

Suzuki–Miyaura Cross-Coupling Reactions of Potassium Boc-Protected Aminomethyltrifluoroborate with Aryl and Hetaryl Mesylates

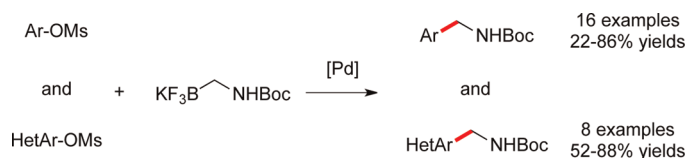
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Received May 4, 2012

ABSTRACT

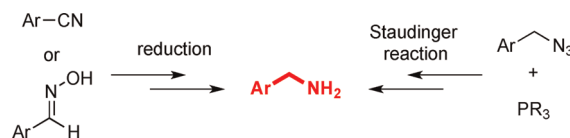


Palladium-catalyzed Suzuki–Miyaura cross-coupling reactions were studied with potassium Boc-protected aminomethyltrifluoroborate through C–O activation of various mesylate derivatives to afford the corresponding products in moderate to good yields.

Aminomethylarenes are encountered in many bioactive materials.¹ Primary aminomethyl subunits are particularly important targets for synthesis because they exhibit useful properties as drugs and inhibitors. Despite their importance, syntheses of aminomethyl substructures are not general. Reduction of aryl cyanides² or oximes³ and the Staudinger reaction⁴ of azides have often been used to build aminomethyl moieties (Scheme 1). However, these approaches have limitations because of their sensitivity to reducible functional groups and the instability of azides,⁵ respectively.

Transition metal mediated cross-coupling reactions of aminomethylmetallic species are one of the most straightforward strategies for preparation of primary

Scheme 1



aminomethyl moieties. To the best of our knowledge, three methods of Suzuki–Miyaura cross-coupling reactions to afford primary aminomethyl arenes using different protecting groups have been reported (Scheme 2).⁶ Among them, *p*-toluenesulfonyl (Ts)^{6a} or *N*-phthalimido (Phth)^{6b,c} groups have been utilized but are not ideal because the amine protecting groups are difficult to remove, requiring relatively harsh reaction conditions.⁷ Recently, we

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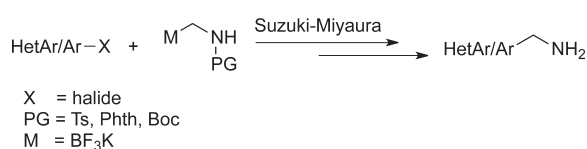
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demonstrated that potassium Boc-protected aminomethyltrifluoroborate can be used as the coupling partner in Suzuki–Miyaura reactions.^{6d} In that contribution, we reported the synthesis and cross-coupling reactions of potassium Boc-protected aminomethyltrifluoroborate **2**. Potassium Boc-protected aminomethyltrifluoroborate, which is a primary aminomethyl equivalent, was synthesized through a ‘one-pot’ synthesis in good yield and is now commercially available. Primary aminomethylarenes are readily available using this method after deprotection of the Boc group. The Boc protecting group is known to be easier to deprotect, compared to Ts or Phth groups, in acidic or even basic conditions.⁸ Therefore, using the Boc group is a more general approach to the primary amines.

Scheme 2



All of the previous reports employed aryl and hetaryl halides as electrophilic partners in cross-coupling reactions. As alternative coupling partners, aryl and hetaryl sulfonate derivatives have been utilized in Suzuki–Miyaura couplings.^{9–11} Sulfonate groups are generally easy to handle and are derived from a complementary set of starting materials. Mesylates are of special interest in terms of atom economy, low cost, and stability, even though they show the lowest reactivity among sulfonate derivatives.¹⁰ Recently, our group has demonstrated the feasibility of Suzuki–Miyaura cross-coupling reactions of aryl and hetaryl mesylates with tertiary ammoniomethyltrifluoroborates and amidomethyltrifluoroborates.^{11c} To extend the scope of this transformation, we investigated the Suzuki–Miyaura cross-coupling reaction of

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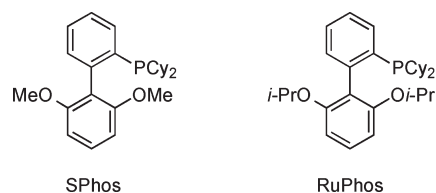


Figure 1. SPhos and RuPhos.

