

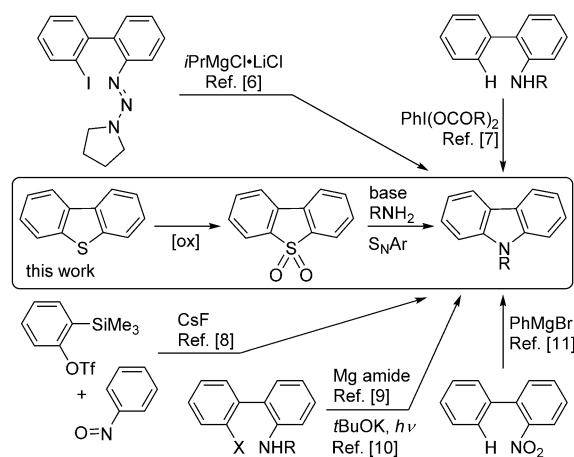
Transition-Metal-Free Synthesis of Carbazoles and Indoles by an S_NAr -Based “Aromatic Metamorphosis” of Thiaarenes**

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Abstract: Dibenzothiophene dioxides, which are readily prepared through oxidation of the parent dibenzothiophenes, undergo nucleophilic aromatic substitution with anilines intermolecularly and then intramolecularly to yield the corresponding carbazoles in a single operation. The “aromatic metamorphosis” of dibenzothiophenes into carbazoles does not require any heavy metals. This strategy is also applicable to the synthesis of indoles. Since electron-deficient thiaarene dioxides exhibit interesting reactivity, which is not observed for that the corresponding electron-rich azaarenes, a combination of a thiaarene-dioxide-specific reaction with the S_NAr -based aromatic metamorphosis allows transition-metal-free construction of difficult-to-prepare carbazoles.

Carbazoles represent an important class of heteroaromatic compounds, many of which find a wide range of applications as biologically active agents in medicinal chemistry^[1] and as hole-transport and light-emitting materials in organic electronics.^[2] For the last two decades, transition-metal catalysis has been used for constructing a carbazole skeleton by catalytic C–N bond formation, and includes Buchwald–Hartwig or Ullmann-type amination of haloaryls,^[3] oxidative C–H amination of aminobiaryls,^[4] or nitrene insertion from azidobiphenyls.^[5] Despite the high efficiency and reasonably wide scope, residual transition-metal impurities could adversely affect the biological properties or device performance of the final products.

Transition-metal-free synthesis of carbazoles has therefore undergone a renaissance for the last few years (Scheme 1). Liu and Knochel disclosed that halogen–magnesium exchange of 2-iodo-2'-triazenobiaryls resulted in formation of carbazoles.^[6] The groups of Chang and Antonchick reported intramolecular oxidative C–H amination of amino-



Scheme 1. Recent strategies for transition-metal-free synthesis of carbazoles.

biaryls with hypervalent iodine reagents.^[7] Studer and co-workers invented an interesting approach, that is, reactions of arynes with nitrosoarenes,^[8] but the yields were moderate and the regioselectivity was not as well controlled as is typically observed in arylene chemistry. Tokuyama and co-workers developed a base-mediated cyclization of 2-amino-2'-bromobiphenyls via arynes.^[9] From similar precursors, photoinduced cyclization occurs by a radical process to yield carbazoles.^[10] Kürti and co-workers disclosed facile reductive cyclization of 2-nitrobiaryls by means of PhMgBr.^[11] Regardless of these elegant protocols, there still remains ample room to develop novel strategies to prepare carbazoles without recourse to transition-metal catalysts.

Recently, our group has been interested in developing “aromatic metamorphosis”, which represents a transformation of an aromatic system into a different one through partial disassembly of the starting aromatic ring.^[12–14] Along these lines, herein we report transition-metal-free S_NAr -based aromatic metamorphosis of dibenzothiophenes to carbazoles via dibenzothiophene dioxides.

