^1\text{H}$  NMR spectrum showed near complete consumption of the aldehyde through loss of the characteristic *CHO* resonance at 10.01 ppm and formation of approximately a 50:50 mixture of  $\text{PhSi}(\text{H})(\text{OCH}_2\text{Ph})_2$  and  $\text{PhSi}(\text{OCH}_2\text{Ph})_3$ ; subsequent heating for a further 30 minutes to ensure full consumption of benzaldehyde resulted in no change to the product distribution. Simple hydrolysis through the addition of a small excess of water resulted in full conversion to the benzylalcohol product with concomitant formation of siloxanes (Fig. S42†). Following the same reaction protocol the substrate scope was expanded to other substituted arylaldehydes and alkylaldehydes (Table 3).

The arylaldehyde substrates provided efficient turnover to the corresponding benzylalcohol product upon quenching with water. Aryl substituents containing electron-withdrawing groups (Table 3, entry 2 and 5) proceed with higher TOFs compared to their electron-donating counterparts (Table 3, entry 3 and 4), in line with previous reports of rhenium catalysed hydrosilylation reactions of aldehydes.<sup>76</sup> In contrast the alkylaldehydes containing more sterically demanding substituents require twice the reaction time in comparison with the linear butyl chain derivatives.

Further extension of this perrhenate hydrosilylation catalysis examined the reduction of ketones (Table 4). Using the same reaction protocol with acetophenone, and monitoring by  $^1\text{H}$  NMR spectroscopy, the loss of the singlet resonance for the  $\text{CH}_3$  protons at 2.56 ppm is seen after 21 h at 80 °C with three new doublet resonances appearing in the range 1.49–1.37 ppm that correspond to oligomeric  $\text{PhCH}_2(\text{O}[\text{Si}])\text{CH}_3$  products; upon quenching with water these resonances resolve into a single species at 1.42 ppm ( $J_{\text{HH}} = 6.51$  Hz). The same is also true for the  $\text{CH}(\text{O}[\text{Si}])\text{CH}_3$  proton, visible around 5.1 ppm, but due to the oligomeric nature of the silylether products and the overlapping remaining  $[\text{Si}]-\text{H}$  resonances this appears as a series of multiplets which resolve into the characteristic quartet at 4.85 ppm ( $J_{\text{HH}} = 6.39$  Hz) on quenching. Expanding the scope of this ketone reduction to include substituted acetophenones follows the same trend as observed with the benzaldehyde derivatives. In this case, however, prolonged reaction times are required for the electron-donating substituents, taking 40 h for only 50% consumption

Table 2 *N*-Methylation of amines using 2 bar  $\text{CO}_2$  and  $\text{Ph}_2\text{SiH}_2$  in  $\text{CD}_3\text{CN}$ , heated at 80 °C for 2 h with 2.5 mol%  $[\text{N}(\text{hexyl})_4][\text{ReO}_4]$  (2)

	Amine	Hydrosilane (eq.)	NMR yield <sup>a</sup> (%)	TOF ( $\text{h}^{-1}$ )
1	Diisopropylamine	4	99	20
2	Isopropylamine	8	43 <sup>b</sup>	9
3	Piperidine	4	87	17
4	Pyrrolidine	4	82	16
5	Morpholine	4	85	17
6	Aniline	8	0	0
7	<i>N</i> -Methylaniline	4	0	0
8 <sup>c</sup>	<i>N</i> -Methylaniline	4	40	0.4

<sup>a</sup> NMR yield vs.  $\text{Me}_3\text{SiPh}$ . <sup>b</sup>  $\text{Me}_2\text{N}^i\text{Pr}$ . <sup>c</sup>  $\text{H}_3\text{SiPh}$ , 20 °C, 100 h.





**Table 3** Hydrosilylation of aldehydes using 1.2 eq. PhSiH<sub>3</sub> in CD<sub>3</sub>CN at 80 °C with 2.5 mol% [N(hexyl)<sub>4</sub>][ReO<sub>4</sub>] (2)

	Aldehyde	Time (h)	NMR yield <sup>a</sup> (%)	TOF <sup>b</sup> (h <sup>-1</sup> )
1	Benzaldehyde	1.5	86	23
2	<i>p</i> -Bromobenzaldehyde	1	92	37
3	<i>p</i> -Tolualdehyde	2.5	79	13
4	<i>p</i> -Anisaldehyde	2.5	89	14
5	<i>m</i> -Anisaldehyde	1.5	96	26
6	Trimethylacetaldehyde	4	91	2
7	(1 <i>R</i> )-(-)-Myrtenal	4	47	4
8	Crotonaldehyde	2.5	48	8
9	Butyraldehyde	2	46	9

