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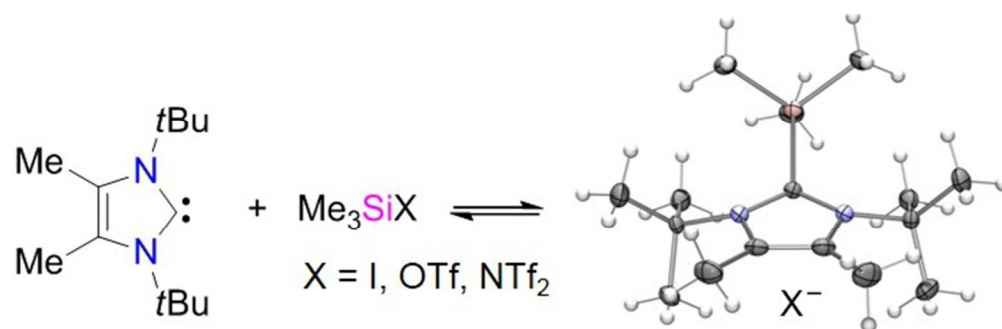


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The reaction of a sterically demanding NHC affords frustrated carbene-trimethylsilylium ion pairs, which exhibit FLP-type reactivity such as carbon dioxide fixation and metal-halide bond activation.
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ARTICLE

Frustrated N-heterocyclic carbene-silylium ion Lewis pairs

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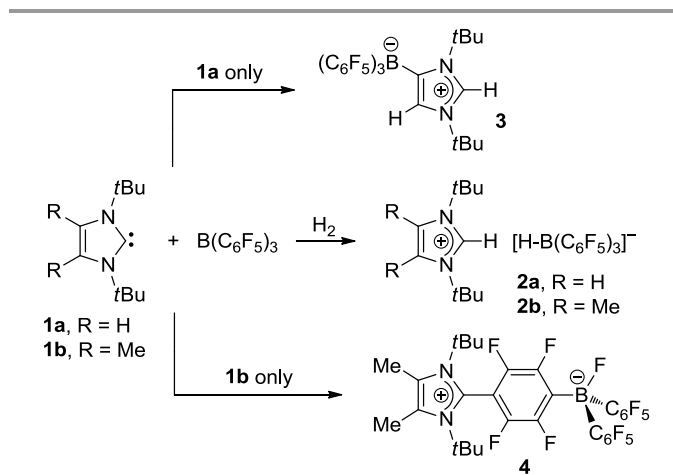
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The reaction of the N-heterocyclic carbene 1,3-di-*tert*-butyl-4,5-dimethylimidazolin-2-ylidene (**1b**) with trimethylsilyl iodide, triflate and triflimidate [Me_3SiX , X = I, CF_3SO_3 (OTf), $(\text{CF}_3\text{SO}_2)_2\text{N}$ (NTf₂)] by mixing the neat, liquid starting materials afforded the corresponding 2-(trimethylsilyl)imidazolium salts $[(1b)\text{SiMe}_3]\text{X}$ as highly reactive, white crystalline solids. Only the triflimidate (X = NTf₂) proved to be stable in solution and could be characterized by means of NMR spectroscopy (in $\text{C}_6\text{D}_5\text{Br}$) and X-ray diffraction analysis, whereas dissociation into free **1b** and Me_3SiOTf was observed for the triflate system, in agreement with the trend derived by DFT calculations; the iodide was too insoluble for characterization. The compounds $[(1b)\text{SiMe}_3]\text{X}$ showed the reactivity expected for frustrated carbene-silylium pairs, and treatment with carbon dioxide, *tert*-butyl isocyanate and diphenylbutadiene gave the 1,2-addition products $[(1b)\text{CO}_2\text{SiMe}_3]\text{X}$ (X = I, OTf, NTf₂), $[(1b)\text{C}(\text{N}t\text{Bu})\text{OSiMe}_3]\text{OTf}$ and $[(1b)\text{C}(\text{Ph})\text{C}(\text{SiMe}_3)\text{CCPh}]\text{OTf}$, respectively. Upon reaction with $[\text{AuCl}(\text{PPh}_3)]$, metal-chloride bond activation was observed, with formation of the cationic gold(I) complexes $[(1b)\text{Au}(\text{PPh}_3)]\text{X}$ (X = OTf, NTf₂).

Introduction

Frustrated Lewis pairs (FLPs), such as the well-studied phosphine-borane combination $\text{P}(t\text{Bu})_3/\text{B}(\text{C}_6\text{F}_5)_3$,^{1,2,3,4,5} represent a growing class of systems that are capable of activating small molecules, most notably dihydrogen, as a result of the unquenched and mutual reactivity of the Lewis base and acid.⁶ Accordingly, exposure of an equimolar solution of $\text{P}(t\text{Bu})_3$ and $\text{B}(\text{C}_6\text{F}_5)_3$ in toluene to an atmosphere of H_2 swiftly affords the phosphonium borate $[t\text{BuPH}][\text{HB}(\text{C}_6\text{F}_5)_3]$ under ambient conditions.^{1,2} Extensive theoretical studies^{3,4,5} of heterolytic dihydrogen splitting with this system consistently indicate the importance of secondary noncovalent interactions for the formation of an encounter complex (or prepared Lewis pair) in which the donor and acceptor sites are suitably pre-organized for synergistic interaction with an incoming H_2 molecule. A similar mechanistic proposal had been developed for the frustrated carbene-borane Lewis pair **1a**/ $\text{B}(\text{C}_6\text{F}_5)_3$,⁷ which exhibits a particularly strong propensity for heterolytic dihydrogen cleavage;^{7,8} this can be ascribed to an enhanced cumulative acid-base strength and to the strongly exergonic formation of the imidazolium salt **2a** (Scheme 1).⁹



Scheme 1 Reactivity of frustrated carbene-borane Lewis pairs.