Our strategy begins with smooth oxidation of dibenzothiophenes with aqueous hydrogen peroxide (see the Supporting Information; Scheme 1).^[15] This conventional process is useful as the first step for the aromatic metamorphosis. Although aryl sulfones are known to undergo S_NAr reactions, the scope of the substitution is limited.^[16–19] Another difficulty could originate from the second intramolecular S_NAr reaction which would involve formal elimination of K_2SO_2 , which is a high-energy compound.^[20] The realization of the second, and the last step, sequential inter/intramolecular nucleophilic

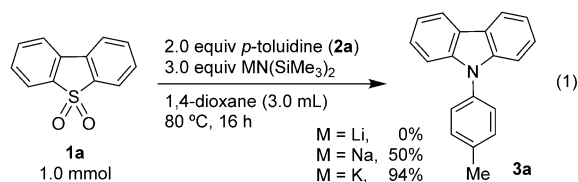
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aromatic substitution, required screening of the reaction conditions. We selected the reaction of dibenzo[*b,d*]thiophene 5,5-dioxide (**1a**) with *p*-toluidine (**2a**) as a model reaction for optimization [Eq. (1)]. The reaction of **1a** (1.0 equiv) with **2a** (2.0 equiv) by means of lithium hexamethyldisilazide (3.0 equiv) in 1,4-dioxane at 80 °C for 16 hours did not produce the desired carbazole **3a**. Interestingly, **3a** was obtained in 50 % yield when NaN(SiMe₃)₂ was employed. To our delight, KN(SiMe₃)₂ (potassium hexamethyldisilazide; KHMDS, 0.5 M toluene solution) gave the best result for isolating **3a** in 94 % yield. Other potassium bases such as K₂CO₃, K₃PO₄, KOAc, and KO^{*t*}Bu were totally ineffective. 1,4-Dioxane has proven to be the best cosolvent for toluene from the KHMDS solution. The reactions in 1,2-dimethoxyethane/toluene and in toluene as a sole solvent delivered **3a** in 91 and 61 % yield, respectively. Reducing the amount of **2a** or KHMDS led to lower yields of **3a**.



With the optimized the reaction conditions in hand, the scope with respect to the amines was examined (Table 1). The

Table 1: Scope with respect to the anilines.

R	t [h]	3	Yield [%] ^[a]
H	24	3b	77
4-MeO	16	3c	94
4- <i>t</i> Bu	16	3d	92
4-Me ₂ N	17	3e	91
4-CH ₂ =CH	17	3f	60
4-TIPSO	17	3g	28 (72 ^[b])
4-HO	24	3h	26 ^[c]
3-MeO	16	3i	70
3,5- <i>t</i> Bu ₂	17	3j	98
3,4-OCH ₂ O	17	3k	65
2,4,6-Me ₃	16	3l	67
4-Cl	18	3m	51 (13 ^[d])
4-F	13	3n	30 ^[e] (9 ^[f])

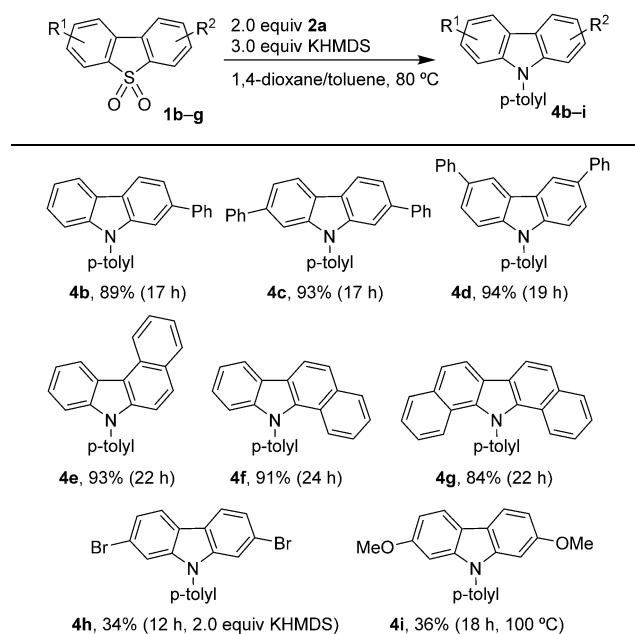
[a] Yield of isolated product. [b] The yield of **3h** as a byproduct. [c] 1,4-Dioxane (6.0 mL), 100 °C. [d] The yield of **3b** as a byproduct. [e] 2.0 equiv of KHMDS. [f] The yield of **3n'** as a byproduct. TIPS = triisopropylsilyl.

