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## Introduction

Organosilicon species are of high interest in materials science<sup>1–5</sup> and biological research<sup>6–10</sup> and represent strategic connections for carbon–carbon or carbon–heteroatom bond formations.<sup>11–13</sup> The introduction of silyl groups in organic substrates is traditionally performed *via* stoichiometric metalation followed by the sequestering of the C-nucleophiles with  $R_3SiX$  (Fig. 1a). During the past decades, significant progress has been made in catalytic C–H bond silylations.<sup>14–16</sup> The available approaches can be classified into transition metal-catalyzed silylation (Fig. 1c), electrophilic aromatic substitution *via* silylium ions (Fig. 1b), nucleophilic substitution *via* silyl anions (Fig. 1d), silyl radical reactions (Fig. 1e), or approaches based on designed reagents as alternative silicon sources.<sup>15–17</sup> Along every individual path, the corresponding silicon reagent  $X-SiR_3$  needs to be activated by either Si–X bond heterolysis, homolysis, or hypercoordination before reacting with the C–H bond. Examples of C–H silylation leading to anionic silicates without the need for prior activation of the silicon reagent are unknown. In contrast, such chemistry is well established for Lewis acidic boranes,<sup>18–25</sup> and was recently extended to phosphonium ions.<sup>26</sup> The primary cause for the lack of reactivity of neutral silanes is their poor electrophilicity, which turns the attack of prospective nucleophilic substrates kinetically

unfeasible and the anionic silicate products thermodynamically unfavorable.

We recently launched a program on the chemistry of highly Lewis acidic silanes based on the catecholate ligand and advanced its use as a catalyst, *e.g.*, in C–O bond metathesis.<sup>27</sup>

A dominant limitation of this substance class was its poor solubility stemming from the polymeric structure formed *via* reversible Si–O dynamic covalent chemistry.<sup>28</sup> Thus, a second

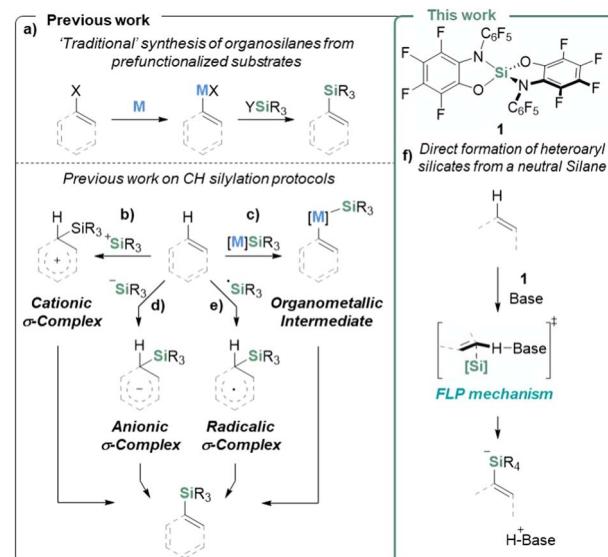


Fig. 1 (a) Previously reported conventional silane syntheses, (b)–(e) direct C–H silylation strategies,<sup>15</sup> and (f) FLP type silylation reported in this work.

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 † Electronic supplementary information (ESI) available. CCDC 2279861. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3sc03488g>



generation of highly Lewis acidic, non-oligomerizing bis(per-fluoro(*N*-phenyl-*ortho*-amidophenolato))silane **1** was developed (Fig. 1f), allowing unprecedented H<sub>2</sub> cleavage with a neutral silane.<sup>29</sup> In the present study, we employ **1** for the first C–H bond silylation reaction with a neutral silane, delivering organosilicates by the co-action of a Brønsted base. Remarkably, this process is reversible and can redeliver the C–H bond upon the addition of silaphilic agents.

## Results and discussion

Reacting *N*-methylindole with one equivalent of 1,2,2,6,6-pentamethylpiperidine (pmp) and **1** in CD<sub>2</sub>Cl<sub>2</sub>, resulted in a fast conversion at rt. NMR spectroscopy suggested clean transformation toward the *N*-methylindol-3-yl silicate [2a]<sup>–</sup> and the [pmpH]<sup>+</sup> cation (Fig. 2a). The molecular structure of the proposed ion pair was confirmed from the SCXRD analysis of the crystals grown by gas diffusion of *n*-pentane into the reaction mixture (Fig. 2b).

