Consecutive \( \beta,\beta' \)-Selective \( \text{C}(\text{sp}^3) - \text{H} \) Silylation of Tertiary Amines with Dihydrosilanes Catalyzed by \( \text{B}(\text{C}_6\text{F}_5)_3 \)

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Dedicated to Professor Siegfried Blechert on the occasion of his 75th birthday

Abstract: Tris(pentafluorophenyl)borane has been found to catalyze the two-fold \( \text{C}(\text{sp}^3) - \text{H} \) silylation of various trialkylamine derivatives with dihydrosilanes, furnishing the corresponding 4-silapiperidines in decent yields. The multi-step reaction cascade involves amine-to-enamine dehydrogenation at two alkyl residues and two electrophilic silylation reactions of those enamines, one inter- and one intramolecular.

Selective functionalization of \( \text{C}(\text{sp}^3) - \text{H} \) bonds is an important goal in synthetic chemistry.\([1]\) One way to achieve this is by transition-metal-catalyzed \( \text{C}(\text{sp}^3) - \text{H} \) silylation,\([2,3]\) and recently selected boron Lewis acids also emerged as catalysts for this purpose.\([4]\) For example, \( \text{B}(\text{C}_6\text{F}_5)_3 \) has been shown to abstract hydride from \( \alpha\)-\( \text{C}(\text{sp}^3) - \text{H} \) bonds of amines to result in the formation of iminium ions and the borohydride;\([5]\) that iminium ion is \( \text{C} - \text{H} \) acidic and can be deprotonated by another molecule of the amine, affording the corresponding enamine along with the ammonium borohydride,\([6,7]\) (Scheme 1, gray box). The net reaction is a dehydrogenation that enables subsequent bond formation with electrophiles in the \( \beta \)-position to the nitrogen atom, thereby representing a formal activation of the \( \beta\)-\( \text{C}(\text{sp}^3) - \text{H} \) bond. This process has already been employed for silylation,\([8]\) alkylation,\([9]\) deuteration,\([10]\) and olefination\([11,12]\) of the \( \beta \)-carbon atom of various (a)cyclic tertiary amines (Scheme 1, top). Of note, Park and Chang merged the \( \text{C}(\text{sp}^3) - \text{H} \) silylation with a \( \text{B}(\text{C}_6\text{F}_5)_3 \)-catalyzed intramolecular Friedel–Crafts-type silylation\([12]\) for the synthesis of bridged silicon-containing nitrogen heterocycles starting from N-arylated piperidines.\([10,11]\) However, the undirected silylation of acyclic tertiary amines\([13] \) as well as their challenging two-fold \( \text{C}(\text{sp}^3) - \text{H} \) silylation are unprecedented. We disclose here a \( \beta,\beta' \)-selective \( \text{C}(\text{sp}^3) - \text{H} \) silylation of acyclic tertiary amines and dihydrosilanes catalyzed by \( \text{B}(\text{C}_6\text{F}_5)_3 \) to directly arrive at sila analogues of piperidines (Scheme 1, bottom left). These are valuable building blocks in medicinal chemistry,\([13]\) for example, for the dopamine receptor antagonist sila-haloperidol (Scheme 1, bottom right).\([14]\) Different from our approach, established syntheses typically start from divinyl-substituted silanes employing a sequence of hydrobromination or hydroboration–oxidation–sulfenylation followed by dialkylation of a primary amine.\([15]\)

We began our investigation with optimizing the two-fold \( \text{C}(\text{sp}^3) - \text{H} \) silylation of benzylidethylamine (1a–3aa; Table 1). Treatment of 1a and PhSiH\(_2 \) (2a, 2.0 equiv) with 20 mol % of \( \text{B}(\text{C}_6\text{F}_5)_3 \) in \( \text{p-xylene} \) at 150 °C afforded 3aa after 15 h in 56% yield (Table 1, entry 1). Previous reports had indicated that the use of a metal oxide\([16,17]\) or a silyl triflate\([18,19]\) as an additive could improve the reactivity.\([16,18]\) However, substoichiometric amounts of CaO or SrO decreased the yield (Table 1, entries 2 and 3). The addition of 40 mol % of a silyl triflate improved the reactivity (Table 1, entries 4–6), and a 75% yield of 3aa was obtained with \( \text{Me}_3\text{SiOTf} \) as the additive. That yield was somewhat lower when using less and more \( \text{Me}_3\text{SiOTf} \), respectively (Table 1, entries 7 and 8). The reaction was completed within 2 h while a further shortened
Table 1: Selected examples of the optimization of B(CF3)3-catalyzed two- 
fold C(sp3)–H silylation.†[a]

