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## Direct formation of 4,5-disubstituted carbazoles via regioselective dilithiation†

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Carbazoles are widely exploited for their interesting photophysical and electronic properties, however bay (4,5-) functionalization is challenging, and previously inaccessible through carbazole C–H activation. We report a simple methodology which introduces a range of versatile 4,5-functionality, enabling the wider investigation of ring annulation and close proximity effects on carbazole properties.

Carbazoles represent a privileged structure within chemistry: their advantageous photophysical and electronic properties have led to multiple applications in organic materials,<sup>1</sup> sensing,<sup>2</sup> and as organic photocatalysts.<sup>3</sup> Carbazoles are also found in a large number of naturally-occurring alkaloids<sup>4</sup> and other biologically active molecules,<sup>5</sup> and hence the development of methods for their preparation remains an important area of research.<sup>6</sup>

Groups at the 4- and 5-positions, or bay region, of carbazole are in close proximity, but systematic investigations into such *peri*-like interactions are limited by the lack of effective preparatory methods and general synthetic building blocks. Thus the relatively few examples of 4,5-disubstituted carbazoles in the literature are prepared from non-carbazole precursors through benzannulation of indoles or pyrroles,<sup>6b</sup> or through formation of the central nitrogen heterocycle from biaryls or diarylamines.<sup>6c</sup> The influence of close proximity, *peri*-like interactions on the structure, reactivity and electronic properties of 4,5-disubstituted carbazoles has been noted in a few cases,<sup>7,8</sup> and carbazoles with additional ring fusion at the 4,5-positions have been investigated in a variety of contexts, but again not synthesized by annulation of carbazole.<sup>9</sup>

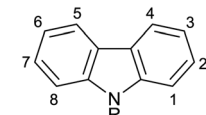
Aromatic C–H functionalization represents the most direct method for the preparation of substituted carbazoles, however the ease of C-substitution is highly positional dependent

(Scheme 1). Functionalization at the 3,6-positions (through electrophilic aromatic substitution<sup>10</sup>) and at the 1,8-positions (through directed lithiation<sup>11</sup> or metal-catalysed C–H activation<sup>12</sup>) are well established. Other substitution patterns are more difficult to access: 2,7-functionalization is rare,<sup>13</sup> and there is only one method for C–H activation at C-4, through the coupling of *N*-pyrimidinyl carbazoles with tertiary  $\alpha$ -bromoesters under Ru catalysis.<sup>14</sup>

The combination of *n*BuLi and *N,N,N',N'*-tetramethylethylenediamine (TMEDA) can be used to regioselectively dilithiate a select range of aromatic systems through directed deprotonation.<sup>15</sup> Given the known effect of placing a triisopropylsilyl (TIPS) group on an indole nitrogen to block *ortho*-lithiation,<sup>16</sup> we hypothesized that the combination of a bulky silyl protecting group on the carbazole nitrogen (to prevent C-1/C-8-lithiation) and the use of *n*BuLi–TMEDA (to dilithiate at C-4 and C-5)<sup>17</sup> would offer a simple and convenient approach to 4,5-difunctionalized carbazoles upon electrophilic trapping (Scheme 1). Based on this hypothesis, we herein report the first method for the direct functionalization at both the 4- and 5-positions of carbazole.

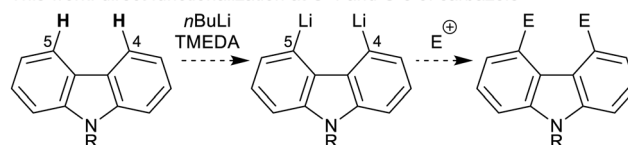
9*H*-Carbazole (**1b**) and silyl-substituted carbazoles **1a** (R = TIPS),<sup>21</sup> **1c** (R = TES) and **1d** (R = TBS) were treated with *n*BuLi under various conditions, followed by quenching with D<sub>2</sub>O, to determine the extent and position of lithiation, and the functional group compatibility of the *N*-silyl group (Table 1).

Previous work: C-functionalization of carbazoles



- Lithiation at C-1
- Metal-catalyzed functionalization at C-1 and C-8
- Metal-catalyzed functionalization at C-2 and C-7
- Electrophilic substitution at C-3 and C-6
- Ru-catalysed functionalization at C-4

This work: direct functionalization at C-4 and C-5 of carbazole



Scheme 1 Considerations in carbazole functionalization.

