

# Experimental and Theoretical Investigations into the Unusual Regioselectivity of 4,5-, 5,6-, and 6,7-Indole Aryne Cycloadditions

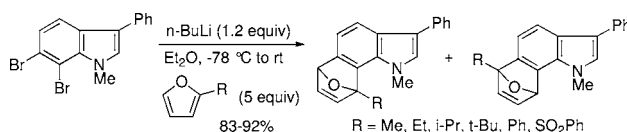
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## ABSTRACT



The 6,7-indolyne shows remarkable regioselectivity in its cycloaddition with 2-substituted furans. Electron-donating groups give predominantly the more sterically crowded product, while electron-withdrawing groups display the opposite regioselectivity. By contrast, 4,5- and 5,6-indolynes show no regioselectivity. Optimized electronic structure calculations using the M06-2X density functional and 6-311+G(2df,p) basis set revealed that the 6,7-indolynes are highly polar structures and that their cycloadditions have substantial electrophilic substitution character that leads to the observed preference for contra-steric products.

Benzyne and other common arynes (e.g., naphthalynes and pyridynes) have fascinated chemists for decades. Surprisingly, arynes occurring at all three benzenoid positions of the ubiquitous indole nucleus (indole arynes or indolynes 1–3, Figure 1) were, until recently, unknown. These newest members of the aryne family represent, inter alia, potentially attractive and powerful new tools for the total synthesis of complex natural products such as the nodulisporic acids,<sup>1</sup> trikentrins,<sup>2</sup> herbindoies,<sup>2a,c</sup> and the teleocidins.<sup>3</sup> (Figure 2). We recently discovered the first successful and general route to these reactive intermediates via metal–halogen exchange

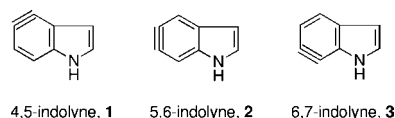


Figure 1. Benzenoid indole aryne (indolyne) systems.

of *o*-dihalides<sup>4</sup> and later via fluoride-induced decomposition of *o*-silyltriflates.<sup>5</sup> We subsequently examined the regiose-

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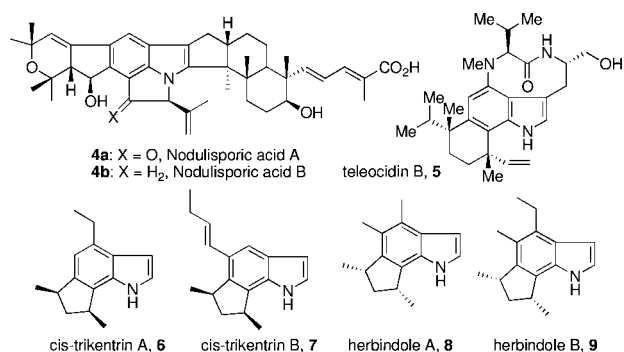
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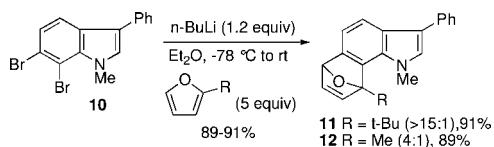


**Figure 2.** Examples of complex indole natural products.

lectivity of indolynes in Diels–Alder reactions with 2-substituted furans.<sup>5a</sup> These studies quickly led to the first application of indolynes in the construction of natural products and culminated in a concise and efficient total synthesis of (±)-*cis*-triketrin A and the (±)-herbindoles A and B.<sup>6</sup> This work and recently that of Garg clearly validated the utility of indolynes to gain quick access to architecturally challenging structures.<sup>7</sup>

During the course of these investigations, we were surprised to find that 2-substituted furans in particular displayed a striking regioselectivity when paired with the 6,7-indolynes (Scheme 1).<sup>5a</sup>

**Scheme 1.** Regioselective 6,7-Indolyne Cycloadditions with 2-Substituted Furans



Using 2-*t*-butylfuran, for example, the contrasteric product **11** was formed in at least a 15:1 preference over the alternate regioisomer. Even with the less sterically demanding 2-methylfuran, there was a 4:1 preference for the indicated isomer. Analysis of the products by various 2D NMR methods (COSY, NOESY, HMBC, and HSQC) allowed for the unequivocal assignment of the structures.