Table 1. Cross-Coupling of Aminomethyltrifluoroborate **2** with Various Electron-Neutral and Electron-Rich Aryl Mesylates

$\text{Ar-OMs} + \text{KF}_3\text{B-CH}_2\text{-NH-Boc} \xrightarrow{[\text{Pd}]^a} \text{Ar-CH}_2\text{-NH-Boc}$ <div style="display: flex; justify-content: space-around; width: 100%;"> 1a-h 2 3a-h </div>				
entry	product	ligand	isolated yield (%)	
1		3a RuPhos	79 (87) ^b	
2		3b RuPhos	66	
3		3c RuPhos	72	
4		3d RuPhos	72	
5		3e RuPhos	22	
6		3f RuPhos	86	
7		3g RuPhos	59	
8		3h RuPhos	84	

^a Reaction conditions: 1.0 equiv of aryl mesylate, 1.1 equiv of trifluoroborate, 5 mol % of $\text{PdCl}_2(\text{cod})$, 10 mol % of ligand, 7 equiv of K_3PO_4 , *t*-BuOH/ H_2O (1:1, 0.2 M), 95 °C, 22 h. ^b 4.0 mmol of mesylate, 3 mol % of $\text{PdCl}_2(\text{cod})$, 6 mol % of RuPhos.

potassium Boc-protected primary aminomethyltrifluoroborate with various aryl and hetaryl mesylates, providing an alternative entry to primary aminomethyl-substituted aromatics.

Initially, when the optimal conditions [$\text{Pd}(\text{OAc})_2$, SPhos or XPhos, K_2CO_3 , and toluene/ H_2O , 85 °C, 22 h] for aryl and hetaryl chlorides with potassium Boc-protected aminomethyltrifluoroborate **2** were applied to mesylates,

Table 2. Cross-Coupling of Aminomethyltrifluoroborate **2** with Various Electron-Poor Aryl Mesylates

$\text{Ar-OMs} + \text{KF}_3\text{B}-\text{CH}_2\text{NHBoc} \xrightarrow{[\text{Pd}]^a} \text{Ar}-\text{CH}_2\text{NHBoc}$ <div> <div>1i-p</div> <div>2</div> <div>3i-p</div> </div>				
entry	product	ligand	isolated yield (%)	
1		3i	SPhos	72
2		3j	RuPhos	46
3		3k	SPhos	42
4		3l	RuPhos	83
5		3m	RuPhos	82
6		3n	SPhos	80
7		3o	RuPhos	81
8		3p	RuPhos	86

^a Reaction conditions: 1.0 equiv of aryl mesylate, 1.1 equiv of trifluoroborate, 5 mol % of PdCl₂(cod), 10 mol % of ligand, 7 equiv of K₃PO₄, *t*-BuOH/H₂O (1:1, 0.2 M), 95 °C, 22 h.

only trace amounts of products were obtained.^{6d} Keeping these results in mind, we screened the Suzuki–Miyaura cross-coupling with the mesylated 1-naphthol **1a** based on the conditions related to C–O activation with potassium organotrifluoroborates reported previously.¹¹ Potassium phosphate tribasic was chosen as a base in a mixture of *t*-BuOH/H₂O. The coupling reactions were screened with different sources of palladium catalysts and ligands. Moreover, the reaction concentrations, ratio of two solvents, and temperature were studied extensively. After the optimization process, the combination of 1 equiv of mesylate, 1.1 equiv of trifluoroborate, 5 mol % of PdCl₂(cod), 10 mol % of SPhos or RuPhos (Figure 1), and 7 equiv of K₃PO₄ in *t*-BuOH/H₂O (1:1, 0.2 M) at 95 °C for 22 h turned out to be the best reaction conditions. Two different ligands were used because neither was general across the entire range of substrates.

With these optimized conditions in hand, we first studied the scope of the coupling reactions with various aryl mesylates (Tables 1 and 2).