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Table 1 Deprotonation screen of carbazoles **1a–d**

Entry	1 (R)	Ligand	Time (h)	Temp. (°C)	% deuteration <sup>a</sup>		
					C1/8	C2–6	C4/5
1	<b>1a</b> (TIPS)	TMEDA	3	60	—	—	70
2	<b>1a</b> (TIPS)	TMEDA	6	60	—	—	83
3	<b>1a</b> (TIPS)	TMEDA	9	60	—	—	75
4	<b>1a</b> (TIPS)	TMEDA	3	68	—	—	83
5	<b>1a</b> (TIPS)	TMEDA	6	68	—	—	79
6	<b>1b</b> (H)	TMEDA	6	60	57	—	12
7	<b>1c</b> (TES)	TMEDA	6	60	100% desilylation		
8	<b>1d</b> (TBS)	TMEDA	6	60	88% desilylation		
9	<b>1a</b> (TIPS)	—	6	60	—	—	—
10	<b>1a</b> (TIPS)	PMDTA	6	60	Decomposition		

<sup>a</sup> R = TIPS; b R = H; c R = TES; d R = TBS

<sup>a</sup> Determined by relative integration of aromatic hydrogen signals in the <sup>1</sup>H NMR.

Subjecting *N*-TIPS carbazole **1a** to standard *n*BuLi-TMEDA conditions<sup>15a,c</sup> gave 70% deuterium incorporation at the 4,5-positions, as estimated by <sup>1</sup>H NMR (Table 1, entry 1),<sup>18</sup> thus establishing the feasibility of the approach. Extending the reaction time (entries 2 and 3) and increasing the reaction temperature (entries 4 and 5) gave increased levels of deuterium incorporation with the same C-4,5 regioselectivity; however evaporation of hexanes proved troublesome, thus favouring longer reaction times as the more practical conditions (entry 2). A screen of *N*-substituents showed that the free *9H*-carbazole (**1b**) gave deuteration preferentially at C-1 (entry 6), complementing prior literature in the absence of TMEDA.<sup>11a</sup> Smaller *N*-silyl substituents resulted in extensive desilylation (entries 7 and 8). Finally, exploring the role of the ligand, attempting the lithiation

in the absence of TMEDA gave no deuterium incorporation (entry 9), while using PMDTA, a known ligand for remote lithiation,<sup>19</sup> resulted in decomposition of **1a** (entry 10).

The conditions from Table 1, entry 2 were screened with a range of different electrophiles to give 4,5-disubstituted *N*-TIPS carbazoles directly from *N*-TIPS carbazole **1a** (Fig. 1). Alkylation with iodomethane gave a mixture of starting material, mono- and dimethylated products from which the major component, 4,5-dimethylcarbazole **2a**, could be separated in 38% yield. However the larger allyl bromide and benzyl bromide electrophiles failed to react.<sup>20</sup> Results of trapping with other carbon electrophiles also highlighted the difficulty in incorporating two groups in close proximity in the carbazole bay region. Although reaction with benzophenone was unsuccessful, use of paraformaldehyde gave separable bis-alcohol **3a**<sup>21</sup> in 43% yield, along with 23% of the monoalcohol. Reaction with CO<sub>2</sub> gave anhydride **4a** in a pleasing 63% yield. Trapping with hexafluorobenzene allowed for annulation to the *4H*-naphtho[1,2,3,4-*def*]carbazole **5a** in 34% yield.<sup>21</sup>

Reaction with heteroatom electrophiles was also investigated. Chlorination with C<sub>2</sub>Cl<sub>6</sub> gave a mixture of monochloride and dichloride, which were separable after removal of the TIPS group from the carbazole nitrogen using TBAF, with dichloride **6b**<sup>21</sup> isolated in 35% yield, along with 13% of the monochloride. Attempted incorporation of larger bromine or iodine were, however, very low yielding, again highlighted the difficulty in incorporating two groups in close proximity in the carbazole bay region.<sup>20</sup> More successful was where the electrophile was particularly small, or where a ring could be formed *in situ*, relieving potential *peri*-like steric and electronic repulsions in the hindered carbazole bay region. Reaction with B(OMe)<sub>3</sub> gave boronic anhydride **7a** in 66% yield after acidic work-up.<sup>22</sup> Reaction with dimethyldichlorosilane gave silafluorene **8a** in 74% yield.<sup>21</sup> Germafluorene **9a** was similarly formed in 39% yield.