The high reactivity of this FLP was also exploited for the activation and fixation of numerous other small molecules,¹⁰ *i.e.* ammonia,⁸ tetrahydrofuran,⁷ alkynes,¹¹ white phosphorus,¹² sulfur,¹³ carbon dioxide and nitrous oxide,¹⁴ and it was also used as a dehydrogenation reagent.¹⁵ In the absence of substrates, however, **1a**/ $\text{B}(\text{C}_6\text{F}_5)_3$ does not become isolable as a weakly bound “normal” carbene-borane adduct, but slowly undergoes self-deactivation within 2 h at room temperature to

form the significantly more stable “abnormal” adduct **3**.⁷ Although initially unwanted, this observation triggered the development of anionic N-heterocyclic carbenes with a weakly coordinating anionic moiety (WCA-NHC), which can be formally derived from **3** by deprotonation.^{16,17} During our quest to prevent such deactivation reactions, numerous FLPs were studied with variation of the N-heterocyclic carbene (NHC),^{18,19} and also of the borane component.²⁰ Thereby, carbene **1b** proved particularly useful,²¹ since the presence of methyl groups in the 4,5-positions affords a sterically even more demanding carbene ligand, and self-deactivation of **1b**/ $\text{B}(\text{C}_6\text{F}_5)_3$ is significantly slowed down in comparison with **1a**/ $\text{B}(\text{C}_6\text{F}_5)_3$. Nevertheless, formation of the zwitterionic fluoroborate **4** by C-F activation was found to proceed within 4 d at room temperature (Scheme 1).¹⁴

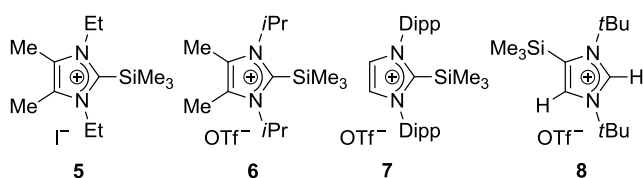


Figure 1 Reported examples of (trimethylsilyl)imidazolium salts.

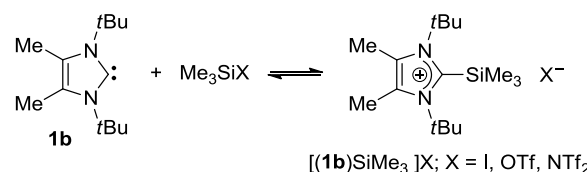
As an alternative to boranes, more strongly Lewis acidic silylium ions have also been introduced to FLP chemistry,²² and it was shown that triarylsilylium salts of the type $[\text{Ar}_3\text{Si}][\text{B}(\text{C}_6\text{F}_5)_4]$ react with bulky tertiary phosphines to afford frustrated Lewis pairs that are able to split dihydrogen irreversibly if an encounter complex with appropriate spatial requirements is formed.²³ More recently, the thermally robust phosphine-silylium adduct $[(t\text{Bu})_3\text{P}-\text{Si}(i\text{Pr})_3][\text{B}(\text{C}_6\text{F}_5)_4]$ was reported to undergo H_2 and D_2 cleavage at 90 °C with formation of the silane $(i\text{Pr})_3\text{SiH}$ and the phosphonium borate $[(t\text{Bu})_3\text{PH}][\text{B}(\text{C}_6\text{F}_5)_4]$.²⁴ In view of these results, it was an obvious choice to aim at combining NHCs with silylium ions; however, for the present study, more readily available silylium sources such as the commercially available trimethylsilyl iodide, triflate and triflimidate $[\text{Me}_3\text{SiX}, \text{X} = \text{I}, \text{CF}_3\text{SO}_3 (\text{OTf}), (\text{CF}_3\text{SO}_2)_2\text{N} (\text{NTf}_2)]$ were chosen. In previous reports, the reactions of Me_3SiI and Me_3SiOTf with sterically less encumbered NHCs furnished the stable 2-(trimethylsilyl)imidazolium salts (“normal carbene-silylium adducts”) **5**,²⁵ **6**,²⁶ and **7**,^{27,28} whereas treatment of the more bulky carbene **1a** with Me_3SiOTf resulted in rapid formation of the 4-(trimethylsilyl)imidazolium triflate **8** (“abnormal carbene-silylium adduct”, Figure 1).²⁹ The latter reaction clearly indicates that the formal addition of Me_3Si^+ to the C2 carbene atom in **1a** would afford a frustrated Lewis pair, which rearranges to the more stable abnormal adduct **8** in a similar fashion as described for **3** (*vide supra*). Therefore, the reactivity of **1b** towards Me_3SiX was investigated in the expectation that the presence of the 4,5-methyl substituents might render the corresponding carbene-silylium adducts $[(\mathbf{1b})\text{SiMe}_3]\text{X}$ ($\text{X} = \text{I}, \text{OTf}, \text{NTf}_2$), more stable and even isolable. It should be noted that these studies are also highly relevant in view of the interest in NHC-mediated

activation of tetravalent silicon compounds, since NHC-Si interactions have been hypothesized for the mechanistic course of various catalytic reactions such as cyanosilylation, CO_2 reduction with silanes or polymerisation of silicon-containing compounds.³⁰

