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Copper-Catalyzed Remote C–H Arylation of Polycyclic Aromatic Hydrocarbons (PAHs)

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Abstract

The regioselective C–H arylation of substituted polycyclic aromatic hydrocarbons (PAHs) is a desired but challenging task. Herein, a copper-catalyzed C7–H arylation of 1-naphthamides has been developed by using aryliodonium salts as arylating reagents. This protocol does not need to use precious metal catalysts and can tolerate wide functional groups. Under standard conditions, the remote C–H arylation of other PAHs including phenanthrene-9-carboxamide, pyrene-1-carboxamide and fluoranthene-3-carboxamide has also accomplished, which provides an opportunity for the development of diverse organic optoelectronic materials.

Keywords

Copper-Catalyzed; C–H Arylation; Polycyclic Aromatic Hydrocarbons (PAHs); Non-precious Metal Catalyst; Regioselectivity

Introduction

Polycyclic aromatic hydrocarbons (PAHs) with rigid planar structure, such as naphthalene, phenanthrene, pyrene and their derivatives, can usually emit relatively strong fluorescence, and have been widely applied in many scientific areas including chemistry, biomedicine and material science.[1–6] The arylation reaction of PAHs is an important strategy to further extend the π-conjugation length, which can effectively adjust the photophysical properties of molecules, thus having drawn much attention. Transition metal-catalyzed C–X/C–M cross-coupling reactions such as Suzuki and Stille couplings are main approach to achieve the arylation of PAHs.[7–11] However, the selective arylation on the C7-position of 1-naphthoic acid derivatives remains a challenging task due to the inaccessibility of the corresponding 7-halonicthalamethane substrates.[12]

Recently, transition metal-catalyzed C–H bond functionalization has emerged as a powerful tool to construct various biaryl skeletons.[13–17] The direct C7–H arylation of 1-naphthoic acid derivatives is undoubtedly a more effective route for the synthesis of 7-aryl naphthalene derivatives. Although the transition metal-catalyzed C2–H and C8–H arylations of 1-naphthoic acid derivatives have been widely reported, the studies on their C7–H arylation remain rare.[18–25] Our group has recently reported F" reagent-promoted Pd-catalyzed C7–H arylation of 1-naphthamides, but this method still suffers from a few disadvantages (Scheme 1).[26] First, precious metal palladium is employed as a catalyst. Moreover, stoichiometric F" reagent is needed to oxidize Pd(II) species to more electrophilic high-valent cationic Pd(IV). In addition, this protocol is not compatible with other PAHs except naphthalene, such as phenanthrene, pyrene and fluoranthene, and cannot tolerate some special functional groups, such as alkenyl and alkynyl groups. As a component part of our ongoing research on direct C–H bond functionalization,[20,27–29] we herein represent a copper-catalyzed remote C–H arylation of PAHs with aryliodonium salts as arylating reagents (Scheme 1). This protocol is compatible with different PAH substrates including 1-naphthamides, phenanthrene-9-carboxamide, pyrene-1-carboxamide and fluoranthene-3-carboxamide, which provides an opportunity for the development of diverse organic optoelectrical materials.

Scheme 1. Direct C–H Arylation of PAHs

Our recent work: Pd-catalyzed C7–H arylation of 1-naphthamides

This work: Cu-catalyzed remote C–H arylation of PAHs

Our investigation commenced with the reaction between N-(tert-butyl)-1-naphthamide 1a and mesityl(phenyl)iodonium...
triflate 2a (for detailed optimization, see Table S1, ESI†).

Initially, the reaction was performed in 1,2-dichloroethane (DCE, 1 mL) at 80 °C for 24 h in the presence of Cu(OTf)₂ (10 mol%) as a catalyst. The direct C7−H arylation product N-(tert-butyl)-7-phenyl-1-naphthamide (3a) was gained in 79% yield (Table 1, entry 1). Gratifyingly, when the reaction temperature was reduced to 70 °C, 3a was obtained in 92% yield (Table 1, entry 2). The C7−H arylation could also occur with active copper powder as a catalyst (Table 1, entry 5). Other copper sources including CuO, CuCl and Cu(OAc)₂ were also found to be effective catalysts in this reaction, albeit with slightly lower yields (Table 1, entries 6-8). The control experiment confirmed that this transformation did not occur in the absence of Cu catalyst (Table 1, entry 9). The screening of other solvent, such as dichloromethane (DCM), ortho-dichlorobenzene (ODCB), CHCl₃ and PhCF₃, indicated that DCE was still the best effective (Table 1, entries 10-13). Finally, the optimal reaction system was established, which composed of Cu(OTf)₂ (10 mol%) in DCE (1.0 mL) at 70 °C under a nitrogen atmosphere for 24 hours.