reactions proceeded efficiently with both electron-rich and electron-neutral anilines to yield **3b–l** in high yields. The vinyl functionality was tolerated under the reaction conditions to afford **3f** in moderate yield. The TIPS group of **2g** partly survived under the reaction conditions to yield a mixture of

the protected **3g** and unprotected **3h** quantitatively in 1:2.6 ratio. Unfortunately, unprotected *p*-aminophenol (**2h**) reacted sluggishly. Despite its steric hindrance, mesitylamine (**2l**) participated in the reaction to give **3l** in high yield. Moderately electron-withdrawing halo groups retarded the reaction and also underwent dehalogenation. The reaction with 4-chloroaniline (**2m**) yielded a 4:1 mixture of the desired **3m** and the hydrodechlorinated byproduct **3b** in 64 % yield. The fluoro analogue **2n** was less reactive and afforded **3n** in 30 % yield. In addition, the fluoro group was substituted with **2n** through an S_NAr reaction to yield *N*-(4-(9-carbazolyl)-phenyl)-4-fluoroaniline (**3n'**) as a byproduct. Alkylamines such as *p*-methoxybenzylamine reacted moderately (34 % yield with *p*-methoxybenzylamine in 18 h), possibly because of the less acidic NH protons. The attempted phenylation of carbazole with diphenyl sulfone under similar reaction conditions resulted in no conversion, probably because of the lower nucleophilicity of potassium carbazolate.

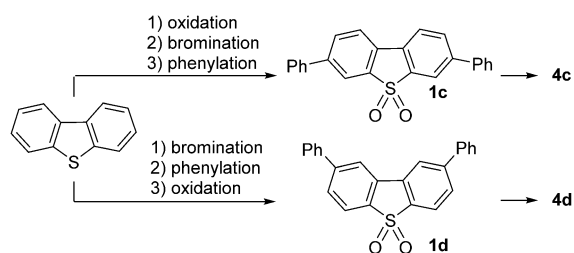
The scope with respect to the dibenzothiophene dioxide derivatives is broad (Table 2). Substituted or π -extended

Table 2: Scope with respect to dibenzothiophenes.



Yield is that of isolated product.

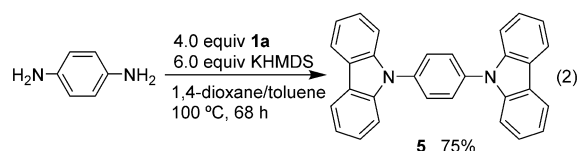
carbazoles were obtained efficiently. Gratifyingly, the sterically demanding **4g**^[21] was obtained in high yield. The reactions of substrates bearing bromo and methoxy groups gave slightly complex mixtures, but **4h** and **4i**, respectively, were isolated in moderate yield. Notably, we could synthesize 2,7-diphenyl-9-*p*-tolylcarbazoles (**4c**) in a controlled manner. While electron-rich dibenzothiophene, as well as carbazole, undergoes electrophilic bromination at the 3,6-positions with exclusive regioselectivity, dibenzothiophene dioxide is electron-deficient and delivers the 2,7-dibromo product (Scheme 2).^[15a,22] By changing the oxidation state of sulfur, we alter



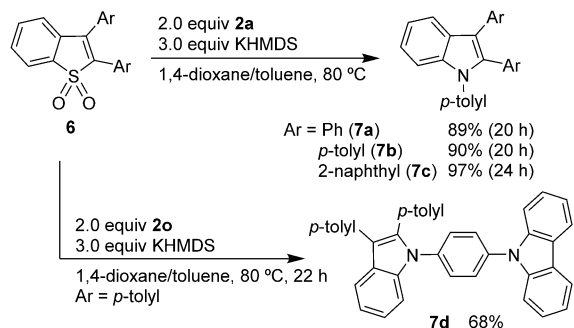
Scheme 2. Selective synthesis of two diphenyl isomers.^[15a, 22b]

the positions of the substituents in the final carbazoles **4c** and **4d**.^[21]

