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Palladium-Catalyzed Arylation and Heteroarylation of Indolizines

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Abstract

A highly effective protocol for palladium-catalyzed selective arylation and heteroarylation of indolizines at C-3 has been developed. Mechanistic studies unambiguously support an electrophilic substitution pathway for this transformation.

Indolizines substituted at C-3 are very attractive heterocyclic units, as a number of representatives of this class¹ and, especially their partially or completely reduced analogues, indolizidine alkaloids² and related unnatural compounds,³ exhibit important biological properties. As a part of our program on developing efficient methods toward differently substituted indolizines, we were particularly interested in elaborating efficient approaches toward C-3-arylated indolizines. To date, no general convenient methods for synthesis of C-3-arylated indolizines exist.⁴ Scattered reports on synthesis of C-3-arylated indolizines via various modes of Chichibabin-type cyclocondensation⁵ are limited to particular substitution patterns⁶ and generally provide low to moderate yields of the products.⁷ Other methods such as dipolar cycloaddition of pyridinium ylides with alkynes,⁸ cyclopropanones,⁹ or cyclopropenes¹⁰ are limited to use of symmetric substrates, producing indolizines with two identical substituents at the pyrrole ring. Finally, synthesis of indolizines via cycloisomerization of alkynyl pyridines,^{11,12} a method recently developed within our group, potentially can give access to 3-arylindolizines; however, it would require employment of rather expensive 3-aryl-1-propynes and, in this case, is unlikely to be considered as synthetically useful. Thus, we decided to explore the possibility of direct, last-stage methods of arylation of the indolizine core. Although analogous reactions on various heterocyclic systems such as furans,¹³ thiophenes,^{13a,14} pyrroles,¹⁵ and indoles¹⁶ have long been known, no reports on arylations involving indolizines have been reported to date.¹⁷ Herein we report efficient palladium-catalyzed arylation and heteroarylation of indolizines, which allows for selective incorporation of aromatic or heteroaromatic substituent at the C-3 position of this heterocycle.

Initial experiments revealed that indolizine **1a** indeed underwent arylation in the presence of different palladium sources in DMSO; however, a 3-fold excess with respect to arylbromide

was required to attain high yields, and the reaction was rather sluggish, as it took 2 days for completion. Optimization experiments¹⁸ indicated that arylation of indolizines proceeded much faster (1–3 h) in the presence of catalytic amounts of PdCl₂(PPh₃)₂, KOAc (2 equiv), and H₂O (2 equiv) in NMP at 100 °C. Under these reaction conditions, a variety of differently substituted indolizines **1** underwent smooth arylation with an equimolar amount of **2** to afford C-3-arylated products **3** in good to excellent yields (Table 1). A variety of substituents are tolerated both on the indolizine and aryl halide. Nitro phenyl bromide **2a** reacted efficiently with all indolizines tested (Table 1, entries 1, 3, 5, 6, and 9). Notably, electron-rich bromoanisole **2c** underwent cross-coupling with both 2-cyanoindolizine **1b** and 2-methylindolizine **1d** (entries 4 and 8). Most remarkably, naphthyl bromide **2d** and even bromothiophene **2e** were able to undergo cross-coupling with **1e** to produce naphthyl- and hetaryl-substituted indolizines **3ec** and **3ed** in reasonable yields (entries 11 and 12).

Naturally, we were interested in elucidating the mechanism for this arylation reaction. Different mechanisms, depending on substitution patterns and reaction conditions, have been proposed for analogous arylations of other heterocycles (vide infra). We were particularly intrigued by Sharp's recent report on the regioselective Heck arylation of 3-ester-substituted furans and thiophenes (Scheme 1).^{13a} It occurred to us that arylation of indolizines **1a–c**, possessing EWG at C-2, and thus resembling Sharp's substrate **4**, may potentially proceed via a similar Heck-type process. Thus, we attempted a cascade Heck reaction¹⁹ of **1f**, aiming for spirocyclic compound **5** (Scheme 2). If successful, this process would not only bring firm support for the involvement of a Heck-type mechanism but also allow for straightforward access to polycyclic, indolizine-containing heteroaromatic compounds. However, trials on this cascade transformation with allyl ether **1f** were unsuccessful, producing a mixture of arylated indolizines **6** and **7** instead (Scheme 2). We also attempted to perform a reductive Heck reaction²⁰ on **1a** toward dihydroindolizine derivative **8** (Scheme 2). However, numerous efforts to perform this transformation under a variety of reducing conditions failed. We also attempted the arylation of 6-carboethoxyindolizine **1g**, expecting to obtain the product of the Heck reaction, indolizine **9** (Scheme 2). To our surprise, however, **1g** was selectively arylated at C-3 instead, producing **10** as a single regioisomer.

Failure to obtain any support for involvement of the Heck-type protocol in the arylation of indolizines prompted us to consider alternative mechanistic routes for this reaction. Different pathways proposed for arylation of various heterocycles, which can be adopted for arylation of indolizines, are depicted in Scheme 3. In addition to the Heck mechanism (**A**)²¹ discussed above, several other pathways have been suggested.