Electrophiles with both electron-neutral and electron-rich substituents on the aryl rings represented good coupling partners in the desired reactions (Table 1). The reactions revealed that RuPhos was the most efficient ligand for all substrates containing electron-neutral and

Table 3. Cross-Coupling of Aminomethyltrifluoroborate **2** with Various Hetaryl Mesylates

$\text{HetAr-OMs} + \text{KF}_3\text{B}-\text{CH}_2\text{NHBoc} \xrightarrow{[\text{Pd}]^a} \text{HetAr}-\text{CH}_2\text{NHBoc}$ <div> <div>4a-h</div> <div>2</div> <div>5a-h</div> </div>				
entry	product	ligand	isolated yield (%)	
1		5a	SPhos	52
2		5b	RuPhos	78
3		5c	SPhos	84
4		5d	RuPhos	87
5		5e	SPhos	57
6		5f	SPhos	75
7		5g	RuPhos	86
8		5h	RuPhos	88

^a Reaction conditions: 1.0 equiv of hetaryl mesylate, 1.1 equiv of trifluoroborate, 5 mol % of PdCl₂(cod), 10 mol % of ligand, 7 equiv of K₃PO₄, *t*-BuOH/H₂O (1:1, 0.2 M), 95 °C, 22 h.

electron-donating groups on the aryl ring. With more sterically demanding substituents *ortho* to the mesylate group, the coupling yields dropped dramatically (Table 1, entries 4 and 5). The more sterically encumbered di-*ortho* substituted electrophile gave only a 22% isolated yield, perhaps because of a slow oxidative addition step (Table 1, entry 5).¹² The reactions were also successful with electron-donating groups on the aryl ring (Table 1, entries 7 and 8). By increasing the reaction scale to 4 mmol of mesylated 1-naphthol **1a**, the reaction could be carried out with a lower catalyst loading [3 mol % of PdCl₂(cod), and 6 mol % of RuPhos] to obtain the corresponding product **3a** with an 87% isolated yield (Table 1, entry 1).

We then investigated electron-poor aryl mesylates as electrophilic coupling partners (Table 2). All electron-deficient aryl mesylates gave the desired products **3i–p** in moderate to good yields. In these cases, two different ligands (RuPhos and SPhos) were utilized to obtain the products, the yields of which depended on the nature of the

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functional groups and ligand utilized. As shown, a wide variety of functional groups, such as nitriles, aldehydes, esters, and ketones, were compatible with the reaction conditions (Table 2). However, lower yields were observed with the aldehyde and methyl ester substituents on the aryl ring compared to other substrates (Table 2, entries 2 and 3).

We next expanded the array of electrophiles to hetaryl mesylates (Table 3). Various hetaryl mesylates were coupled with potassium Boc-protected aminomethyltrifluoroborate **2** in moderate to good yields. Again, two ligands (RuPhos and SPhos) were required to give better results depending on the nature of the hetaryl coupling partners. Nitrogen-containing hetaryl mesylates, such as pyridine, quinoline, isoquinoline, indole, and thiazole, all provided the expected products in good yields (Table 3, entries 1–7). Interestingly, indole was successfully coupled without any protecting group in a 75% isolated yield (Table 3, entry 6). Moreover, sulfur-containing hetaryls also proved to be good coupling partners under the set of reaction conditions developed (Table 3, entries 7 and 8).

In summary, we have shown that potassium Boc-protected aminomethyltrifluoroborate, an equivalent of a primary aminomethyl subunit, was a good coupling partner in Suzuki–Miyaura cross-coupling reactions with aryl and hetaryl mesylates. A broad array of electrophiles, such as functionalized aryl mesylates and hetaryl mesylates, were coupled efficiently in standard coupling reactions. Further efforts to expand the scope of aminomethyl moieties are currently under study.

Acknowledgment. We thank the National Institutes of Health (NIGMS R01 81376) for their support of this research. We acknowledge Dr. Rakesh Kohli (University of Pennsylvania) for obtaining HRMS data.

Supporting Information Available. Experimental procedures and spectral data of all compounds synthesized. This material is available free of charge via the Internet at <http://pubs.acs.org>.

The authors declare no competing financial interest.