

# Copper-Catalyzed, Directing Group-Assisted Fluorination of Arene and Heteroarene C–H Bonds

Thanh Truong, Kristine Klimovica, and Olafs Daugulis\*

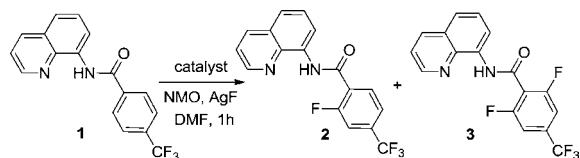
Department of Chemistry, University of Houston, Houston, Texas 77204-5003, United States

**S** Supporting Information

**ABSTRACT:** We have developed a method for direct, copper-catalyzed, auxiliary-assisted fluorination of  $\beta$ - $sp^2$  C–H bonds of benzoic acid derivatives and  $\gamma$ - $sp^2$  C–H bonds of  $\alpha,\alpha$ -disubstituted benzylamine derivatives. The reaction employs a CuI catalyst, a AgF fluoride source, and DMF, pyridine, or DMPU solvent at moderately elevated temperatures. Selective mono- or difluorination can be achieved by simply changing reaction conditions. The method shows excellent functional group tolerance and provides a straightforward way for the preparation of ortho-fluorinated benzoic acids.

Fluoroaromatic compounds possess inertness; high chemical, thermal, and metabolic stability; and unique electronic

**Table 1. Optimization of Reaction Conditions<sup>a</sup>**



entry	catalyst (mol %)	T, °C	1, %	2, %	3, %
1 <sup>b</sup>	Cu(OAc) <sub>2</sub> (25%)	100	4	17	5
2 <sup>b,c</sup>	Cu(OAc) <sub>2</sub> (25%)	100	<2	13	4
3	Cu(OAc) <sub>2</sub> (25%)	100	20	42	8
4	CuI (25%)	100	11	54	6
5	CuI (25%)	80	20	60	5
6	CuI (15%)	80	18	70	3
7 <sup>d</sup>	CuI (15%)	80	4	79	7
8 <sup>d,e</sup>	CuI (15%)	80	16	80	3
9 <sup>f</sup>	CuI (20%)	80	<2	17	38
10 <sup>e,f</sup>	CuI (20%)	80	10	52	33
11 <sup>f,g</sup>	CuI (20%)	80	<2	5	78

<sup>a</sup>Amide 0.25 mmol, AgF 3 equiv, NMO 4 equiv, DMF 1 mL. Yields were determined by GC analysis. <sup>b</sup>DMSO solvent. <sup>c</sup>PhI(OPiv)<sub>2</sub> oxidant instead of NMO. <sup>d</sup>AgF 4 equiv, NMO 5 equiv. <sup>e</sup>Pyridine solvent. <sup>f</sup>AgF 6 equiv, 8 equiv NMO, 1.5 h. <sup>g</sup>Pyridine additive 2 equiv.

properties. They are widely used as pharmaceuticals, agrochemicals, and imaging materials.<sup>1</sup> Classical methods for synthesis of fluoroaromatics such as the Balz–Schiemann reaction employ prefunctionalized starting materials and typically require harsh reaction conditions, thus limiting the scope of transformations.<sup>2,3</sup> More recently, aryl-fluorine bonds have been created by using transition-metal catalysis.