Results and discussion

Generation of carbene-silylium adducts

Our initial efforts to isolate the carbene-silylium adducts $[(\mathbf{1b})\text{SiMe}_3]\text{X}$ ($\text{X} = \text{I}, \text{OTf}, \text{NTf}_2$) by mixing **1b** with Me_3SiX were ambiguous, and only the reaction with $\text{Me}_3\text{SiNTf}_2$ produced a precipitate of the triflimidate $[(\mathbf{1b})\text{SiMe}_3]\text{NTf}_2$ from hexane or toluene solution in satisfactory yield. Since the silyl reagents and also carbene **1b** are liquid at room temperature, equimolar amounts of **1b** and Me_3SiX were mixed by blending in a Teflon vial under argon, resulting in immediate solidification to afford $[(\mathbf{1b})\text{SiMe}_3]\text{I}$, $[(\mathbf{1b})\text{SiMe}_3]\text{OTf}$ and $[(\mathbf{1b})\text{SiMe}_3]\text{NTf}_2$ as colourless materials in quantitative yield and analytically pure form after careful washing with cold hexane in order to remove any trace of unreacted starting materials (Scheme 2). NMR spectroscopic characterisation in solution was hampered by the high reactivity of the compounds, which decompose rapidly in deuterated solvents such as CDCl_3 , CD_2Cl_2 and $\text{THF}-d_6$ and are only sparingly soluble in toluene- d_8 . However, instructive NMR spectra could be recorded for the triflate and triflimidate salts in *deutero*-bromobenzene ($\text{C}_6\text{D}_5\text{Br}$), although the samples always showed contamination with the corresponding imidazolium salt, in which a hydrogen atom replaces the Me_3Si group. The ^{13}C NMR spectra reveal distinct differences, since the spectrum of the $[(\mathbf{1b})\text{SiMe}_3]\text{OTf}$ shows a resonance at 210.5 ppm, indicating the presence of the free carbene **1b** ($\delta = 210.2$ ppm, in $\text{C}_6\text{D}_5\text{Br}$; $\delta = 212.2$ ppm in C_6D_6).²¹ Accordingly, the ^1H NMR spectrum exhibits the resonances for the free carbene at 1.55 (*t*Bu) and 2.09 ppm (Me) and also for the methyl groups of Me_3SiOTf at 0.20 ppm. The formation of only minor amounts of the expected adduct $[(\mathbf{1b})\text{SiMe}_3]\text{OTf}$ can be deduced from the observation of characteristic ^{13}C NMR signals at 155.5 and 5.4 ppm, which can be assigned to the C2 and SiCH_3 carbon atoms of the 2-(trimethylsilyl)imidazolium cation. Integration of the corresponding signals in the ^1H NMR spectrum reveals more than 90% dissociation into the starting materials.



Scheme 2. Reaction of **1b** with trimethylsilyl iodide, triflate and triflimidate; mixing liquid **1b** and liquid Me_3SiX affords quantitatively solid $[(\mathbf{1b})\text{SiMe}_3]\text{X}$.

In contrast, the NMR spectra of $[(\mathbf{1b})\text{SiMe}_3]\text{NTf}_2$ (in $\text{C}_6\text{D}_5\text{Br}$) indicate complete consumption of the free carbene **1b**, and full conversion is indicated by strong ^{13}C NMR signals at

156.0 (C2) and 5.3 ppm (SiCH₃). The ¹H NMR spectrum shows signals at 1.99, 1.35 and 0.23 ppm in a 6:18:9 ratio, attributable to the different methyl groups in [(**1b**)SiMe₃]⁺, and the ²⁹Si NMR spectrum gives rise to a signal at 4.5 ppm, which falls in the range observed for the related compounds **5** (-5.1 ppm),²⁵ **6** (4.5 ppm),²⁶ and **7** (-1.0 ppm),^{27,28b} and is significantly upfield from the chemical shift of silylating agents including Me₃SiNTf₂.³¹ It should be noted that the high reactivity of the systems **1b**/Me₃SiX has so far precluded further investigation of these putative equilibrium reactions, e.g. by temperature-dependent NMR studies.

Single crystals of [(**1b**)SiMe₃]NTf₂ suitable for X-ray diffraction analysis were obtained as a hemisolvate from C₆D₅Br solution, and the molecular structure is shown in Figure 2. Remarkably, the cation is strongly twisted and exhibits a long Si-C1 bond of 1.9671(13) Å, with the silicon atom lying 1.078(2) Å above the imidazole (C1-N1-C2-C3-N1) plane (r.m.s. deviation 0.02 Å).[#] The *t*Bu groups are forced to the opposite side of the heterocycle with deviations of 0.554(2) Å (C4) and 0.473(2) Å (C10) of the tertiary carbon atoms from the plane. A similarly distorted structure is found for the same cation with the azidodiborate counterion [(C₆F₅)₃BN₃B(C₆F₅)₃]⁻, whereas the sterically less congested compound **7** (Figure 1) shows a shorter Si-C1 bond length of 1.9330(12) Å and a smaller displacement (0.232 Å) of the Si atom, indicating a smaller degree of “frustration”.³² For comparison, significantly shorter bonds are found for unperturbed 2-silylimidazolium cations of the type [NHC-SiH₃]⁺ (1.920(2) Å)³³ and [NHC-SiI₃]⁺ (1.911(3) Å).³⁴

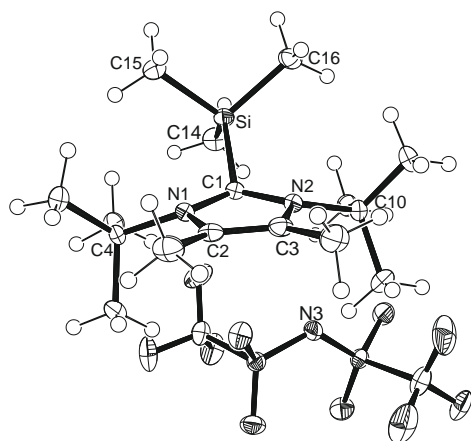


Figure 2. ORTEP diagram of [(**1b**)SiMe₃]NTf₂ with thermal ellipsoids drawn at 50% probability. Selected bond lengths (Å) and angles (°): Si-C14 1.8655(15), Si-C15 1.8650(14), Si-C16 1.8681(14), Si-C1 1.9671(13), N1-C1 1.3653(16), N1-C2 1.3949(17); C15-Si-C14 113.36(7), C15-Si-C16 101.00(6), C14-Si-C16 109.96(7), C15-Si-C1 105.10(6), N1-C1-N2 106.35(11). Solvent is omitted.