By contrast, there was no preference for either regioisomer in the 4,5- and 5,6-indolyne cycloaddition examples involving 2-*t*-butylfuran. In each case, a nearly 1:1 ratio of cycloadducts was formed in excellent yield (Scheme 2).<sup>5a</sup>

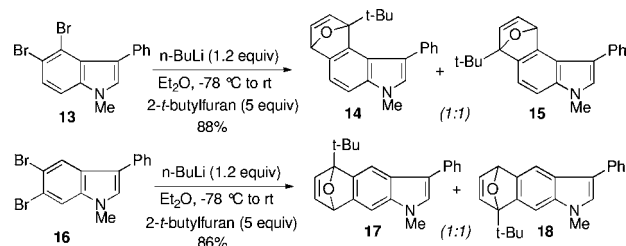
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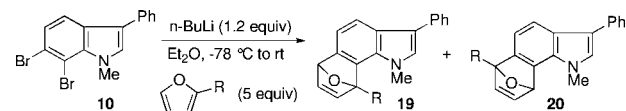
**Scheme 2.** Furan and Azide Cycloadditions with 4,5-, and 5,6-Indolynes



Intrigued by these observations, we wished to identify the fundamental electronic and steric parameters that account for these fascinating reaction profiles. The information obtained from this study will provide a better understanding of the structure and reactivity of the indole aryne system. This knowledge in turn should result in more versatile strategies and benefits for complex molecule total synthesis.

We began our investigation by examining a series of 2-substituted furans featuring electron-donating straight chain alkyl, branched alkyl, and aryl groups and electron-withdrawing groups (Table 1).<sup>8</sup> The electron-donating groups

**Table 1.** Regioselective 6,7-Indolyne Cycloadditions with 2-Substituted Furans



entry	furan, R	19	20	yield, %
1	Me	80	20	89
2	Et	84	16	90
3	<i>i</i> -Pr	94	6	88
4	<i>t</i> -Bu	98	2	91
5	Ph	>99	<1	92
6	SO <sub>2</sub> Ph	<1	>99	83

(entries 1–5) showed a remarkable preference for the more crowded cycloadduct **19**, while the opposite regiochemistry was found for an electron-withdrawing case (entry 6).

In earlier studies involving the reaction of monosubstituted 3-benzynes with 2-substituted furans (Me, Et, *i*-Pr, *t*-Bu),<sup>9</sup> regioselectivity correlated well with the strength of inductively electron-withdrawing groups such as F and OMe on the benzyne to favor contrasteric products, while it was reversed with inductively electron-donating groups such as Me.

(8) 2-Methyl-, 2-ethyl-, and 2-*t*-butylfuran are readily available from commercial sources. The others were prepared as follows. (a) 2-*i*-Propylfuran: see ref 8c. (b) 2-Phenylfuran: Negishi, E.; Takahashi, T.; King, A. O. *Org. Synth.* **1988**, *66*, 67–72. (c) 2-Phenylsulfonylfuran: Malanga, C.; Aronica, L. A.; Lardicci, L. *Synth. Commun.* **1996**, *26*, 2317–2327.

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To characterize this phenomenon in additional detail, electronic structure calculations were undertaken to predict the structures and reactivities of the *N*-methyl-4,5-, 5,6-, and 6,7-indolynes with furan and 2-alkylfurans. Similar calculations were also undertaken for the benzofuran and benzothiophene analogues of the indolynes as these heterocycles may prove useful for the construction of other interesting natural (or unnatural) products. With respect to methodology, all structures were optimized using the M06-2X density functional<sup>10</sup> and 6-311+G(2df,p) basis set<sup>11</sup> as implemented in MN-GFM,<sup>12</sup> a locally modified version of the Gaussian03 software package.<sup>13</sup> Analytical frequency calculations were employed to characterize the nature of all gas-phase structures as minima or transition states. In select instances, the effects of diethyl ether solvation were taken into account using the SMD<sup>14</sup> implicit solvation model.