Benzene-1,4-diamine reacted with **1a** smoothly to afford the bis(carbazolyl)benzene **5**, an important structural unit which has found diverse applications in electronic devices^[23] [Eq. (2)]. Usually, bis(carbazolyl)benzenes are obtained in moderate yields through palladium-catalyzed or copper-mediated reactions of 1,4-dihalobenzene with carbazole at 170–200 °C.



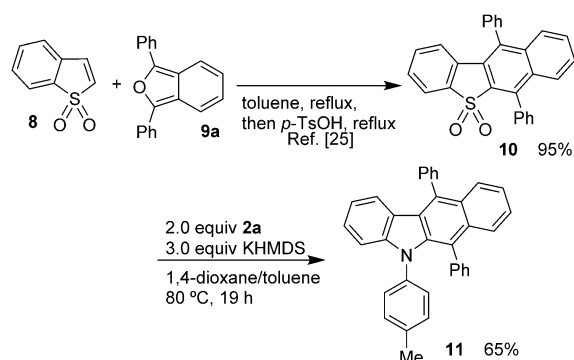
Our S_NAr -based amination was applicable to the synthesis of indoles (Scheme 3). Although 2,3-diaryl groups are neces-



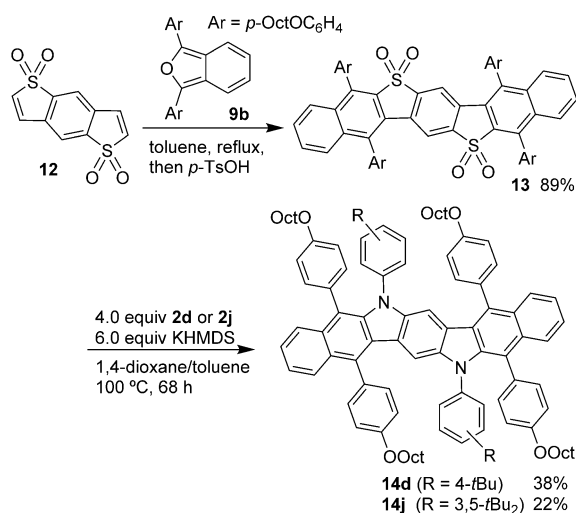
Scheme 3. S_NAr -based synthesis of indoles.

sary to avoid undesired conjugate addition onto the vinyl sulfoxide unit, densely arylated indoles (**7a–c**)^[21] were formed in excellent yields. The reaction of **6** with 4-(9-carbazolyl)aniline (**2o**) yielded an unsymmetric indole-carbazole hybrid (**7d**), which will find applications in organic electroluminescence.^[24]

The electron-deficient benzothiophene dioxide **8** showed uniquely high reactivity as a dienophile in the Diels–Alder reaction with isobenzofuran (**9a**),^[25] whereas indoles do not react with **9a** (Scheme 4). By taking advantage of the Diels–Alder reaction, we could extend the π -conjugation to form benzonaphthothiophene dioxide (**10**). Eventually, the π -extended benzocarbazole **11** was obtained in good overall yield from commercially available **8**.



Scheme 4. Sequential Diels–Alder reaction/carbazole formation.



Scheme 5. Concise synthesis of π -extended dibenzindolo[3,2-*b*]carbazoles.

We used this Diels–Alder-based approach to achieve significantly π -extended dibenzindolo[3,2-*b*]carbazoles (**14**; Scheme 5). The precursor bis(sulfone) **13** was obtained from a rather small aromatic compound (**12**) in 89% yield. Subsequently, expected quadruple C–N bond formations occurred to give **14d** and **14j** in 38 and 22% yield, respectively. In comparison with elegant yet burdensome construction of similar dibenzindolo[3,2-*b*]carbazoles for OFET devices,^[26, 27] this overall transformation highlights the synthetic potential of our carbazole synthesis.