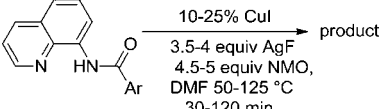
Specifically, Buchwald has shown that aryl (Ar) triflates can be converted to Ar fluorides under Pd catalysis.<sup>4</sup> Ritter has developed methods for stannane, boronic acid, and phenol conversion to fluoroaromatics.<sup>5</sup> Hartwig has shown that Ar iodides can be converted to Ar fluorides by a combination of a stoichiometric copper(I) complex and AgF.<sup>6</sup> Aryl stannanes and Ar borates have been converted to Ar fluorides by employing Cu(I) in combination with an electrophilic fluoride source.<sup>7</sup> Many of these processes have been developed on the basis of mechanistic studies;<sup>8</sup> however, these methods require the use of prefunctionalized starting materials. Nondirected fluorination of  $sp^3$  C–H bonds by radical methods has also been investigated.<sup>9</sup> Only a few groups have reported directed catalytic  $sp^2$  C–H bond fluorination. In a pioneering work, Sanford has shown that a pyridine directing group can be employed for a Pd-catalyzed arene fluorination by electrophilic fluorine sources. Mechanistic studies support involvement of high-valent Pd complexes in these reactions.<sup>10</sup> Yu has demonstrated Pd-catalyzed benzylamine triflamide and benzoic acid perfluoroaniline amide ortho-fluorination by *N*-fluoropyridine derivatives.<sup>11</sup> Recently, Sanford has shown that Pd-(OAc)<sub>2</sub>/AgF/PhI(OPiv)<sub>2</sub> system<sup>12</sup> can be employed for 8-methylquinoline benzylic C–H bond fluorination.<sup>13</sup> A general method for directed arene C–H bond fluorination by using a first-row transition metal catalyst has not been reported.<sup>15</sup> We disclose here a method for aminoquinoline and picolinamide-directed benzoic acid and benzylamine derivative ortho-fluorination under Cu catalysis.

In 2005, we introduced 8-aminoquinoline and picolinic acid auxiliaries for Pd-catalyzed  $sp^2$  and  $sp^3$  C–H bond arylation.<sup>14a</sup> Recently Cu-catalyzed sulfonylation and amination of C–H bonds in 8-aminoquinoline benzamides and benzylamine picolinamides was demonstrated.<sup>14b,c</sup> We hypothesized that 8-aminoquinoline and picolinic acid auxiliaries would promote Cu-catalyzed ortho-fluorination of  $sp^2$  C–H bonds on the basis of the following considerations: (1) Cu-catalyzed Ar halide fluorination has been reported in a macrocyclic polyamine system,<sup>16a</sup> (2) Cu-promoted C–H activation has been reported in the same system,<sup>16b</sup> and (3) it appears that both macrocyclic amine and 8-aminoquinoline benzamide ligands stabilize high-valent Cu intermediates.

Reaction of 8-aminoquinoline *p*-trifluoromethyl-benzamide was investigated with respect to Cu catalyst, fluorine source, oxidant, and solvent (Table 1). Best results for monofluorination were obtained by using CuI catalyst, NMO oxidant, and

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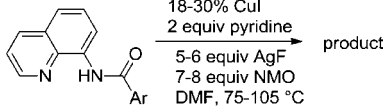
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Table 2. Monofluorination of Carboxylic Acid Derivatives<sup>a</sup>


entry	Ar	product	yield, %
1	4-CF <sub>3</sub> C <sub>6</sub> H <sub>4</sub>		71
2	4-MeC <sub>6</sub> H <sub>4</sub>		75
3	4-MeO <sub>2</sub> CC <sub>6</sub> H <sub>4</sub>		56
4	4-NCC <sub>6</sub> H <sub>4</sub>		62 60 <sup>b</sup>
5	2-CF <sub>3</sub> C <sub>6</sub> H <sub>4</sub>		80
6	2-MeC <sub>6</sub> H <sub>4</sub>		63
7 <sup>c</sup>	4-NO <sub>2</sub> C <sub>6</sub> H <sub>4</sub>		60
8	3-CF <sub>3</sub> C <sub>6</sub> H <sub>4</sub>		71
9	3-( <i>N</i> -Me-indolyl)		54
10 <sup>c</sup>	4-Pyridyl		62

<sup>a</sup>Amide 0.25 mmol, DMF 1 mL. Yields are isolated yields. Please see Supporting Information (SI) for details. <sup>b</sup>Reaction scale: 5 mmol. <sup>c</sup>Pyridine solvent.