Theoretical calculations of carbene-silylium adduct formation

To rationalize the reactivity of the carbene **1b** towards the reagents Me₃SiX (X = I, OTf, NTf₂), the thermodynamics of adduct formation were calculated for the ion pairs [(**1b**)SiMe₃]X and also for the corresponding adducts [(**1c**)SiMe₃]X containing the sterically less encumbered carbene 1,3,4,5-tetramethylimidazolin-2-ylidene (**1c**). The anions were

generally placed underneath the plane of the resulting 2-(trimethylsilyl)imidazolium salt, and the structures were freely refined. We employed the functionals M05-2X, M06-2X and B97-D, which were developed by Zhao and Truhlar³⁵ and by Grimme³⁶ to describe conveniently noncovalent and long-range dispersion interactions that can be expected to contribute significantly to the overall binding, in particular for the sterically congested adducts of carbene **1b**. Table 1 summarizes the enthalpies (ΔH) and Gibbs free energies (ΔG) at 298 K for the formation of the adducts [(**1b**)SiMe₃]X and [(**1c**)SiMe₃]X, and Figure 3 shows a bar diagram for the values derived with the B97-D functional. In addition, the gas-phase association energies were calculated for the reaction of the carbenes **1b** and **1c** with a “naked” [Me₃Si]⁺ cation in the absence of any counterion (Table 1). Since the reverse reaction, the dissociation of [(**1b**)SiMe₃]⁺ and [(**1c**)SiMe₃]⁺ into the free carbenes and [Me₃Si]⁺, can be defined as the trimethylsilylium affinity (TMSA = $-\Delta H$),³⁷ high TMSA values of ca. 75 kcal mol⁻¹ (for **1b**) and ca. 92 kcal mol⁻¹ (for **1c**) can be derived for both carbenes as expected for the combination of very strong nucleophiles and electrophiles. Nevertheless, the TMSA value of **1b** is significantly smaller than that of **1c**, which confirms the lower stability (by ca. 17 kcal mol⁻¹) of the sterically overcrowded system. Notably, all calculations reproduce the highly distorted structure of the cation [(**1b**)SiMe₃]⁺ found in the solid state (Figure 2).[†]

Table 1 Enthalpies (ΔH_{298K}) and Gibbs free energies (ΔG_{298K}) in kcal mol⁻¹ for the formation of carbene-silylium adducts^a

Compound ^b	M05-2X		M06-2X		B97-D	
	ΔH	ΔG	ΔH	ΔG	ΔH	ΔG
[(1b)SiMe ₃]I	-0.2	14.6	-1.8	13.1	-5.8	9.3
[(1b)SiMe ₃]OTf	-0.3	13.8	-5.4	12.7	-9.0	6.1
[(1b)SiMe ₃]NTf ₂	-11.2	4.1	-14.0	1.6	-19.0	-3.9
[(1b)SiMe ₃] ⁺	-73.7 ^c	-56.8	-75.4 ^c	-58.3	-77.5 ^c	-60.2
[(1c)SiMe ₃]I	-22.8	-10.2	-23.4	-10.3	-26.1	-13.2
[(1c)SiMe ₃]OTf	-24.0	-11.2	-26.2	-11.0	-27.1	-13.5
[(1c)SiMe ₃]NTf ₂	-35.3	-22.4	-36.5	-22.2	-36.9	-24.7
[(1c)SiMe ₃] ⁺	-92.3 ^c	-77.2	-93.0 ^c	-77.5	-92.1 ^c	-77.1

^a M05-2X, M06-2X, and B97-D/6-311G(d,p) enthalpies and Gibbs free energies at 298 K. ^b **1b** = 1,3-di-*tert*-butyl,4,5-dimethylimidazolin-2-ylidene; **1c** = 1,3,4,5-tetramethylimidazolin-2-ylidene; if present, the anion X was placed underneath the plane of the resulting 2-trimethylimidazolium salt, and the structure was freely refined. ^c $-\Delta H$ = TMSA (trimethylsilylium affinity).³⁷

The calculations for the ion pairs [(**1b**)SiMe₃]X and [(**1c**)SiMe₃]X reveal a marked influence of the counterion X, and in particular, the triflimidates (X = NTf₂) of both systems are significantly more stable than the corresponding triflates and iodides; this reflects the effect of a highly delocalized counteranion,³⁸ but might also indicate a favourable interaction with the 2-(trimethylsilyl)imidazolium cation. Since such interactions also play an important role in the solid state, we believe that the gas-phase calculations at hand will allow us to predict or substantiate trends in reactivity. All three functionals produce qualitatively similar results, with the B97-D consistently yielding lower energies. The following discussion will be restricted to the B97-D values, which are assembled in Figure 3. For [(**1c**)SiMe₃]X, the formation of all adducts is

markedly exothermic and exergonic, which clearly suggests that frustrated Lewis pair behaviour cannot be expected for the combinations **1c**/TMSX and that complete formation of ion pairs can be expected. In contrast, the adducts [(**1b**)SiMe₃]X are significantly less stable, with their stability following the order [(**1b**)SiMe₃]I ($\Delta H = -5.8$ kcal mol⁻¹) < [(**1b**)SiMe₃]OTf ($\Delta H = -9.0$ kcal mol⁻¹) < [(**1b**)SiMe₃]NTf₂ ($\Delta H = -19.0$ kcal mol⁻¹). Although entropy effects should not be overestimated, it can be stated that the formation of the triflimidate was calculated to be slightly exergonic ($\Delta G = -3.9$ kcal mol⁻¹), whereas the triflate ($\Delta G = 6.1$ kcal mol⁻¹) and iodide ($\Delta G = 9.3$ kcal mol⁻¹) form endergonically. These theoretical results are in full agreement with the experimental observation that dissolution of the triflate in bromobenzene leads to predominant dissociation into **1b**/Me₃SiOTf, whereas the starting materials **1b**/Me₃SiNTf₂ are not observed for the triflimidate; this allowed us to characterize the latter by means of NMR spectroscopy and X-ray diffraction analysis (*vide supra*).

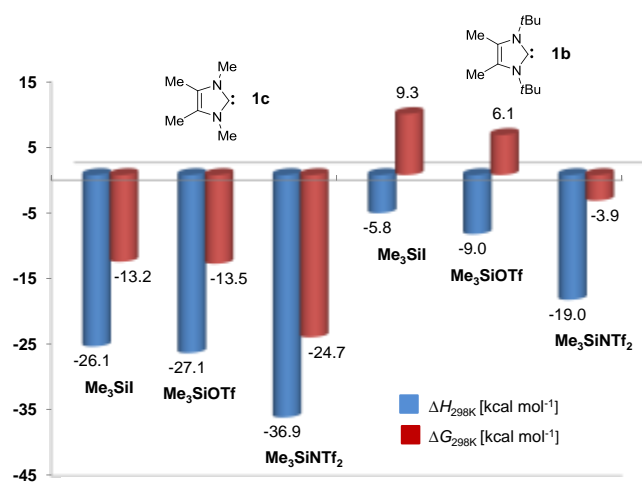


Figure 3. Bar diagram showing the enthalpies (ΔH) and Gibbs free energies at 298 K (ΔG) for the reaction of **1b** (right) and **1c** (left) with Me₃SiX (X = I, OTf, NTf₂) at the B97D/6-311G(d,p) level of theory.