Table 2 lists the activation free energies computed for the reaction of the *N*-methylindolynes with the furans. With unsubstituted furan,  $\Delta G^\ddagger$  is similar for the 4,5- and 5,6-indolyne isomers, and slightly smaller for the 6,7-isomer. Considering 2-*t*-butylfuran, the predicted free energies of activation are reduced by 3–4 kcal/mol compared to reaction with furan itself in the 4,5- and 5,6-indolyne cases. Regioselection becomes possible upon 2-substitution of the furan, and the differential free energies of activation in these two instances are predicted to be 1 kcal/mol or less. In the 6,7-indolyne case, by contrast, a free energy of activation similar to those for 4,5 and 5,6 is predicted for the formation of **26a** (which is equivalent to **19** but lacks the 3-phenyl substituent), but no transition-state (TS) structure for the formation of **26b** (corresponding to desphenyl **20**) could be located—the approach of the furan to the indolyne led smoothly and without barrier to the final tetracyclic product in every instance.

To better understand this point, the reactions of other 2-alkylfurans with **23** were modeled. With R = Me, Et, or *i*-Pr, TS structures for formation of **26b** could be found in every instance. Interestingly, the differential free energies of activation for formation of **26a** vs **26b** favored formation

**Table 2.**  $\Delta G^\ddagger$  Values in kcal/mol for the Reaction of Indolynes with Substituted Furans

entry	aryne	furan, R	$\Delta G^\ddagger$ (a)	$\Delta G^\ddagger$ (b)
1	<b>21</b>	H	10.5	
2	<b>22</b>	H	10.0	
3	<b>23</b>	H	8.7 (8.4) <sup>a</sup>	
4	<b>21</b>	<i>t</i> -Bu	6.3	7.3
5	<b>22</b>	<i>t</i> -Bu	6.9	6.1
6	<b>23</b>	<i>t</i> -Bu	7.0 (8.0)	<sup>b</sup>
7	<b>23</b>	Me	9.2 (9.4)	7.5 (7.8)
8	<b>23</b>	Et	9.0 (9.4)	7.8 (8.2)
9	<b>23</b>	<i>i</i> -Pr	9.7 (10.3)	7.6 (8.2)

<sup>a</sup> Values in parentheses include continuum ethereal solvation effects.  
<sup>b</sup> No barrier to reaction is predicted in the gas phase or continuum solution.

of the latter in each case, with the margin being largest (2.1 kcal/mol) for R = *i*-Pr. Ethereal solvation effects on the free energies of activation are predicted from the SMD continuum model to be 1 kcal/mol or smaller in every instance and to have no influence on regioselection.

Inspection of the TS structures for the formation of **26b** with the smaller alkyl groups provides insight into the failure to find such a TS structure for *t*-Bu. There are significant steric interactions between the *N*-Me group of the indolyne and the 2-alkyl group on the furan so that the bond forming to the 6 position is substantially shorter than that to the 7 position. The difference in the two formation bond lengths becomes increasingly long as the substituent is varied from Me to Et to *i*-Pr. This increase, however, is probably *not* associated with larger steric bulk since each of these furan 2-alkyl substituents orients a C–H bond toward the *N*-methyl group to minimize steric clash. Instead, it becomes clear from consideration of the preference for **26b** over **26a** and inspection of partial atomic charges that the 6,7-“cycloaddition” has substantial electrophilic substitution character.

Thus, the electron-poor indolyne attacks the 2-substituted furan to generate the more stable 2-alkyldihydrofurylcarbenium ion. As the 2-substituent is varied from Me to Et to *i*-Pr, the increased stabilization provided by the larger alkyl groups increases this character and leads to increased regioselection and increased electrophilic substitution character so that bond formation becomes decreasingly synchronous. Thus, it appears that in the case of R = *t*-Bu the combination of unavoidably increased sterics (there is no longer a C–H bond that can be disposed toward the *N*-Me group) and enhanced carbenium ion stabilization switches

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the mechanisms for a highly asynchronous but still concerted cycloaddition to a “stepwise” electrophilic substitution and subsequent ring closure. We place “stepwise” in quotes here because no TS structures nor intermediates can be located in either the gas phase or ethereal continuum solution for this process. To the extent that the reaction has an actual free energy of activation, it is likely associated with the displacement of discrete solvent molecules between the reacting partners that is not included in the present computational models, which otherwise are in reasonable quantitative agreement for predicted regioselection in **26a** and **26b** compared to **19** and **20**.