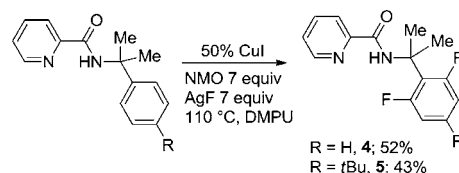
DMF solvent. Use of  $\text{PhI}(\text{OPiv})_2$  oxidant resulted in lower yields (entry 2). DMSO solvent is inferior to DMF due to starting material decomposition (entry 1 vs 4). Reaction can be run in pyridine (entry 8) which slows decomposition of starting material, albeit at the expense of reaction rate. Selective difluorination can be achieved by using higher loading of CuI

Table 3. Difluorination of Carboxylic Acid Derivatives<sup>a</sup>


entry	Ar	product	yield, %
1	4-CF <sub>3</sub> C <sub>6</sub> H <sub>4</sub>		67
2	4-MeO <sub>2</sub> CC <sub>6</sub> H <sub>4</sub>		62
3	NCC <sub>6</sub> H <sub>4</sub>		70
4	2-Naphthyl		70
5	4-MeOC <sub>6</sub> H <sub>4</sub>		75
6	4-FC <sub>6</sub> H <sub>4</sub>		61
7	3-MeC <sub>6</sub> H <sub>4</sub>		77
8 <sup>b</sup>	4-Pyridyl		61

<sup>a</sup>Amide 0.25 mmol, DMF 1 mL. Yields are isolated yields. Please see SI for details. <sup>b</sup>Pyridine solvent.

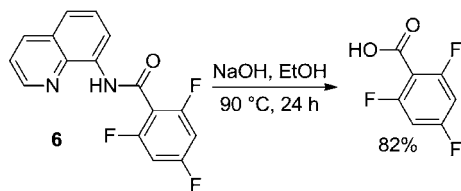
## Scheme 1. Fluorination of Picolinamides



(entries 9–11). Longer reaction times require pyridine additive (2 equiv) to prevent decomposition of amide substrate (entry 11).

Reaction scope with respect to monofluorination of 8-aminoquinoline benzamides is presented in Table 2. Both electron-rich (entries 2 and 6) and electron-poor (entries 1, 3–5, 7, 8) benzamides are reactive. Heterocyclic carboxamides

Scheme 2. Auxiliary Cleavage



containing indole (entry 9) and pyridine (entry 10) moieties are fluorinated in good yields. The reaction is functional-group tolerant, with carboxylate (entry 3), nitrile (entry 4), and nitro groups (entry 7) compatible with the fluorination conditions. For strongly electron-deficient substrates such as 4-nitrobenzoyl and pyridyl derivatives (entries 7 and 10), the reaction must be run in pyridine solvent to prevent decomposition of product. However, somewhat longer reaction times are required if pyridine is employed.

Optimization results in Table 1 show that, by increasing CuI and AgF loading and reaction time, clean difluorination can be obtained. Longer reaction times require the use of pyridine to prevent decomposition of aminoquinoline amides. Difluorination examples are presented in Table 3. Similar to monofluorination, electron-rich (entries 4, 5, 7), electron-poor (entries 1–3, 6), and heterocyclic (entry 8) amides can be efficiently difluorinated in good yields. Interestingly, difluorination of meta-substituted amides is possible (entries 4, 7), but contrasts with Pd-catalyzed C–H bond functionalization, where substitution at more hindered positions is typically not observed.<sup>17</sup> The observation is consistent with results obtained in the Cu-promoted sulfenylation of  $sp^2$  C–H bonds, where functionalization of hindered positions is possible.<sup>14b</sup>

Fluorination of benzylamine derivatives is also possible by using a picolinamide directing group (Scheme 1). However, the reactions are less efficient, requiring 50 mol % CuI catalyst, higher temperature, and DMPU solvent. Reasonable conversions could only be obtained with  $\alpha,\alpha$ -disubstituted benzylamines. This behavior is consistent with Cu-catalyzed amination and sulfenylation of C–H bonds.<sup>14b,c</sup>

Auxiliary can be cleaved by base hydrolysis. Thus, heating amide **6** with NaOH in ethanol for 24 h afforded high yield of trifluorobenzoic acid (Scheme 2).

While speculations about the reaction mechanism are premature at this point, Ribas has shown that Cu-catalyzed nucleophilic Ar fluorination and Ar halide exchange is possible in a geometrically constrained system.<sup>16a</sup> The reactions proceed via Cu(III) intermediates, thus showing that C–F reductive elimination from Cu(III) is possible under very mild conditions.<sup>7b</sup> Given that aminoquinoline amides stabilize high oxidation states in transition metals,<sup>14d</sup> it is likely that Cu-catalyzed aminoquinoline amide fluorination also proceeds via Cu(III) intermediates.