Encouraged by this reactivity, related *N*-methyl- and *N*-phenylpyrrole were tested and found to analogously react to the corresponding silicates under mild conditions. A similar reaction was observed for phenylacetylene, leading to silicate [2d]<sup>–</sup>. For thiophene, an activation product could not be detected at rt; heating **1**/pmp in neat thiophene to 80 °C resulted in the formation of the C–H silylated thiophene [2e]<sup>–</sup>. All products could be isolated up to 91% yields. While the C–H silylation of *N*-heterocycles<sup>30–33</sup> and acetylenes<sup>34–43</sup> that rely on the prior activation of the silicon reagents are established, we assumed a different mechanism in the present case.

Thus, the reaction was examined computationally (DLPNO-CCSD(T)/def2-TZVPP + SMD(CH<sub>2</sub>Cl<sub>2</sub>)/PBE0-D3(BJ)/def2-TZVPP, for details, see ESI†). The thermodynamic data demonstrated the stability of all products [pmpH][2x] ( $\Delta G = -52.8$  to  $-80.1$  kJ mol<sup>–1</sup>). *N*-Methylpyrrole was investigated in detail concerning possible mechanisms. The initial deprotonation of the C–H bond by pmp was assumed implausible from

the consideration of the respective pK<sub>a</sub> values (see Table S1, ESI†). Thus, a Friedel–Crafts-type process *via* the Wheland intermediate, as found with silylium ions,<sup>44</sup> element-ligand-cooperation (ELC), as per the results of bis(catecholato) phosphonium ions,<sup>26</sup> or an FLP process, in accordance with an intramolecular aminoborane,<sup>25</sup> were considered viable. First, the interaction between the substrate and Lewis acid was investigated. The distance between the silicon center and *N*-methylpyrrole in the endergonic precomplex INT-2b ( $\Delta G = 36$  kJ mol<sup>–1</sup>) is relatively large, and the pyrrole carbon do not pyramidalize – overall not indicating the formation of a Wheland intermediate, but rather a van-der-Waals complex (Fig. 3, INT-2b). Four different ELC-type transition states are possible by combining the two basic sites in the ligand of **1** (O and N) with the 2- or 3-position at the pyrrole. Barriers of  $>150$  kJ mol<sup>–1</sup> disagree with the mild reaction conditions (Fig. 3, orange path). Instead, the FLP-type cleavage in the 3-position revealed a low barrier of  $\Delta G^\ddagger = 72$  kJ mol<sup>–1</sup>, well in line with the observed reactivity at rt (Fig. 3, green path). The high selectivity is satisfactorily explained by a significantly higher barrier for the 2-position ( $\Delta G^\ddagger = 94$  kJ mol<sup>–1</sup>, Fig. 3, blue path).

Based on these insights, we concluded that the FLP mechanism is operative. For *N*-methylindole, an even lower barrier for bond cleavage in 3-position was revealed ( $\Delta G^\ddagger = 60$  kJ mol<sup>–1</sup>), whereas C–H silylation at C2 was found to be less favored ( $\Delta G^\ddagger = 120$  kJ mol<sup>–1</sup>). The C2 selectivity in thiophene is in line with a barrier of  $\Delta G^\ddagger = 98$  kJ mol<sup>–1</sup>, and a significant barrier of  $\Delta G^\ddagger = 122$  kJ mol<sup>–1</sup> at C3. Given the simultaneous C–H deprotonation in the transition state, it is also interesting to compare the pK<sub>a</sub> values of the C2–H bonds in this context: thiophene = 33.0, *N*-methylindole = 38.1, and *N*-methylpyrrole = 39.5.<sup>45</sup> Since the pK<sub>a</sub> values do not correlate with the computed barriers, it can be concluded that it is not the C–H bond acidity, but the donor-interaction with the silane that determines the barrier.