To establish a potential pathway for the formation of the 2-(trimethylsilyl)imidazolium salts [(**1b**)SiMe₃]X, we assumed that the initial binding between the Lewis basic carbene **1b** and the Lewis acidic Me₃SiX reagent will occur according to the established model of n-σ* interaction,^{30,39} which leads to polarization of the adjacent Si-X bond by formation of a hypervalent silicon species. In the case of strong polarization, this may be followed by subsequent heterolytic Si-X bond cleavage and ionization. Since the same considerations are valid for the reverse reaction, we obtained suitable starting geometries based on the calculated structures of [(**1b**)SiMe₃]X (**C**), which were modified by placing the counterion X opposite to the carbene ligand. Full refinement without any geometrical constraints afforded a second set of ionic minimum structures **B** with weak Si-X interactions, which are 15.8 (X = I), 13.4 (X = OTf) and 13.0 kcal mol⁻¹ (X = NTf₂) higher in energy than the global minima. For all three systems, we were then able to

locate transition states (**TS**), which connect the ionic local minima **B** with covalent adducts of the type **1b**/Me₃SiX (**A**). The transition state structures exhibit hypercoordinate silicon atoms with distorted trigonal-bipyramidal geometries ($C_{\text{carbene}}\text{-Si-I} = 178.2^\circ$, $C_{\text{carbene}}\text{-Si-O} = 178.1^\circ$ and $C_{\text{carbene}}\text{-Si-N} = 170.5^\circ$), in which the trigonal-planar [Me₃Si]⁺ cation (C-Si-C angle sums = 360°) is flanked by loosely bound counterions and carbene ligands ($C_{\text{carbene}}\text{-Si} = 2.434$ Å for X = I, $C_{\text{carbene}}\text{-Si} = 2.401$ Å for X = OTf, $C_{\text{carbene}}\text{-Si} = 2.612$ Å for X = NTf₂). It is particularly noteworthy that very low barriers are found for the iodide and triflate systems ($\Delta\Delta E = 0.8$ kcal mol⁻¹ X = I; $\Delta\Delta E \approx 0$ kcal mol⁻¹),⁴⁰ which can be expected to rearrange readily to **1b**/Me₃SiX, whereas moderately higher barriers are found for both the formation and cleavage of [(**1b**)SiMe₃]NTf₂. The resulting adducts **A** are located -8.8 (X = I), -10.8 (X = OTf) and -9.0 kcal mol⁻¹ (X = NTf₂) below the energy of the free, isolated starting materials, but, surprisingly, they do not display any direct C-Si contacts between the carbene carbon and the silicon atoms, as indicated by very long $C_{\text{carbene}}\text{-Si}$ distances of 3.965 Å for X = I, 4.117 Å for X = OTf and 3.933 Å for X = NTf₂). Instead, the TMS groups display a series of short intermolecular C-H...C and C-H...N contacts in the range 2.53–2.78 Å, which probably contribute significantly to the overall binding energy.

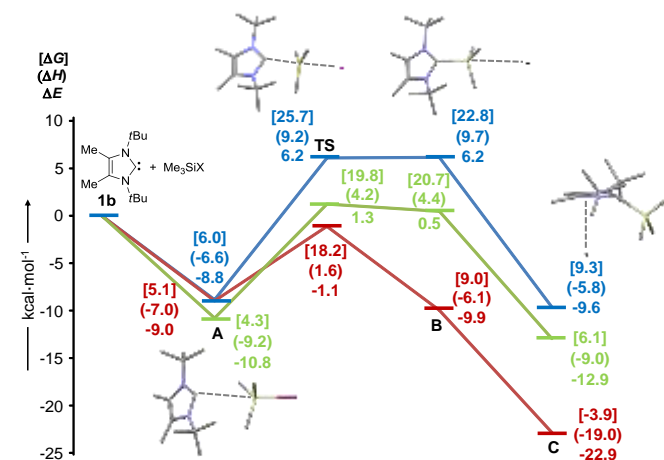


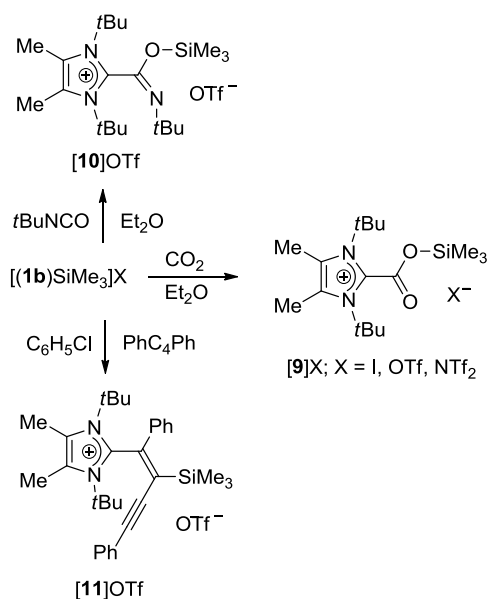
Figure 4 Potential-energy profile for the reaction of **1b** with Me₃SiX (X = I, blue; X = OTf, green; X = NTf₂, red). Values correspond to $\Delta E =$ zero-point uncorrected B97-D/6-311G(d,p) electronic energies, $\Delta H =$ enthalpies at 298 K (round brackets), and $\Delta G =$ Gibbs free energies at 298 K (square brackets). Stick models are provided for the stationary points of the **1b**/Me₃SiI system. TS = transition state.