<sup>a</sup> NMR ratio vs. Me<sub>3</sub>SiPh after water quench. <sup>b</sup> TOF = (conversion/catalyst loading)/time.

of the starting ketone (Table 4, entries 2 and 4 vs. 3 and 5). Reduction of benzophenone to the corresponding alcohol proceeds efficiently, and again the use of a more sterically demanding alkyl ketone requires prolonged reaction times. Interestingly, a small amount of H/D exchange was seen to take place during extended heating. Hydrogen formation occurs during the initial stages of the reaction where some of the hydrosilane is initially consumed in the reduction of trace water to form H<sub>2</sub> and siloxanes and, in the case of the acetophenone, HD is seen in the <sup>1</sup>H NMR spectrum as a characteristic 1:1:1 triplet at 4.56 ppm (*J*<sub>HD</sub> = 42.6 Hz) (Fig. S58†). Further H/D exchange was seen at the acetyl position of the acetophenone substrates and is thought to occur through keto–enol tautomerisation in the presence of the [Si]–D which results from the reaction of [Si]–H and D<sub>2</sub>O. The use of dry d-solvents negates this HD exchange.

## Experimental

### Synthesis of [N(hexyl)<sub>4</sub>][ReO<sub>4</sub>] (2)

An aqueous solution of ammonium perrhenate (1.26 g, 4.7 mmol) was added to a chloroform solution of tetrahexylammonium bromide (2.00 g, 4.6 mmol) and stirred together for 6 hours at room temperature after which the two layers were separated and the aqueous layer extracted with chloroform (2 × 10 mL). The combined chloroform extracts were dried over magnesium sulphate and the solvent removed under reduced pressure to provide a colourless solid of tetrahexylammonium perrhenate (2.10 g, 75% yield).

**Table 4** Hydrosilylation of ketones using 1.2 eq. PhSiH<sub>3</sub> in CD<sub>3</sub>CN at 80 °C with 2.5 mol% [N(hexyl)<sub>4</sub>][ReO<sub>4</sub>] (2)

	Ketone	Time (h)	NMR yield <sup>a</sup> (%)	TOF <sup>b</sup> (h <sup>-1</sup> )
1	Acetophenone	21	72	1
2	<i>p</i> -Bromoacetophenone	40	45	0.5
3	<i>p</i> -Acetyltoluene	21	66	1
4	<i>p</i> -Acetanisole	40	39	0.4
5	<i>m</i> -Acetanisole	20	81	2
6	Benzophenone	21	66	1
7	Pinacolone	37	22	0.2

<sup>a</sup> NMR ratio vs. Me<sub>3</sub>SiPh after water quench; <sup>b</sup> TOF = (conversion/catalyst loading)/time.

<sup>1</sup>H NMR (500 MHz; d<sub>3</sub>-MeCN, 300 K): δ<sub>H</sub> (ppm) 3.07 (m, 8H, NCH<sub>2</sub>), 1.60 (m, 8H, CH<sub>2</sub>), 1.33 (m, 24H, CH<sub>2</sub>), 0.91 (t, 12H, CH<sub>3</sub>).

<sup>13</sup>C{<sup>1</sup>H} NMR (500 MHz; d<sub>3</sub>-MeCN, 300 K): δ<sub>C</sub> (ppm) 59.48 (NCH<sub>2</sub>), 31.84 (CH<sub>2</sub>), 26.57 (CH<sub>2</sub>), 26.57 (CH<sub>2</sub>), 23.10 (CH<sub>2</sub>), 22.34 (CH<sub>2</sub>), 14.21 (CH<sub>3</sub>); FTIR (ATR): ν<sub>max</sub>/cm<sup>-1</sup> 902 (Re–O); ESI-MS: M<sup>+</sup> *m/z* found (calculated): 354.41068 (354.40943 C<sub>24</sub>H<sub>52</sub>N<sub>2</sub>), 959.77918 (959.75512 (C<sub>24</sub>H<sub>54</sub>N)<sub>2</sub>ReO<sub>4</sub>), 1565.16124 (1565.10132 (C<sub>24</sub>H<sub>54</sub>N)<sub>3</sub>(ReO<sub>4</sub>)<sub>2</sub>). M<sup>-</sup> *m/z* found (calculated): 250.93270 (250.93597 ReO<sub>4</sub>), 854.26749 (854.27860 (C<sub>24</sub>H<sub>52</sub>N)(ReO<sub>4</sub>)<sub>2</sub>), 1459.59185 (1459.62427 (C<sub>24</sub>H<sub>52</sub>N)<sub>2</sub>(ReO<sub>4</sub>)<sub>3</sub>), 2064.90770 (2064.97026 (C<sub>24</sub>H<sub>52</sub>N)<sub>3</sub>(ReO<sub>4</sub>)<sub>4</sub>); analysis: C<sub>24</sub>H<sub>52</sub>NO<sub>4</sub>Re found (calc.); C: 47.55% (47.65), H: 8.72% (8.67), N: 2.38% (2.32).