To conclude, we developed a method for direct, Cu-catalyzed, auxiliary-assisted fluorination of  $\beta$ - $sp^2$  C–H bonds of benzoic acid derivatives and  $\gamma$ - $sp^2$  C–H bonds of benzylamine derivatives. The reaction uses catalytic CuI or AgF as the nucleophilic fluoride source, and DMF, pyridine, or DMPU solvent at moderately elevated temperatures. The method allows for selective mono- or difluorination of benzamide substrates, shows excellent functional group tolerance, and provides a straightforward way for the preparation of ortho-fluorinated benzoic acids. Future work involves mechanistic

studies of the transformation and attempts to isolate reaction intermediates.

## ■ ASSOCIATED CONTENT

### Supporting Information

Detailed experimental procedures and characterization data for new compounds. This material is available free of charge via the Internet at <http://pubs.acs.org>.

## ■ AUTHOR INFORMATION

### Corresponding Author

olafs@uh.edu

### Notes

The authors declare no competing financial interest.

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## ■ REFERENCES

- (1) (a) Hollingworth, C.; Gouverneur, V. *Chem. Commun.* **2012**, 48, 2929. (b) Coenen, H. H.; Ermert, J. *Curr. Radiopharm.* **2010**, 3, 163. (c) Zhang, X.-J.; Lai, T.-B.; Kong, R. Y.-C. *Top. Curr. Chem.* **2012**, 308, 365. (d) Purser, S.; Moore, P. R.; Swallow, S.; Gouverneur, V. *Chem. Soc. Rev.* **2008**, 37, 320. (e) Jeschke, P. *ChemBioChem* **2004**, 5, 570.
- (2) (a) Balz, G.; Schiemann, G. *Chem. Ber.* **1927**, 60, 1186.
- (3) (a) Dawood, K. M. *Tetrahedron* **2004**, 60, 1435. (b) Adams, D. J.; Clark, J. H. *Chem. Soc. Rev.* **1999**, 28, 225. (c) Rozen, S. *Adv. Synth. Catal.* **2010**, 352, 2691. (d) Yamada, S.; Knochel, P. *Synthesis* **2010**, 2490. (e) Umemoto, T.; Tomizawa, G. *J. Org. Chem.* **1995**, 60, 6563.
- (4) (a) Watson, D. A.; Su, M.; Teverovskiy, G.; Zhang, Y.; Garcia-Fortanet, J.; Kinzel, T.; Buchwald, S. L. *Science* **2009**, 325, 1661. (b) Maimone, T. J.; Milner, P. J.; Kinzel, T.; Zhang, Y.; Takase, M. K.; Buchwald, S. L. *J. Am. Chem. Soc.* **2011**, 133, 18106.
- (5) (a) Furuya, T.; Kaiser, H. M.; Ritter, T. *Angew. Chem., Int. Ed.* **2008**, 47, 5993. (b) Tang, P.; Wang, W.; Ritter, T. *J. Am. Chem. Soc.* **2011**, 133, 11482. (c) Tang, P.; Furuya, T.; Ritter, T. *J. Am. Chem. Soc.* **2010**, 132, 12150. (d) Furuya, T.; Ritter, T. *J. Am. Chem. Soc.* **2008**, 130, 10060. Review: (e) Furuya, T.; Kamlet, A. S.; Ritter, T. *Nature* **2011**, 473, 470.
- (6) Fier, P. S.; Hartwig, J. F. *J. Am. Chem. Soc.* **2012**, 134, 10795.
- (7) (a) Ye, Y.; Sanford, M. S. *J. Am. Chem. Soc.* **2013**, 135, 4648. (b) Fier, P. S.; Luo, J.; Hartwig, J. F. *J. Am. Chem. Soc.* **2013**, 135, 2552.
- (8) (a) Grushin, V. V. *Acc. Chem. Res.* **2010**, 43, 160. (b) Vignalok, A. *Organometallics* **2011**, 30, 4802. (c) Dubinsky-Davidchik, I. S.; Potash, S.; Goldberg, I.; Vignalok, A.; Vedernikov, A. N. *J. Am. Chem. Soc.* **2012**, 134, 14027. (d) Zhao, S.-B.; Wang, R.-Y.; Nguyen, H.; Becker, J. J.; Gagné, M. R. *Chem. Commun.* **2012**, 48, 443. (e) Mankad, N. P.; Toste, F. D. *Chem. Sci.* **2012**, 3, 72.
- (9) (a) Bloom, S.; Pitts, C. R.; Miller, D. C.; Haselton, N.; Holl, M. G.; Urheim, E.; Lectka, T. *Angew. Chem., Int. Ed.* **2012**, 51, 10580. (b) Liu, W.; Huang, X.; Cheng, M.-J.; Nielsen, R. J.; Goddard, W. A., III; Groves, J. T. *Science* **2012**, 337, 1322. (c) Yin, F.; Wang, Z.; Li, Z.; Li, C. *J. Am. Chem. Soc.* **2012**, 134, 10401. (d) Amaoka, Y.; Nagatomo, M.; Inoue, M. *Org. Lett.* **2013**, 15, 2160. (e) Katcher, M. H.; Doyle, A. G. *J. Am. Chem. Soc.* **2010**, 132, 17402. Review: (f) Sibi, M. P.; Landais, Y. *Angew. Chem., Int. Ed.* **2013**, 52, 3570.
- (10) (a) Hull, K. L.; Anani, W. Q.; Sanford, M. S. *J. Am. Chem. Soc.* **2006**, 128, 7134. (b) Ball, N. D.; Sanford, M. S. *J. Am. Chem. Soc.* **2009**, 131, 3796.
- (11) (a) Wang, X.; Mei, T.-S.; Yu, Y.-Q. *J. Am. Chem. Soc.* **2009**, 131, 7520. (b) Chan, K. S. L.; Wasa, M.; Wang, X.; Yu, J.-Q. *Angew. Chem., Int. Ed.* **2011**, 50, 9081. Review: (c) Engle, K. M.; Mei, T.-S.; Wang, X.; Yu, J.-Q. *Angew. Chem., Int. Ed.* **2011**, 50, 1478.