C–H activation motif **B** has been proposed by Miura for the “coordination-assisted” C-3 arylation of 2-carbamoyl-substituted thiophenes^{14a} and also by Sames as a possible route for arylation of unsubstituted indoles.¹⁵ To elucidate possible involvement of C–H activation (path **B**) in the Pd-catalyzed arylation of indolizines, a kinetic isotope effect study was performed (Scheme 4). We subjected deuteriumlabeled 2-carboethoxyindolizine **1a** to our arylation conditions. If C–H activation path **B** is in operation, a substantial isotope effect would be expected.²²

Experiments revealed, however, no isotope effect ($k_{H/D} = 1$), thus disfavoring possible involvement of pathway **B**.

Another related mechanism, cross-coupling pathway **C**, has been proposed by Miura²³ and Sames²⁴ for the copper-assisted C-2 arylation of azoles. Thus, we looked at cross-coupling (path **C**) as a possible route for our arylation reaction. Since it has been reported that arylation of acidic sites of the heterocycles is promoted by the addition of Cu salts,^{23,24} we

attempted arylation of **1a** in the presence of CuI. Contrary to our expectations, the presence of CuI greatly lengthened reaction times and reduced yields, thus disfavoring the involvement of mechanism **C**.

The fourth possible mechanism (**D**), electrophilic aromatic substitution, was initially proposed by Miura²³ and since then has often been considered as the most probable mechanism for arylation of heterocycles.^{13a,15,17,25} Accordingly, we drew our attention to mechanism **D**. Importantly, the absence of an isotope effect in the arylation of **1a**, mentioned above (Scheme 4), does not contradict with electrophilic mechanism **D**. In this case, C–H cleavage is a fast step,²⁶ and thus no isotope effect could be expected. Furthermore, it is generally believed that the pyrrole ring of indolizines is rather electron-rich, as it easily undergoes electrophilic substitution reactions.⁴ Our DFT calculations (B3LYP/6-31G*) perfectly confirmed this point, revealing that the pyrrole ring has an extended HOMO, whereas the LUMO mostly resides at the pyridine ring (Figure 1). Thus, it is quite reasonable that Pd-catalyzed electrophilic arylation selectively occurs at the electron-rich pyrrole ring. Similar arguments were provided by Li for the selective arylation of the imidazole ring in imidazopyrimidines.¹⁷ It is noteworthy that our calculations revealed that introduction of an EWG at C-6 of the pyridine ring does not qualitatively change the HOMO/LUMO distribution, which perfectly explains arylation of **1g** at the pyrrole ring to give **10** (Scheme 2).

Finally, to gain further support for electrophilic mechanism **D**, we performed kinetic studies. It was expected that electron-donating substituents at C-2 of indolizine would accelerate arylation at C-3, whereas electron-withdrawing groups should suppress the reaction. To test this hypothesis, competition experiments were run between indolizine (**1e**), 2-methylindolizine (**1d**), and 2-carboethoxyindolizine (**1a**). The relative rates of Pd-catalyzed arylation (method **A**) of **1e:1d:1a** were found to be: 1.00:0.97:0.66. The EWG-substituted indolizine **1a** underwent the slowest arylation, as expected; however, 2-methylindolizine **1d** was arylated slightly more slowly than parent indolizine **1e**. Potentially, this could be attributed to the steric bulk of the methyl group. To get a direct comparison of relative rates of Pd-catalyzed arylation with traditional Friedel–Crafts-type reactions, we set up electrophilic acylation of the same substrates with acyl chloride in the presence of AlCl₃ (method **B**). Experiment revealed relative rates of C-3 acylation of **1e:1d:1a** equal to 1:0.67:0.33. Although the values are not identical, the trend in rates for Friedel–Crafts acylation of indolizines reflects the trend in rates found for the Pd-catalyzed arylation reaction. We believe that these kinetic results, in combination with all experimental and computational data presented above, strongly support the electrophilic mechanism for the Pd-catalyzed arylation of indolizines.

In conclusion, the direct, efficient, regioselective C-3 aryl- and heteroarylation of indolizines has been developed. A variety of substituents on both the indolizine and aryl bromide are tolerated, providing rapid access to substituted indolizines in good to very high yields. Electrophilic substitution has been strongly supported as the mechanism through a combination of experimental and computational data.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1.
HOMO and LUMO Densities of **1e**.

Scheme 1. Proposed Heck Arylation of Heterocycles^{13a}

Scheme 2. Attempts to Perform Heck Reaction on Indolizines

Scheme 3. Possible Mechanisms for Pd-Catalyzed Arylation of Indolizines

Scheme 4. Isotope Effect Study

Table 1
Arylation and Heteroarylation of Indolizines

entry	indolizine	ArBr	product	yield (%) ^a
1				91
2				96
3				74
4				51
5				82
6				72

^aYields refer to pure isolated yields.

^bNMR yield.

^cReactions were run in NMP at 100 °C by using a 0.5 M solution of **1** with the following molar ratio: **1**:**2**:PdCl₂(PPh₃)₂:H₂O: KOAc = 1:1.2:0.05:2:2.

Table 2
Relative Rates of Pd-Catalyzed Arylation and Electrophilic Acylation of Selected Indolizines

relative rates			
R ¹		method A ^a	method B ^b
H	(e)	1.00	1.00
Me	(d)	0.97	0.67
CO ₂ Et	(a)	0.66	0.33

^a Performed with 5% PdCl₂(PPh₃)₂, **2** = 0.75 equiv of *p*-NO₂C₆H₄Br, KOAc, H₂O in NMP, 100 °C.

^b Performed with 1 equiv of AlCl₃, **2** = 0.75 equiv of AcCl, CH₂Cl₂, -78 °C.