Results and Discussion

Table 1. Optimization of Reaction Conditions

<table>
<thead>
<tr>
<th>Entry</th>
<th>Solvent</th>
<th>[Cu]</th>
<th>T (°C)</th>
<th>Yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>DCE</td>
<td>Cu(OTf)₂</td>
<td>80</td>
<td>79</td>
</tr>
<tr>
<td>2</td>
<td>DCE</td>
<td>Cu(OTf)₂</td>
<td>70</td>
<td>92</td>
</tr>
<tr>
<td>3</td>
<td>DCE</td>
<td>Cu(OTf)₂</td>
<td>90</td>
<td>53</td>
</tr>
<tr>
<td>4</td>
<td>DCE</td>
<td>Cu(OTf)₂</td>
<td>60</td>
<td>41</td>
</tr>
<tr>
<td>5</td>
<td>DCE</td>
<td>Cu</td>
<td>70</td>
<td>54</td>
</tr>
<tr>
<td>6</td>
<td>DCE</td>
<td>CuO</td>
<td>70</td>
<td>81</td>
</tr>
<tr>
<td>7</td>
<td>DCE</td>
<td>CuCl</td>
<td>70</td>
<td>84</td>
</tr>
<tr>
<td>8</td>
<td>DCE</td>
<td>Cu(OAc)₂</td>
<td>70</td>
<td>80</td>
</tr>
<tr>
<td>9</td>
<td>DCE</td>
<td>-</td>
<td>70</td>
<td>nd</td>
</tr>
<tr>
<td>10</td>
<td>DCM</td>
<td>Cu(OTf)₂</td>
<td>70</td>
<td>38</td>
</tr>
<tr>
<td>11</td>
<td>ODCB</td>
<td>Cu(OTf)₂</td>
<td>70</td>
<td>77</td>
</tr>
<tr>
<td>12</td>
<td>CHCl₃</td>
<td>Cu(OTf)₂</td>
<td>70</td>
<td>trace</td>
</tr>
<tr>
<td>13</td>
<td>PhCF₃</td>
<td>Cu(OTf)₂</td>
<td>70</td>
<td>trace</td>
</tr>
</tbody>
</table>

Table 1. Optimization of Reaction Conditions

*Reaction conditions: 1a (0.2 mmol, 1.0 equiv), 2a (0.3 mmol, 1.5 equiv), [Cu] (10 mol%) and solvent (1 mL) under N₂ for 24 h. Isolated yield. DCE = 1,2-dichloroethane. DCM = dichloromethane. ODCB = ortho-dichlorobenzene. nd: not detected.

With the optimal conditions in hand, we first examined the scope of aryliodonium salts. We were very pleased to find that a range of aryliodonium salts could be employed as arylation reagents, affording 7-arylated 1-naphthamides (3a–3r) in moderate to excellent yields (Table 2). This protocol tolerated wide functional groups, including electron-donating methyl, methoxy and phenoxo groups, as well as electron-withdrawing ester, trifluoromethyl, fluoro, chloro, bromo, iodo and formyl group. The arylation reactivity of aryliodonium salts with various substituents varied greatly due to the different electronic effect and steric hindrance. Arylation reagents with ortho-substituents led to slightly reduced yields (Table 2, 3g and 3j). Aryliodonium salts containing halogen substituents, especially bromo and iodo groups, could afford the desired products in moderate to good yields (Table 2, 3k–3o), making it possible to introduce useful functional groups into the products through the further transformation of corresponding aryl halides. 2-Naphthylidonium salt could react smoothly with 1a to provide 3p in 66% yield (Table 2, 3p). Moreover, thio-phen-2-yl iodonium salt could be tolerated, albeit with a lower yield (Table 2, 3r).

Table 2. Scope of Aryliodonium Salts

We next examined the scope of various naphthalene substrates (Table 3, 4a–4l). The electronic effect of C4-substituents on N-(tert-butyl)-1-naphthamide was not obvious. The 4-substituted 1-naphthamide substrates, whether with electron-donating methyl and methoxy groups, or with electron-withdrawing phenyl, ester, fluoro and bromo groups, gave the corresponding products in good to excellent yields (Table 3, 4a–4f). Substrates with C2-substituents also exhibited excellent reactivity, providing the desired products 4h and 4i in 85% and 80% yields, respectively (Table 3, 4h and 4i). Notably, 1-naphthamides with alkenyl (1m) and alkynyl (1n) groups were also suitable substrates for this direct C7−H arylation, affording 4j and 4k in good yields (Table 3, 4j and 4k).
Furthermore, this Cu-catalyzed direct C−H arylation could tolerate other PAH substrates. The regioselective arylation of PAHs is challenging, and so far, there are no examples on the selective remote C−H arylation of phenanthrene-9-carboxamide, pyrene-1-carboxamide and fluoranthene-3-carboxamide. Gratifyingly, the remote C−H arylation of these PAH substrates occurred smoothly, giving the corresponding arylation products in moderate to good yields (Table 3, 4m-4o).