| Entry | Additive (mol%) | Solvent | t [h] | Yield [%][b] |
|-------|----------------|---------|------|-------------|
| 1     | –              | p-xylene | 15   | 56          |
| 2     | CaO (50)       | p-xylene | 15   | 48          |
| 3     | SrO (50)       | p-xylene | 15   | 50          |
| 4     | Me3SiOTf (40)  | p-xylene | 15   | 75          |
| 5     | iPrMe3SiOTf (40)| p-xylene | 15   | 66          |
| 6     | Me3SiOTf (20)  | p-xylene | 15   | 67          |
| 7     | Me3SiOTf (80)  | p-xylene | 15   | 60          |
| 8     | Me3SiOTf (40)  | p-xylene | 2     | 75 (73)    |
| 9     | Me3SiOTf (40)  | p-xylene | 1     | 42          |
| 10    | Me3SiOTf (40)  | toluene  | 2     | 74          |
| 11    | Me3SiOTf (40)  | benzene  | 2     | 62          |
| 12    | Me3SiOTf (40)  | CH2Cl2   | 2     | 55          |
| 13    | Me3SiOTf (40)  | p-xylene | 2     | 0           |
| 14[b] | Me3SiOTf (40)  | p-xylene | 2     | 49          |
| 15[h] | Me3SiOTf (40)  | p-xylene | 2     | 68          |
| 16[h] | Me3SiOTf (40)  | p-xylene | 1     | 61          |
| 17[i] | Me3SiOTf (40)  | p-xylene | 2     | 60          |
| 18[i] | Me3SiOTf (40)  | p-xylene | 2     | 34          |
| 19[i] | Me3SiOTf (40)  | p-xylene | 2     | 65          |
| 20[i] | Me3SiOTf (40)  | p-xylene | 2     | 65          |

[a] All reactions were performed on a 0.050 mmol scale in a 10 mL sealed tube. [b] Yields determined by 1H NMR spectroscopy using mesitylene as an internal standard; isolated yields in parentheses. [c] Without B(CF3)3. [d] 10 mol% B(CF3)3 used. [e] 1.5 equiv Ph2SiH2 (2a). [f] Run at 120°C. [g] 5.0 mL sealed tube used. [h] 1.0 mL sealed tube used. [i] Open system with a continuous flow of nitrogen gas. [j] 5.0 mmol scale.

reaction time to 1 h resulted in a lower yield (Table 1, entries 9 and 10). Other arene solvents were tested but none provided a better outcome (Table 1, entries 11–13). A control experiment verified that Me3SiOTf is unable to mediate the reaction in the absence of B(CF3)3 (Table 1, entry 14). Less B(CF3)3 or Ph3SiH2 (2a) as well as lowering the temperature to 120°C led to a decreased reactivity (Table 1, entries 15–17). The volume of the reaction vessel was also examined, and the results indicate that vessels smaller than 10 mL are detrimental (Table 1, entries 18 and 19). We ascribe this to catalyst inhibition by dihydrogen at high pressure.†[7] A good yield was restored on a 5.0 mmol scale when performing the two-fold C(sp3)–H silylation of substrate 1o in an open system with a continuous flow of nitrogen gas (Table 1, entry 20).

We continued exploring the scope under the optimized reaction setup (Scheme 2; cf. Table 1, entry 9). It must be noted that reductive C(sp3)–N bond cleavage†[27] is competing in any of the reactions summarized in Scheme 2, and secondary amines are the major byproducts (not quantified because of their volatility). N-Benzylated diethylamine derivatives bearing various electron-donating or-withdrawing substituents on the aryl moiety reacted with Ph3SiH2 (2a) to furnish the corresponding 4-silapiperidines in moderate to good yields (1b–l → 3ba–la; gray box). All halo groups (1g–j) and a trifluoromethyl group (1k) were compatible. Tertiary amine 1l containing a methyl ether underwent demethylation/silylation, and the free phenol was isolated in 50% yield after purification by flash chromatography on silica gel (1l → 3la). A lower yield was obtained for a naphth-2-ylmethyl instead of the benzyl group (1m → 3ma). The bis(4-silapiperidine) 3na was formed in 47% yield by four-fold C(sp3)–H silylation of 1n. Replacing the benzyl group by an alkyl group was feasible (1o–q → 3oa–qa). Notably, the two-fold C(sp3)–H silylation of substrate 1o bearing two ethyl groups and one cyclohexyl group proceeded chemoselectively at the ethyl groups to form 3oa. Substituted 4-silapiperidine derivatives were obtained from tertiary amines with groups other than ethyl (1r–t → 3ra–ta). As expected, 1t gave 3ta with essentially no diastereoselectivity (cis/trans = 58:42). Attempted but failed cyclizations included tertiary benzylamines as precursors having two isopropyl, cyclohexyl, isobutyl, or phenethyl groups as well as 1-benzylazepane (see the Supporting Information for details). We also tried the silylation of the tertiary aniline derivative 1u which did not react under Park’s and Chang’s catalytic system (Scheme 3).†[34] Bicyclic 4ua and tricyclic 5ua formed in yields of 50% and 8%, respectively. The proportion of 5ua increased at longer reactions times, for example, 44% yield of 4ua and 25% yield of 5ua after 3 h. As for the aforementioned method†[15] intramolecular Friedel–Crafts C(sp3)–H silylation†[28] is favored over intramolecular C(sp3)–H silylation.
We next assessed the dihydrosilane scope in the reaction of model substrate 1a (Table 2). Diarylsilanes 2b–e exhibited good reactivity, furnishing the corresponding products in the same yield range as compared to 2a (1a → 3ab–ae; Table 2, entries 1–4). No reaction was seen with sterically hindered dimethylsilane (2f; Table 2, entry 5). Modest yield was obtained with MePhSiH₂ (2g in 1a → 3ag; Table 2, entry 6) but the synthesis of a spirocyclic derivative with 1-silaindan-1,1-diyl traces (3a i) in moderate yield (Table 2, entry 8), but again, there was no reaction with bulky Bu₂SiH₂ (2j; Table 2, entry 9). The reaction of the primary hydrosilane PhSiH₂ yielded only trace amounts of the 4-silapiperidine (not shown).