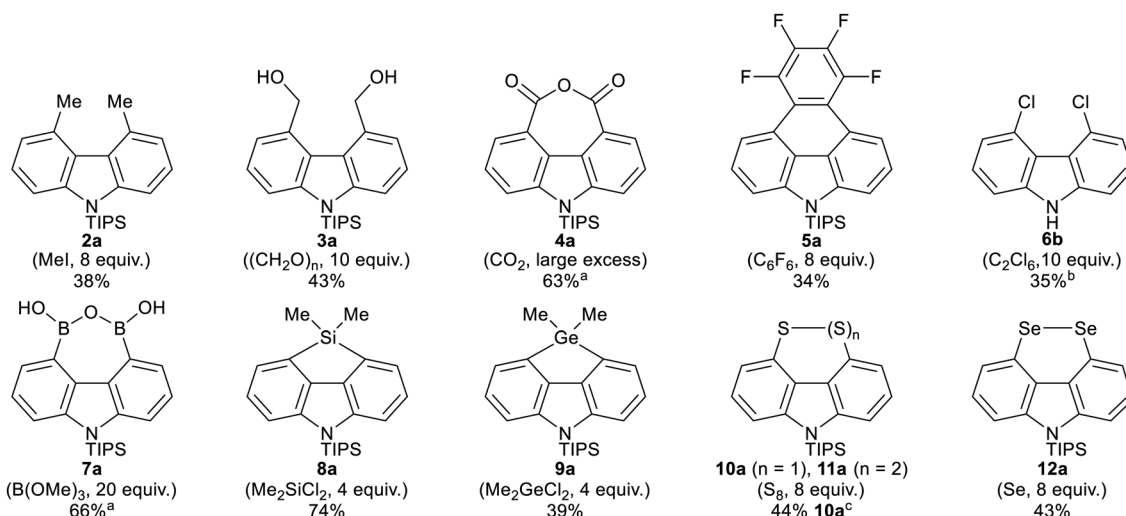
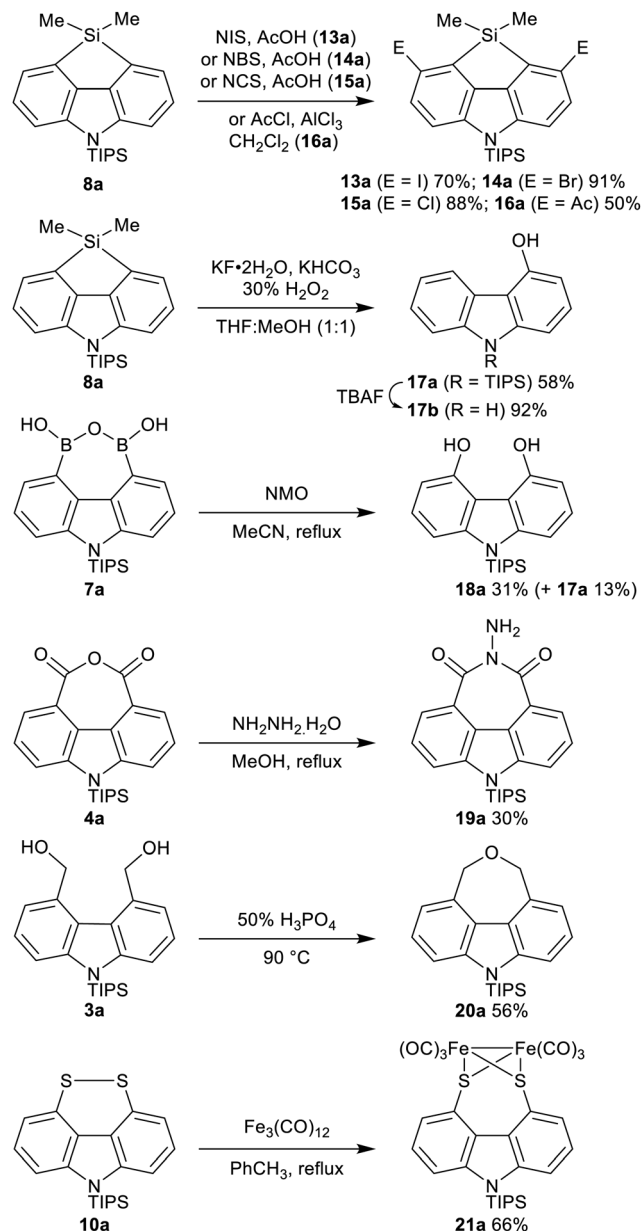