Figure 1 shows the UV/visible and fluorescence spectra of **14j** in dichloromethane.^[28] The lowest energy absorption maximum was observed at $\lambda = 496$ nm, which is significantly red-shifted compared to typical absorption of indolo[3,2-*b*]carbazoles ($\lambda = 420$ – 440 nm).^[29] The compound **14j** showed bright-green fluorescence which has its intensity maximum at $\lambda = 511$ nm with a fluorescence quantum yield of 34%. Our S_NAr -based approach is thus useful for exploring π -extended carbazoles which show interesting functions.

The most likely mechanism would be the sequential inter/intramolecular S_NAr reactions with highly nucleophilic potassium amides (Scheme 6). The first intermolecular S_NAr reaction would be faster than the second intramolecular

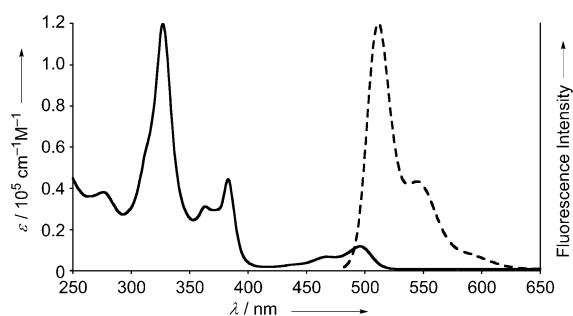
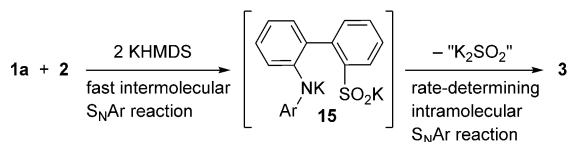
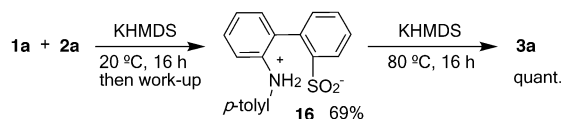


Figure 1. UV/visible (solid line) and fluorescence (dotted line) spectra of **14j** in CH_2Cl_2 .



Scheme 6. Plausible mechanism.



Scheme 7. Elucidation of mechanism.

reaction of **15**,^[30] and the carbazole formation necessitates a high temperature. The following experiments strongly support the mechanism (Scheme 7): **1a** reacted with **2a** in the presence of KHMDS at room temperature and the zwitterionic **16** was isolated.^[21] Subsequent exposure of **16** to KHMDS at 80 °C afforded **3a** quantitatively. Although there are several debatable reaction pathways for aromatic substitution of aryl sulfones,^[18] the isolation of **16** strongly indicates the intermediacy of **15** and hence the $\text{S}_{\text{N}}\text{Ar}$ pathway in our case.

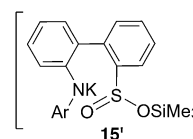
In conclusion, we have reported a new powerful and practical strategy to prepare carbazoles without recourse to transition-metal catalysts. Dibenzothiophene dioxide derivatives undergo sequential inter/intramolecular nucleophilic aromatic substitution with anilines in the presence of KHMDS to afford a wide range of carbazoles in a single operation. Considering that dibenzothiophene dioxides are readily available by oxidation of the parent dibenzothiophenes, one can concisely achieve aromatic metamorphosis of dibenzothiophenes to carbazoles. Since electron-deficient thiophene dioxides exhibit reactivity different from that of the corresponding electron-rich azaarenes, coupling a thiaarene-dioxide-specific reaction with the $\text{S}_{\text{N}}\text{Ar}$ carbazole synthesis opens up a new route to difficult-to-access carbazoles. Current efforts are directed toward exploring new aromatic metamorphosis in our laboratory.

Keywords: amination · aromatic substitution · fluorescence · sulfur heterocycles · synthetic methods

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