It should be emphasized that the potential-energy profiles shown in Figure 4 can be expected to be strongly affected by solvent effects, since a transition from covalent to ionic structures is undergone, which proceeds via strongly polarized transition states. Nevertheless, we believe that the gas-phase calculations at hand allow us to rationalise the different reactivities of the **1b**/Me₃SiX systems, with the most weakly coordinating triflimidate ion affording a persistent ionic compound [(**1b**)SiMe₃]NTf₂, whereas the corresponding iodide

and triflate only form by mixing the neat starting materials, but are readily cleaved upon dissolution.

Reactivity of carbene-silylium Lewis pairs – addition reactions

The structural distortions determined experimentally and theoretically for the carbene-silylium adducts $[(\mathbf{1b})\text{SiMe}_3]\text{X}$, together with their obvious lability, prompted us to investigate their behaviour as potential frustrated Lewis pairs. However, dihydrogen cleavage with the triflate and triflimidate systems proved unsuccessful under the conditions described for isolable phosphine-silylium adduct $[(t\text{Bu})_3\text{P-Si}(i\text{Pr})_3][\text{B}(\text{C}_6\text{F}_5)_4]$.²⁴ In view of the general interest in the reduction of carbon dioxide by silanes, which can for instance be mediated by frustrated Lewis pairs,⁴¹ silylium ions,⁴² and N-heterocyclic carbenes,⁴³ carbon dioxide fixation was studied by purging diethyl ether suspensions of $[(\mathbf{1b})\text{SiMe}_3]\text{X}$ ($\text{X} = \text{I}, \text{OTf}, \text{NTf}_2$) with CO_2 . The adducts $[\mathbf{9}]\text{X}$ were isolated as voluminous white precipitates in excellent yields for $\text{X} = \text{I}, \text{OTf}$, whereas for $\text{X} = \text{NTf}_2$, the reaction was accompanied by the formation of significant amounts of imidazolium triflimidate (Scheme 3). Although slow decomposition in chlorinated solvents was observed in a similar fashion as described for carbene-borane systems,^{14,44} satisfactory NMR spectra could be recorded in CD_2Cl_2 , affording ^{13}C NMR signals for the carboxylic group at ca. 158 ppm, similar to the value for the adducts $\mathbf{1b}\cdot\text{CO}_2$ (165.5 ppm)¹⁴ and $\mathbf{1b}\cdot\text{CO}_2\cdot\text{B}(\text{C}_6\text{F}_5)_3$ (157.7 ppm)¹⁴ and shifted by approximately 33 ppm to lower field compared to free CO_2 .⁴⁵ Crystals of $[\mathbf{9}]\text{OTf}$ suitable for X-ray diffraction analysis could be obtained from THF solution at -30°C ; the resulting molecular structure is shown in Figure 5. The structural features of the CO_2 moiety, with C-O bonds of 1.2083(16) and 1.3090(15) Å, are identical within error limits to the values found for the carbene-borane adduct $\mathbf{1b}\cdot\text{CO}_2\cdot\text{B}(\text{C}_6\text{F}_5)_3$, and the same perpendicular orientation (86.5°) to the imidazole plane is also observed.¹⁴



Scheme 3. Addition reactions of $[(\mathbf{1b})\text{SiMe}_3]\text{X}$ with carbon dioxide, *tert*-butyl isocyanate and diphenylbutadiyne.

The fixation of the CO_2 congener *tert*-butyl isocyanate was also studied, and its addition to a suspension of $[(\mathbf{1b})\text{SiMe}_3]\text{OTf}$ in diethyl ether afforded the adduct $[\mathbf{10}]\text{OTf}$ as a colourless solid in 79% yield (Scheme 4). The ^1H NMR spectrum (in CD_2Cl_2) exhibits four singlets at 0.22, 1.35, 1.73 and 2.42 ppm in a 9:9:18:6 ratio, which can be assigned to the Me_3Si and the three different types of methyl groups, respectively. Single crystals of $[\mathbf{10}]\text{OTf}$ were obtained from $\text{Et}_2\text{O}/\text{CH}_2\text{Cl}_2$ solution at -30°C , and the molecular structure was established by X-ray diffraction analysis. The compound crystallizes with two independent cations and anions in the asymmetric unit with very similar structural parameters, which will be discussed for cation 1 only. The trimethylsilyl group is bound to the oxygen atom with $\text{Si-O} = 1.7103(10)$ Å, which affords a shorter C1-C14 bond length of 1.5038(18) Å and a longer C14-O1 bond length of 1.3564(16) Å in comparison to neutral NHC-isocyanate adducts.⁴⁶ The entire isocyanate moiety (Si-O1-C14-N3-C15) is planar and adopts a perpendicular orientation (89.4°) to the imidazole ring.

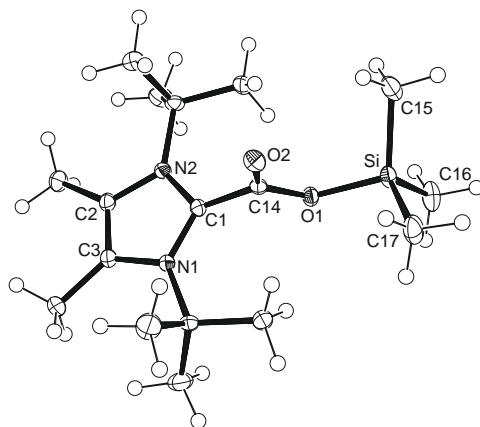


Figure 5 ORTEP diagram of cation in $[\mathbf{9}]\text{OTf}$ with thermal ellipsoids at 50% probability. Selected bond lengths (Å) and angles ($^\circ$): Si-O1 1.7399(10), Si-C16 1.8417(16), Si-C17 1.8469(15), Si-C15 1.8507(15), C1-C14 1.5129(17), N1-C1 1.3506(16), C14-O1 1.3090(15), C14-O2 1.2083(16); O1-Si-C16 101.30(6), O1-Si-C17 108.30(6), C14-O1-Si 123.60(9), N1-C1-C14 124.78(11), N1-C1-N2 109.79(11).