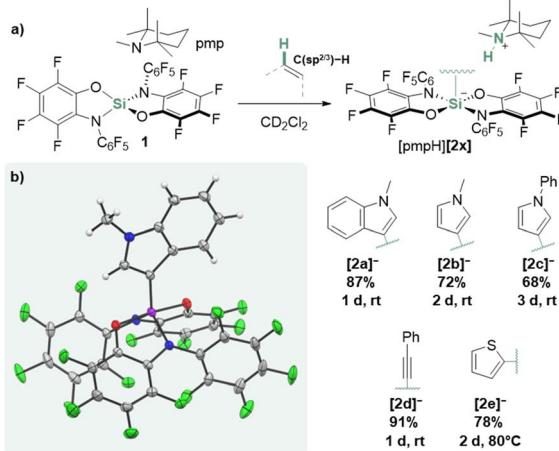


Fig. 2 (a) General synthesis of silicates [pmpH][2x] from **1** and (b) SCXRD-derived structure of [2a]<sup>–</sup> ([pmpH]<sup>+</sup> cation omitted for clarity, ellipsoids shown at 50% probability).

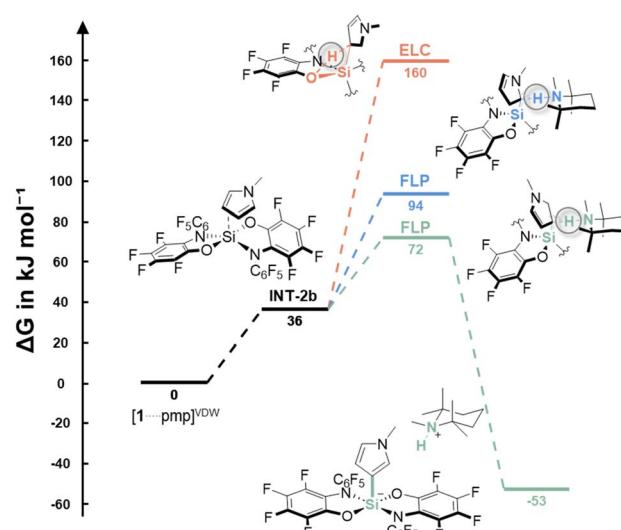


Fig. 3 Free-energy profile for the C–H silylation of *N*-methylpyrrole with **1** and pmp (bottom, DLPNO-CCSD(T)/def2-TZVPP + SMD(CH<sub>2</sub>Cl<sub>2</sub>)/PBE0-D3(BJ)/def2-TZVPP). For details and the energies of other ELC transition states, see ESI†.



Accordingly, for the even weaker donor benzene, a calculated barrier of  $\Delta G^\ddagger = 154 \text{ kJ mol}^{-1}$  explains the absence of an activation product even at elevated temperatures.

To corroborate the C–H bond cleavage as the rate-determining step, we measured the kinetic isotope effect (KIE). Equimolar amounts of the respective 3-proteo- and 3-deutero- *N*-methylindole were dissolved in  $\text{CD}_2\text{Cl}_2$  and reacted with 0.5 eq. of pmp and **1**. The quantification of the H/D ratios before and after the reaction revealed a significantly faster reaction for the proteo derivative (KIE = 3.5, for details, see ESI†). This value is in line with an FLP-type C–H bond cleavage, as observed previously by Fontaine *et al.*<sup>25</sup> Another experimental indication for the FLP-type mechanism was obtained by using 2,6-di-*tert*-butyl-pyridine as the base, in which case, no reaction occurred. The reactivity of FLPs is decisively influenced by weak association through secondary interactions,<sup>46,47</sup> for which the choice of the base can be crucial.<sup>18</sup> In a Friedel–Crafts or ELC mechanism, the base would not be involved in the rate-determining step and less sensitivity on their nature would be expected.