Fig. 1 4,5-Dilithiation and electrophilic trapping of *N*-TIPS carbazole **1a**. Reaction conditions: (i) *n*BuLi (4 equiv.), TMEDA (4 equiv.), 60 °C, 6 hours, (ii) THF, –78 °C then (electrophile, equiv.). % yields refer to isolated yields after purification by column chromatography unless otherwise stated. <sup>a</sup> Isolated yield after purification by trituration with *n*-hexane. <sup>b</sup> Isolated yield over two steps after removal of TIPS group with TBAF. <sup>c</sup> Yield of **10a** after NaBH<sub>4</sub> reduction of mixture of **10a** and **11a**.



Our interest in *peri*-substituted naphthalene disulfides and diselenides led us to investigate incorporation of sulfur and selenium at C-4 and C-5.<sup>23</sup> Reaction with elemental sulfur gave disulfide **10a** in 54% yield on a 3.1 mmol scale. On larger scale (6.2 mmol **1a**), mixtures of disulfide **10a** and trisulfide **11a** were obtained, which could be converged to **10a** in 44% overall yield by treatment with NaBH<sub>4</sub>. Reaction with elemental selenium gave diselenide **12a** in 43% yield without evidence of any triselenide formation. X-Ray analyses of disulfide **10b** and diselenide **12b**, obtained after removal of the *N*-TIPS groups with TBAF, are reported in the ESI.<sup>†21</sup>

The functionality embedded in these novel 4,5-disubstituted carbazoles can be further manipulated, increasing the range of carbazoles accessible from this methodology (Scheme 2).



Scheme 2 Selective transformations of 4,5-disubstituted carbazoles.

Regioselective electrophilic aromatic substitution of the fused carbazole-silafluorene **8a** provided the corresponding 3,6-disubstituted carbazoles **13a–16a**. Silafluorenes have broad application in materials and synthesis,<sup>24</sup> and the preparation of dihalides **13a–15a** in good yields opens up the investigation of these novel heteroatom-bridged heterofluorenes<sup>25</sup> in further cross-coupling reactions. Oxidation of **8a** under Dudley's conditions gave desymmetrized carbazole monoalcohol **17a**.<sup>26</sup> *N*-Desilylation of **17a** with TBAF gave 4-hydroxycarbazole (**17b**), a known starting point for natural product syntheses.<sup>27</sup> Complementary to the mono-oxidation of **8a**, the diol **18a** was the major product formed in the oxidation of boronic anhydride **7a** using NMO.<sup>28</sup> The potential of anhydride **4a** was demonstrated through reaction with hydrazine hydrate to give the *N*-iminoimide **19a** in 30% yield (unoptimized). Dehydration of diol **3a** to the carbazole-annulated cyclic ether **20a**<sup>21</sup> was achieved using H<sub>3</sub>PO<sub>4</sub>.<sup>29</sup> Oxidative insertion of Fe<sub>3</sub>(CO)<sub>12</sub> into the disulfide **10a** gave the Fe<sub>2</sub>(CO)<sub>6</sub> complex **21a**, of interest as a structural mimic of the active site of the enzyme [FeFe] hydrogenase.<sup>23d,30</sup>

In conclusion, the first methodology for the direct functionalization of the sterically hindered bay positions of carbazoles is reported, through the simple combination of a bulky TIPS group on the carbazole nitrogen and the use of *n*BuLi-TMEDA for deprotonation and directed lithiation. With 4,5-substitution being relatively unrepresented in the carbazole literature, and substitution in this bay region effecting, for example, the planarity of the carbazole,<sup>8,31</sup> this methodology opens up new opportunities for more systematic investigations and future applications in synthesis, catalysis, organic materials and biology.

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## Conflicts of interest

There are no conflicts to declare.

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- 21 CCDC 2079909–2079912 (**1a**, **8a**, **10b** and **12b** respectively) and CCDC 2080733–2080736 (**3a**, **5a**, **6b** and **20a** respectively) contain the supplementary crystallographic data for this paper†.
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- 31 The C4–C–C5 dihedral angle in disulfide **10b** (11.4(3)°, 10.5(3)°) and diselenide **12b** (13.6(4)°) can be compared with the essentially planar carbazole ring system in **1a** (–0.8(5)°). See ref. 21 for crystallography data.