Table 3. Scope of PAHs*

<table>
<thead>
<tr>
<th>Substituted naphthalenes</th>
<th>2a</th>
<th>Cu(OTf)₂ (10 mol%)</th>
<th>DCE, 73 °C, 24 h</th>
<th>Cu(OTf)₂ (10 mol%)</th>
<th>DCE, 73 °C, 24 h</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ph</td>
<td>O</td>
<td>Ph</td>
<td>4a, 89%</td>
<td>Ph</td>
<td>O</td>
</tr>
<tr>
<td>Ph</td>
<td>O</td>
<td>Ph</td>
<td>4c, 81%</td>
<td>Ph</td>
<td>O</td>
</tr>
<tr>
<td>Ph</td>
<td>O</td>
<td>Ph</td>
<td>4e, 71%</td>
<td>Ph</td>
<td>O</td>
</tr>
<tr>
<td>Ph</td>
<td>O</td>
<td>Ph</td>
<td>4g, 85%</td>
<td>Ph</td>
<td>O</td>
</tr>
<tr>
<td>Other PAH substrates</td>
<td>4i, 80%</td>
<td>Ph</td>
<td>O</td>
<td>Ph</td>
<td>4j, 53%</td>
</tr>
<tr>
<td>Other directing groups</td>
<td>4k, 73%</td>
<td>Ph</td>
<td>O</td>
<td>Ph</td>
<td>4l, 52%</td>
</tr>
</tbody>
</table>

*Reaction conditions: 1 (0.2 mmol), 2a (0.3 mmol) in DCE (1 mL) at 70 °C under N₂ for 24 h. Isolated yield. DCE = 1,2-dichloroethane.

This catalytic system was also compatible with substrates bearing other directing groups except tert-butylaminocarbonyl (Table 3, 4p-4s). When employing methylaminocarbonyl and cyclohexylaminocarbonyl as directing groups, the 7-arylation products of naphthalene rings were obtained in good yields (Table 3, 4p and 4q). Keto carbonyl group was also proved to be suitable directing groups, affording the corresponding arylation products in moderate yields (Table 3, 4r and 4s).

Considering that Cu(0), Cu(I) and Cu(II) all could catalyze this C−H arylation reaction and referring to previous research results,[30-31] a Cu(I)/Cu(III) catalytic cycle was proposed (Scheme 2). First, Cu(I) is formed by the reduction or disproportionation of Cu(II). Then, arylidonium salt oxidizes Cu(I) to highly electrophilic Cu(III)-aryl intermediate I. The coordination of carbonyl oxygen to I gives intermediate II, which undergoes an aryl-transfer reaction via a Heck-like four-membered-ring transition state III to form the intermediate IV with Cu(III) and aryl group added at the C8- and C7-positions of naphthalene ring, respectively. Finally, the breakdown of C8–Cu bond delivers Cu(I), meantime, OTf anion takes away the proton from the C7-position, affording the desired product 3 or 4.

Scheme 2. Proposed Catalytic Cycle

Subsequently, the photophysical property of the arylation products 4k, 4n and 4o was investigated (Figures 1). Their absorption bands cover from 300 nm to 400 nm, which corresponds to π−π* electron transition (Figure 1a and Table S2). The measurement of emission spectra demonstrates that 4k and 4n emit violet fluorescence with emission maxima at 395 nm and 390 nm, respectively, while 4o exhibits a sky-blue emission with emission maximum at 477 nm (Figure 1b and Table S2).

Figure 1. a) UV-visible absorption spectra of 4k, 4n and 4o in toluene (1×10⁻³ mol/L). b) Emission spectra of 4k, 4n and 4o in toluene (1×10⁻³ mol/L).

Conclusion

In summary, we have developed a highly efficient strategy to accomplish the direct C7−H arylation of 1-naphthamides by the usage of Cu(II) as a catalyst and arylidonium salts as arylating reagents, which features mild reaction conditions, excellent functional group tolerance, and moreover, does not need to use precious metal catalysts. This protocol is also compatible with other PAH substrates including phenanthrene-9-carboxamide, pyrene-1-carboxamide and fluoranthene-3-carboxamide, which provide an opportunity for the development of diverse organic photoelectrical materials.

Supporting Information

Detailed experimental procedures, characterization data and copies of ¹H and ¹³C NMR spectra of products.
Acknowledgment

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References