### General procedure for catalytic reactions of hydrosilanes with CO<sub>2</sub>

[N(hexyl)<sub>4</sub>][ReO<sub>4</sub>] (2) (3 mg, 2.5 mol%) in 0.6 mL d-solvent, hydrosilane (0.2 mmol) and trimethylphenylsilane (2 μL) as an internal standard (unless otherwise stated) were added to a Teflon-tapped NMR tube. The solution was freeze–pump–thaw degassed three times before being refilled with 1 bar of CO<sub>2</sub>. The <sup>1</sup>H NMR spectrum was recorded and then the tube was placed in a preheated oil bath (80 °C). The reactions were monitored by <sup>1</sup>H NMR spectroscopy regularly until the hydrosilane was consumed. At this point the reaction was quenched with water (10 μL, 0.56 mmol), heated for 2 h and the <sup>1</sup>H NMR spectrum recorded.

### General procedure for catalytic *N*-methylation of amines with CO<sub>2</sub>

[N(hexyl)<sub>4</sub>][ReO<sub>4</sub>] (2) (3 mg, 2.5 mol%) in 0.6 mL d-solvent, amine (0.2 mmol), silane (0.8 mmol) and trimethylphenylsilane (2 μL) as an internal standard (unless otherwise stated) were added to a Teflon-tapped NMR tube. The solution was freeze–pump–thaw degassed three times before being refilled with 2 bar of CO<sub>2</sub>. The <sup>1</sup>H NMR spectrum was recorded and then the tube was placed in a preheated oil bath (80 °C) for 2 h, after which the <sup>1</sup>H NMR spectrum was recorded.



## General procedure for catalytic hydrosilylation of aldehydes and ketones

[N(hexyl)<sub>4</sub>][ReO<sub>4</sub>]<sub>2</sub> (3 mg, 2.5 mol%) in 0.5 mL CD<sub>3</sub>CN, phenylsilane (0.24 mmol), carbonyl (0.2 mmol) and trimethylphenylsilane (2 μL) as an internal standard were added to the NMR tube. The <sup>1</sup>H NMR spectrum was recorded and then the tube was placed in a preheated oil bath (80 °C). The reactions were monitored by <sup>1</sup>H NMR spectroscopy regularly until the hydrosilane or carbonyl was consumed. At this point the reaction was quenched with water (10 μL, 0.56 mmol), heated for 15 min and the <sup>1</sup>H NMR spectrum recorded.

## Computational details

All calculations have been performed with the software Gaussian09.<sup>77</sup> The hybrid density functional B3LYP<sup>78–81</sup> has been used together with dispersion correction GD3BJ<sup>82</sup> and the triple zeta basis set 6-311++G\*\*<sup>83,84</sup> for all elements except Re. The Re atoms are described with the Stuttgart-Dresden-ECP.<sup>85</sup> The energies reported are free energies in gas-phase or for the solvents benzene and DMF with SMD solvent calculations at 298.15 K.<sup>86</sup>

## Conclusions

We have shown that the simple, lipophilic perrhenate salt [N(hexyl)<sub>4</sub>][ReO<sub>4</sub>]<sub>2</sub> acts as a catalyst for the reduction of CO<sub>2</sub> by hydrosilanes. This is the first time that perrhenate has acted as a catalyst for CO<sub>2</sub> reduction, and is likely facilitated by dissolution of the lipophilic assembly into a hydrophobic, organic solvent. The calculated mechanism shows that direct attack of CO<sub>2</sub> on a Si–H bond of a perrhenate hypervalent silicate occurs instead of alternative mechanisms through rhenium hydride formation. Furthermore, the observation that *N*-methylation of amines using CO<sub>2</sub> and organic carbonyl reduction can also be undertaken using a perrhenate catalyst expands considerably the scope of this chemistry. The ease of transfer of perrhenate from an organic phase back into an aqueous phase by the addition of base (*e.g.* NaOH) should lead to straightforward rhenium recycling, an essential step if this precious metal is to be further exploited in catalytic chemistry.

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