Despite considerable recent progress with group 14 FLPs,<sup>48–58</sup> the cleavage of C–H bonds states a novum. Thus, we were interested in how other silanes would perform in such reactivity. We computed the FLP mechanism for  $\text{SiCl}_4$  as ‘archetypal’ silicon Lewis acid with *N*-methylpyrrole as a substrate. A barrier of  $\Delta G^\ddagger = 181 \text{ kJ mol}^{-1}$  renders the reaction very unlikely. Accordingly, a control experiment with  $\text{SiCl}_4$ /pmp did not result in any observable reaction after 7 d at rt or 2 d at 60 °C. To account for a stronger silicon Lewis acid, we computed the process with  $\text{Si}(\text{cat}^{\text{Cl}})_2$ ,<sup>72</sup> which revealed a sufficiently low calculated barrier ( $\Delta G^\ddagger = 63 \text{ kJ mol}^{-1}$ ). Indeed, similar reactivity could be observed for the FLP  $[\text{Si}(\text{cat}^{\text{Cl}})_2]_n/\text{pmp}$ , but with substantially prolonged reaction times (56% conversion after 5 d), compared to **1**/pmp (78% after 16 h). This discrepancy arises from the polymeric nature of  $[\text{Si}(\text{cat}^{\text{Cl}})_2]_n$ , which causes extremely poor solubility in non-donor solvents.<sup>59</sup> While the monomeric, Lewis superacidic  $\text{Si}(\text{cat}^{\text{Cl}})_2$  is easily formed with donor-substrates (*e.g.*, ethers, amines), the cleavage of C–H bonds is favorable only with **1**, which is monomeric and soluble right away.

Next, we probed the reactivity of the ammonium silicate products. Interestingly, when  $[\text{pmpH}][\mathbf{2a}]$  was handled in strong donor solvents such as DMSO, we observed partial reformation of *N*-methylindole. Aiming for a complete reconstruction of the C–H bond,  $[\text{pmpH}][\mathbf{2a}]$  was dissolved in  $\text{CD}_2\text{Cl}_2$ , and one equivalent of 1,3-dimethylimidazolidin-2-one (DMI) was added. Heating the mixture to 60 °C resulted in the formation (>90%) of *N*-methylindole and pmp, along with the DMI adduct of **1** (Fig. 4a). Experiments to transfer a benzyl group, instead of a proton, to the silylated C3 in  $[\mathbf{2a}]^-$  using trimethylbenzylammonium chloride were unsuccessful, but again led to *N*-methylindole.

To probe the potential of the C–H bond silylation for subsequent electrophilic functionalization, the reaction of  $[\text{pmpH}][\mathbf{2a}]$  with acrylonitrile was tested (Fig. 4b).<sup>60,61</sup> The formation of **3a** could be observed, but the process occurred only slowly and incompletely at 60 °C (38%, details see ESI†).