Support for the polarization of the 6,7-indolyne that makes the 6 position substantially more electrophilic than the 7 position may be obtained from inspection of atomic polar tensor partial charges<sup>15</sup> and the molecular geometry. Thus, C(6) is predicted to have a charge 0.26 au more positive than C(7), and the C(5)C(6)C(7) bond angle is predicted at the M06-2X level to be 135.3° while the C(6)C(7)C(7a) angle is predicted to be 117.2°. The former value is more consistent with carbocationic character, while the latter is more consistent with carbanionic character. For comparison, the corresponding bond angles in the analogous indyne (i.e., the hydrocarbon generated by replacing the *N*-methyl group with a methylene) are predicted at the DFT level to be 128.2 and 125.5°.<sup>16</sup> Thus, the C(6)–C(7) bond is strongly polarized in the expected direction by the nearby C(7a)–N bond dipole.

At the M06-2X level, the C(7a)–N bond dipole is predicted to lead to some polarization of the formal triple bonds in the 4,5- and 5,6-indolynes as well, although to a substantially smaller degree than in the 6,7 case (cf. entries 4 and 5 in Table 2). The failure to observe any regioselection under experimental conditions (reactions of Scheme 2) may reflect reduced polarization associated with the 3-phenyl group in the experimental substrate, or some degree of solvent leveling of the gas-phase predictions, or simply overpolarization predicted at the DFT level, but the effect in any case is predicted to be only about 1 kcal/mol.

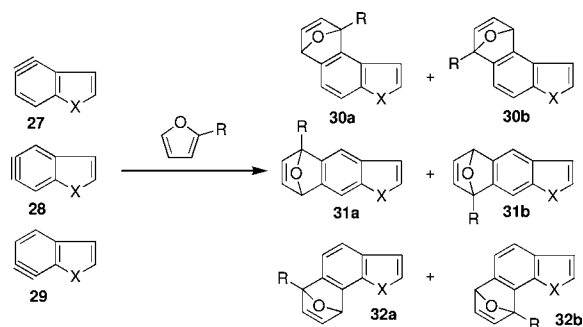
Given the substantial synthetic utility of the indolyne cycloadditions, we modeled the analogous cycloadditions of furan and 2-*t*-butylfuran with benzofuran and benzothiophene; results are presented in Table 3. Very similar reactivity profiles are predicted for all three heterocyclic systems, with the benzofuran case being predicted to have the lowest free energies of activation by a small margin, as would be expected given the greater electrophilicity of this heterocycle. This suggests that synthetic efforts based on these other heterocycles should be equally fruitful.

Benzynes derived from benzannulated heterocycles are thus revealed to be powerful synthetic tools for the construction of highly functionalized tetracycles through [4 + 2]

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**Table 3.**  $\Delta G^\ddagger$  Values in kcal/mol for the Reaction of Benzofurans and -Thiophenes with Substituted Furans



entry	aryne, X	furan, R	$\Delta G^\ddagger$ (a)	$\Delta G^\ddagger$ (b)
1	<b>27</b> , O	H	9.2	
2	<b>28</b> , O	H	9.1	
3	<b>29</b> , O	H	7.3	
4	<b>27</b> , O	<i>t</i> -Bu	4.9	6.4
5	<b>28</b> , O	<i>t</i> -Bu	5.8	4.9
6	<b>29</b> , O	<i>t</i> -Bu	6.2	<sup>a</sup>
7	<b>27</b> , S	H	9.6	
8	<b>28</b> , S	H	8.9	
9	<b>29</b> , S	H	8.4	
10	<b>27</b> , S	<i>t</i> -Bu	5.4	6.5
11	<b>28</b> , S	<i>t</i> -Bu	5.5	4.8
12	<b>29</b> , S	<i>t</i> -Bu	5.8	<sup>a</sup>

<sup>a</sup> No barrier to reaction is predicted in the gas phase or continuum solution.

cycloaddition reactions. Dipolar effects can induce regioselection with asymmetrically substituted dienes when polarization of the benzyne triple bond imparts substantial asynchronous electrophilic substitution character to the initial bond forming step.

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**Supporting Information Available:** NMR and other spectroscopic data for compounds used in this study and experimental details for their preparation as well as optimized cartesian coordinates for all computed structures. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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