- (12) Wu, T.; Yin, G.; Liu, G. *J. Am. Chem. Soc.* **2009**, *131*, 16354.
- (13) McMurtrey, K. B.; Racowski, J. M.; Sanford, M. S. *Org. Lett.* **2012**, *14*, 4094.
- (14) (a) Zaitsev, V. G.; Shabashov, D.; Daugulis, O. *J. Am. Chem. Soc.* **2005**, *127*, 13154. (b) Tran, L. D.; Popov, I.; Daugulis, O. *J. Am. Chem. Soc.* **2012**, *134*, 18237. (c) Tran, L. D.; Roane, J.; Daugulis, O. *Angew. Chem., Int. Ed.* **2013**, *52*, 6043. (d) Shabashov, D.; Daugulis, O. *J. Am. Chem. Soc.* **2010**, *132*, 3965.
- (15) CuF<sub>2</sub>-mediated fluorination of benzene: Subramanian, M. A.; Manzer, L. E. *Science* **2002**, *297*, 1665.
- (16) (a) Casitas, A.; Canta, M.; Solá, M.; Costas, M.; Ribas, X. *J. Am. Chem. Soc.* **2011**, *133*, 19386. (b) Ribas, X.; Jackson, D. A.; Donnadieu, B.; Mahía, J.; Parella, T.; Xifra, R.; Hedman, B.; Hodgson, K. O.; Llobet, A.; Stack, T. D. P. *Angew. Chem., Int. Ed.* **2002**, *41*, 2991. (c) Huffman, L. M.; Stahl, S. S. *J. Am. Chem. Soc.* **2008**, *130*, 9196. (d) Chen, X.; Hao, X.-S.; Goodhue, C. E.; Yu, J.-Q. *J. Am. Chem. Soc.* **2006**, *128*, 6790.
- (17) Daugulis, O.; Do, H.-Q.; Shabashov, D. *Acc. Chem. Res.* **2009**, *42*, 1074.