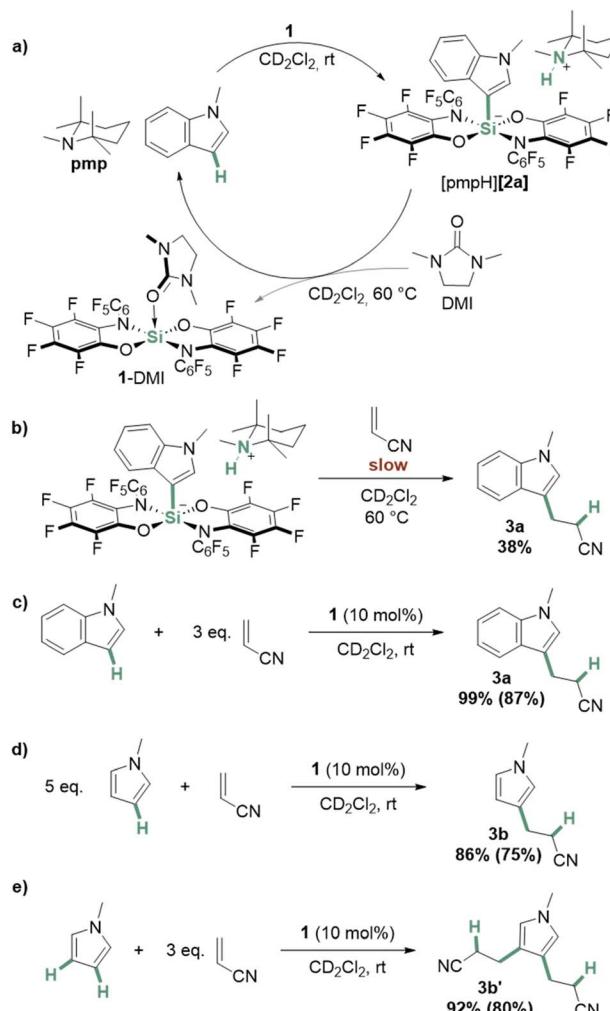


Fig. 4 Schematic depiction of (a) the synthesis of  $[\text{pmpH}][\mathbf{2a}]$  from *N*-methylindole and the reverse reaction triggered by DMI, (b) the reaction of acrylonitrile and  $[\text{pmpH}][\mathbf{2a}]$ , the addition of (c) *N*-methylindole and (d)/(e) *N*-methylpyrrole to acrylonitrile catalyzed by **1**. Yields were determined against cyclooctane as an internal standard, yields in parentheses are isolated yields (details see ESI†).

Based on the computed energies (Fig. 3), we attributed the poor reactivity of  $[\text{pmpH}][\mathbf{2a}]$  to its high thermodynamic stability. Thus, we considered the absence of a base to avoid this energetic sink. Indeed, the reaction between *N*-methylindole and acrylonitrile proceeded at rt to quantitative conversion using 10 mol% of **1** (Fig. 4c).

To our delight, we found **1** also to catalyze the one- and two-fold addition of *N*-methylpyrrole to acrylonitrile to yield **3b** and **3b'** at room temperature (Fig. 4d and e, respectively). Of course, in the absence of a base, the Lewis acid activation of acrylonitrile by **1** is more likely than the involvement of a C–H silylated intermediate. Thus, the formation of  $[\text{pmpH}][\mathbf{2a}]$  would represent a ‘bond activation’ that corresponds to an overall substrate deactivation – a situation always to be suspected if one aims for catalysis with highly reactive reagents. Instead, the observed reactivity ultimately exploits the ability of this class of Lewis acids to bind even weakly donating nitriles<sup>62–66</sup> – and

unlocks a substrate class that is challenging to activate with less reactive catalysts.

Indeed, while the Lewis acid-mediated addition of *N*-heteroarenes to Michael systems has been investigated for nitro-olefins or acrolein derivatives,<sup>67–70</sup> only a few examples have been reported for catalytic additions to acrylonitrile. For the synthesis of **3a** from *N*-methylindole, we only found a gold(III) catalyzed protocol resulting in a 52% yield.<sup>71</sup> For less nucleophilic pyrroles, such reactivity is even more sparsely reported. Using osmium catalysis, **3b** could be synthesized in 40% yield.<sup>72</sup> Related protocols for the synthesis of **3b'** were, to the best of our knowledge, not reported.

## Conclusions

A C–H bond cleavage using a neutral Lewis superacidic silane **1** and the amine pmp is introduced as a valuable extension to existing C–H silylation protocols. The process occurs *via* an FLP-type mechanism, which represents a novum for silicon Lewis acid–base pairs. Lewis acid **1** offers superior properties over other neutral silicon Lewis acids due to solubility-enhanced kinetic advantages, improving the young class of Lewis superacidic silanes. The bond cleavage at *N*-methylindole can be reversed using a donor, such as DMI. Because carbon-substituted bis(catecholato)silicates have proven to be versatile for generating C–radicals,<sup>73–78</sup> or as transmetalation reagents in coupling reactions,<sup>79–81</sup> strategic combinations with the herein-discovered C–H silylation can be considered. However, this study also emphasizes that an apparent “bond activation” does not necessarily correspond to “substrate activation”, as exemplified in the electrophilic alkylation of *N*-methylindole.

## Data availability

Cartesian coordinates of the computed structures and characterization data of the described compounds are available in the ESI.† Crystallographic data have been deposited at the Cambridge Crystallographic Data Centre (CCDC: 2279861).

## Author contributions

L. G. and T. T. devised the project and designed the experiments. T. T. carried out the calculations, performed the experimental work, and analyzed the data. L. G. and T. T. wrote the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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