The benzyl group in 4-silapiperidines such as 3aa serves as a linchpin for further manipulations (Scheme 4). Debenzylation was achieved by treatment with 1-chloroethyl chloroformate followed by the reaction of the resulting carbamate with MeOH (3aa → 6). The benzyl group can also be converted into a benzoyl group by oxidation with KMnO₄ in the presence of BnNEt₃Cl (3aa → 7).

To gain insight into the reaction mechanism of this two-fold C(sp³)–H silylation, deuterium-labeling experiments and stoichiometric experiments were performed (Scheme 5). The reaction of 1a with Ph₂SiOTf (2a–d₃) under standard conditions gave 3aa–d₃ in the expected yield with 41% deuterium incorporation in the benzylic position as well as at the α carbon atoms (Scheme 5, top). This result confirms the known reversible hydride abstraction from C(sp³)–H bonds α to an amine nitrogen atom. Importantly, 6% deuterium incorporation was also detected for the β carbon atoms, which is evidence for hydrogenation of the enamine intermediate. In the case of diethyl-substituted 1a, silylation is faster than this backward reaction. Conversely, di-n-butyl-substituted 1v shows a different outcome (Scheme 5, top). None of the hypothetical 4-silapiperidine 3aa–d₃ was found (not shown) but instead 1v–d₃ with the usual deuteration in the α-positions. However, the deuteration grade in the β-positions was 24%, demonstrating that enamine hydrogenation is now a competitive if not the only reaction pathway for more hindered alkyl chains. To inspect the influence of the Me₂SiOTf additive, we mixed 1a and B(C₆F₅)₃ in an equimolar ratio (Scheme 5, bottom). This known reaction led to the formation of the three boron species 8a–10a in 52%, 30%, and 17% yield, respectively, and this product distribution was not affected by the addition of 1.0 equiv of Me₂SiOTf.

On the basis of the above experimental results and the literature precedent as well as DFT calculations by Park and Dang, a plausible reaction mechanism is proposed (Scheme 6). B(C₆F₅)₃ promotes hydride abstraction from the tertiary amine 1a to generate the iminium borohydrides 11a.
and 11a' in equilibrium. Their subsequent deprotonation by unreacted 1a yields enamine 12a and FLP-type dihydrogen adduct 8a; these can regenerate the free amine 1a and the catalyst B(C₆F₅)₃ along with release of dihydrogen.[7,19] The thus-formed enamine 12a then engages in the rate-determining B(C₆F₅)₃-catalyzed intermolecular hydrosilylation through the Piers mechanism[20] with 12a as a carbon nucleophile (B(C₆F₅)₃)–13a=15aa). Alternatively, B(C₆F₅)₃-activated hydrosilane 13a can also react with the amine nitrogen nucleophile 1a to equilibrate with silylaminium borohydride 14aa, the resting species of the overall process.[8a,b] Initially formed 15aa stands in equilibrium with regioisomeric 15a'aa and 15a''aa, and 15a''aa can undergo another deprotonation affording enamine 16aa. That enamine again enters the catalytic cycle of the B(C₆F₅)₃-promoted, now intramolecular hydrosilylation to eventually arrive at the title compound 3aa.

In summary, we have developed a B(C₆F₅)₃-catalyzed two-fold β,β-selective (formal) C(sp³)–H silylation of acyclic tertiary amines with dihydrosilanes to construct 4-silapiperidines and its derivatives. The reaction involves two amine-to-enamine dehydrogenation reactions each followed by an inter- and an intramolecular electrophilic enamine silylation, respectively.

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Conflict of interest

The authors declare no conflict of interest.

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