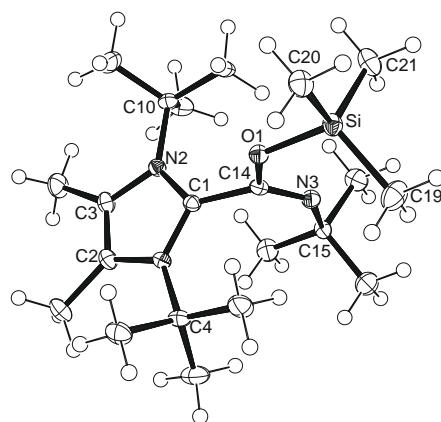


Figure 6 ORTEP diagram of one of the two independent cations in **[10]**OTf with thermal ellipsoids at 50% probability. Selected bond lengths (Å) and angles (°) in cation 1 [cation 2]: Si-O1 1.7103(10) [1.7106(10)], Si-C19 1.8444(17) [1.8486(17)], Si-C20 1.8498(15) [1.8432(18)], Si-C21 1.8509(17) [1.8484(17)], C1-C14 1.5038(18) [1.5019(18)], O1-C14 1.3564(16) [1.3577(16)], C14-N3 1.2555(18) [1.2537(18)], N3-C15 1.4897(16) [1.4873(16)]; O1-Si-C19 111.64(7) [101.71(6)], O1-Si-C20 110.01(7) [110.50(7)], C14-O1-Si 122.23(8) [120.91(8)], N1-C1-C14 124.79(12) [125.56(12)], C14-N3-C15 131.66(12) [131.82(12)].

The reaction of frustrated amine-borane and phosphine-borane Lewis pairs with conjugated diynes has been reported to proceed either by 1,2- or 1,4-addition to afford 1,3-enyne or 1,2,3-butatriene derivatives, respectively.^{47,48,49} Treatment of diphenylbutadiyne with **[(1b)SiMe₃]OTf** in chlorobenzene solution at room temperature gave the *trans*-1,2-addition product **[11]**OTf in 51% yield as a reddish brown solid (Scheme 7). The ¹H NMR spectrum shows three singlets at 0.42, 1.69 and 2.62 ppm in a 9:18:6 ratio for the methyl groups together with two sets of multiplets between 7.16 and 7.52 ppm for the phenyl rings. In the ¹³C NMR spectrum, the signals at 91.9, 106.7, 139.3 and 146.3 ppm can be assigned to a conjugated enyne moiety. 1,2-Addition of the carbene-silylium pair can be confirmed by X-ray diffraction analysis, and the molecular structure exhibits the characteristic features of an enyne framework, *i.e.* C14-C15 = 1.3649(18) Å, C15-C16 = 1.4301(19) Å, C16-C17 = 1.208(2) Å.^{48,49b} It should be noted that addition of amine-borane and phosphine-borane pairs was found to occur in a reverse manner, with the Lewis base, unlike the carbene in **[11]**OTf, attached to the inner carbon atom.

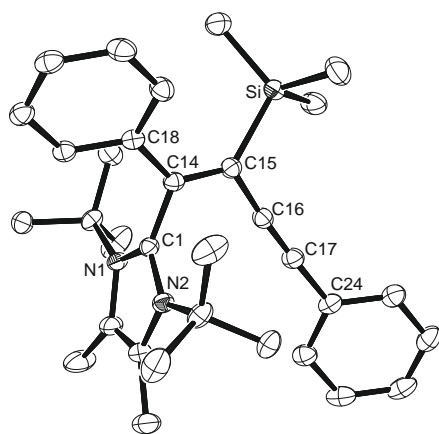
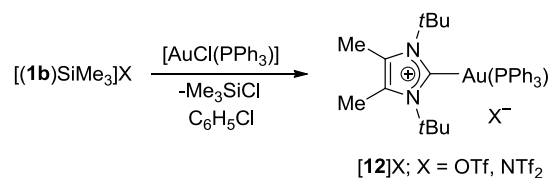


Figure 7 ORTEP diagram of the cation **11** in **[11]**OTf with thermal ellipsoids at 50% probability. Selected bond lengths [Å] and angles: Si-C15 1.9211(14), C1-C14 1.4951(17), C14-C15 1.3649(18), C14-C18 1.4783(18), C15-C16 1.4301(19), C16-C17 1.208(2), C17-C24 1.4316(19); N1-C1-N2 108.66(11), C15-C14-C1 114.89(11), C17-C16-C15 174.63(15).

Reactivity of carbene-silylium Lewis pairs – activation of metal-halide bonds

The high reactivity of the carbene-silylium pairs prompted us to employ **[(1b)SiMe₃]OTf** and **[(1b)SiMe₃]NTf₂** as metal-free carbene-transfer reagents by taking advantage of the favourable formation of volatile trimethylsilyl halides.⁵⁰ In view of the prominent role of cationic gold(I) complexes in homogeneous catalysis,⁵¹ the gold(I) chloride complex **[AuCl(PPh₃)]** was chosen as model system. Treatment of this starting material

with **[(1b)SiMe₃]OTf** and **[(1b)SiMe₃]NTf₂** in chlorobenzene solution furnished the complexes **[(1b)Au(PPh₃)]X**, **[12]X** (X = OTf, NTf₂), in high yield as colourless solids, which were isolated by precipitation with hexane and filtration (Scheme 4). NMR spectra were recorded in CD₂Cl₂ solution, revealing ³¹P NMR signals at ca. 38 ppm and ¹³C NMR signals at ca. 184 ppm with ²J_{CP} = 125, which is in agreement with the data reported for other NHC-phosphine gold(I) complexes.⁵²



Scheme 4. Synthesis of cationic gold-carbene complexes.

The molecular structures of both complexes were established by X-ray diffraction analyses, revealing almost identical structural parameters. Figure 8 shows an ORTEP diagram of the cation in **[12]NTf₂**, and further discussion will be confined to this compound. On first glance, the structural characteristics are unremarkable, with the cation displaying an almost linear C1-Au-P axes of 174.43(10)° and Au-C1 and Au-P distances of 2.063(3) and 2.2777(9) Å, which fall in the ranges found for related systems.^{16,17,52} For instance, gold-carbon and gold-phosphorus bond lengths of 2.044(4) Å and 2.275(1) Å were reported for the closely related complex **[Au(1a)(PPh₃)]PF₆**.⁵³ However, closer inspection of the structures reveals distinct differences, since the carbene ligand in the latter complex shown the expected coplanar orientation with the C-Au-P axis, whereas pronounced twisting is observed for **12**, with the gold atom being displaced by 0.753(6) Å from the imidazole plane, providing a remarkably large “pitch angle” of 20.5°. In addition, two conspicuously short Au···H contacts of ca. 2.4 Å are found, involving a methyl hydrogen atom of each *tert*-butyl group (at C6 and C13, Figure 8). These are among the shortest gold-hydrogen bonds known,⁵⁵ and their occurrence, together with the overall structural distortion, can be ascribed to hindered rotation of the *t*Bu substituents due to steric interaction with the 4,5-methyl groups (buttressing effect).^{18,21,56} It is anticipated that coordination to other metal ions such as Ru(II), Rh(I) or Ir(I) might lead C-H activation and formation of cyclometallated NHC complexes.⁵⁷

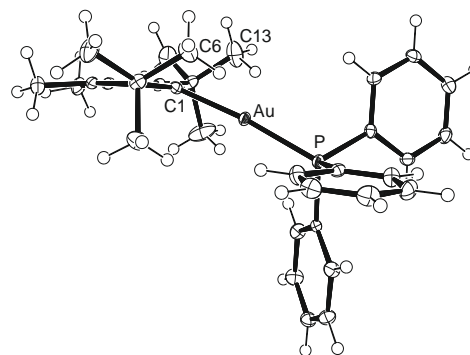


Figure 8. ORTEP diagram of the cation in **[12]**NTf₂ with thermal ellipsoids at 50% probability. Selected bond lengths [Å] and angles in **[12]**OTf/**[12]**NTf₂: Au-C1 2.063(4)/2.068(3), Au-P 2.2777(10)/2.2777(9), C1-N1 1.353(5)/1.355(4), C2-N1 1.362(5)/1.360(4); C1-Au-P 176.52(11)/174.43(10), N1-C1-N2 106.6(3)/106.5(3), N1-C1-Au 125.5(3)/124.7(2), N2-C1-Au 126.8(3)/125.2(2).

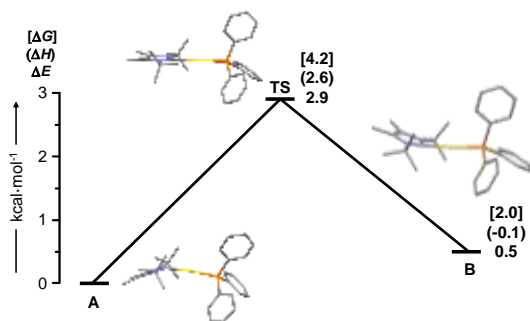


Figure 9. Potential-energy profile for the interconversion between the out-of-plane structures **A** and **B**, which proceeds via a coplanar transition state (**TS**). Values correspond to ΔE = zero-point uncorrected B97-D/6-311G(d,p) electronic energies, ΔH = enthalpies at 298 K (round brackets, and ΔG = Gibbs free energies at 298 K (square brackets).

DFT calculations were carried out to assess the energy difference between the twisted structure found in the solid state and a structure with a coplanar orientation of the imidazole plane towards the C1-Au-P axis. Therefore, the solid state structure of **12** in **[12]**NTf₂ was used as starting coordinates and fully refined to afford the minimum structure **B** with similar structural characteristics, e.g. pitch angle = 20.1° (Figure 9). Attempts to calculate an in-plane minimum structure failed; however, we were able to locate a transition state (**TS**) with a pitch angle close to zero (1.4°), which connects **B** with a second, energetically slightly favoured ($\Delta\Delta E = 0.5$ kcal mol⁻¹) minimum structure **A** (pitch angle = 22.9°). This process represents the inversion of the carbene ligand at the donor carbon atom (analogous to the known inversion process of amines), which interconverts the two twisted structures **A** and **B**, and the small barrier of 2.9 kcal mol⁻¹ indicates that the observation of this process cannot be monitored by NMR spectroscopy.

Conclusions

It was demonstrated that the stability of trimethylsilylium adducts of the sterically demanding N-heterocyclic carbene **1b** is strongly affected by the nature of the counterion and increases with decreasing nucleophilicity of the ion X or with increasing Lewis acidity of the silylating agent Me₃SiX (X = I, OTf, NTf₂).^{31,38} For all systems **1b**/Me₃SiX, the reactivity expected for frustrated Lewis pairs, such as CO₂ fixation, was observed, but it is unclear whether the reactions occur by insertion into the C-Si bond of an isolable carbene-silylium adduct or proceed by stepwise addition of Lewis acid and base, and the reaction path is again likely to depend on the counterion X. Accordingly, the term “frustrated Lewis pair” has to be used with caution, since the carbene and the counterion compete for the Lewis acid component, the trimethylsilylium ion, which cannot be expected to be freely available at any time. This

might also explain the inability of these systems to split dihydrogen. Alternatively, these carbene-silylium combinations can also be regarded as “tamed” silylium ions,⁵⁸ which have found numerous applications in Lewis acid catalysis;⁵⁹ and in view of numerous reports on NHC-mediated activation of silicon(IV) compounds,³⁰ the potential use of carbene-silylium Lewis pairs for applications in catalysis can be envisaged. As demonstrated for the preparation of cationic NHC-gold(I) complexes, the systems **1b**/Me₃SiX promote metal-halide bond activation and allow for the simultaneous introduction of carbene **1b** and a weakly coordinating counterion such as triflate or triflimidate, which might become a useful method for the preparation of coordinatively unsaturated and/or catalytically active transition metal complexes.

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[†] Electronic Supplementary Information (ESI) available: Electronic supplementary information (ESI) available: Experimental section including the synthesis and characterization of all compounds; details of the X-ray crystal structure determination and electronic structure calculations. CCDC 1052442–1052450. For ESI and crystallographic data in CIF or other electronic format, see DOI: 10.1039/c2cc32961a.

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