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Copper-Catalyzed Oxidative Cross-Coupling of Electron-Deficient Polyfluorophenylboronate Esters with Terminal Alkynes

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Abstract: We report herein a mild procedure for the copper-catalyzed oxidative cross-coupling of electron-deficient polyfluorophenylboronate esters with terminal alkynes. This method displays good functional group tolerance and broad substrate scope, generating cross-coupled alkynyl(fluoro)arene products in moderate to excellent yields. Thus, it represents a simple and alternative to the conventional Sonogashira reaction.

Introduction

Functionalized aryl and heteroaryl alkynes are powerful building blocks in chemical synthesis because of their versatility to be transformed into various useful molecules and also their ubiquity in natural product synthesis, pharmaceuticals, and advanced materials.^[11] Consequently, much effort has been expended to develop efficient methods to install various alkynyl groups. Some of the strategies which have been established include: (1) Sonogashira palladium/copper-catalyzed sp²-sp cross-coupling of aryl halides with terminal alkynes;^[2] (2) direct alkynylation of unreactive alkyl and aryl C-H bonds with prefunctionalized alkynating reagents such as alkynyl halides^[3] and hypervalent iodine reagents;^[4] (3) alkynylation of tetra-and penta-fluoroarenes and heteroarenes via C-H bond activation;^[5,6] and (4) cross-coupling of copper(I) acetylides with aryl halides, known as the Castro–Stephens reaction.^[7,8,9] However, some drawbacks remain, such as the use of precious metal catalysts including those of Pd,^[2] Rh,^[4a,b,h] and Au,^[4c,d] strategies that depend on the use of alkynyl halides or hypervalent iodine reagents, which are less readily available than the corresponding terminal alkynes, and the fact that copper(I) acetylides can be heat and shock sensitive when isolated.

It is generally acknowledged that polyfluoroarenes are important fluorinated aromatic cores and key structural units for various organic molecules, such as pharmaceuticals, agrochemicals and organic materials.^[10] The development of efficient methods to introduce fluorine or fluorinated building blocks into organic molecules has been the subject of intense research. Under certain conditions, Sonogashira cross-couplings involving highly fluorinated aryl halides can be problematic, giving low yields^[11a] and side reactions, i.e., hydrodehalogenation accompanied by homocoupling of the terminal alkyne.^[11b] The latter problem seems to arise from the slow reductive elimination of the fluoroaryl alkyne from Pd(II), which leads to competing reverse transmetallation processes, i.e., transfer of aryl groups from Pd to Cu in exchange for a second alkynyl moiety being transferred from Cu to Pd. Thus, an alternative approach would be useful. In 2010, Su and co-workers demonstrated the direct functionalization of polyfluoroarene C-H bonds with terminal alkynes, which has proven to be a viable method to generate the corresponding alkynylated products (Scheme 1a),^[12] but this reaction is limited to C_6F_5H or $4-RC_6F_4H$ substrates. Soon after, the

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oxidative alkynylation of azoles containing acidic C-H bonds with terminal alkynes was reported by the groups of Miura,^[13] Chang,^[14] and others.^[15] Recently, Su and co-workers reported a palladium-catalyzed alkynylation of heterocyclic substrates such as thiophenes and furans.^[16] Although these achievements were promising, they were restricted by elevated temperatures (>90 °C) and limited substrate scope. In 2003, the palladium-catalyzed oxidative cross-coupling of terminal alkynes with arylboronic acids was first disclosed by Zou and co-workers (Scheme 1b).^[17] In the past few years, various modifications of this Pd-catalyzed reaction have been developed.^[18] However, palladium is costly and only a few electron-withdrawing substituents on the aromatic ring of arylboronic acids were employed. Recently, Cheng *et al.* disclosed a copper-catalyzed oxidative coupling of arylboronic acids with terminal alkynes.^[19] However, the reported method sufferers from some disadvantages including high reaction temperature, long reaction time (36 h), and only moderate yields. From a synthetic point of view, the development of an improved procedure employing an inexpensive catalyst for widespread application has remained a highly desirable goal.

We reported the C-F borylation of fluoroarenes using a NHC (N-heterocyclic carbene)-ligated Ni complex as a catalyst to generate fluorinated arylboronic acid pinacol esters (Ar_FBpin) in good to excellent yields.^[20a,b] Very recently, we reported optimized conditions for the Suzuki-Miyaura cross-coupling of Ar_FBpin with aryl iodides and bromides using a combination of CuI and phenanthroline as a catalyst precursor to generate cross-coupled products in moderate to excellent yields.^[20e] We have recently reported the palladium-catalyzed homocoupling of fluorinated arylboronates,^[20d] and the borylation of aryl chlorides, using NHC-stabilized nickel(0) complexes^[20e] or a readily prepared NHC-stabilized Cu catalyst.^[20f] Inspired by these results, we attempted to develop a Cu-catalyst system for the oxidative cross-coupling of Ar_FBpin compounds with terminal alkynes.

Scheme 1. Selected Oxidative Cross-Coupling Reactions of Alkynes

a) Polyfluoroarenes with terminal alkynes^[12]



b) Arylboronic acids with terminal alkynes^[17,18]

R'-B(OH)₂ + H───R ──[Pd], base, oxidant R'───R

c) This work: Polyfluorophenylboronate esters with terminal alkynes



Results and Discussion

We initially investigated the cross-coupling reaction with model substrates pentafluorophenyl-Bpin (1a) and phenylacetylene (2a), using Ag₂O as the oxidant and phenanthroline (Phen) as the ligand. During our initial experiments, no reaction occurred when CuBr₂ was employed as the metal source, with *t*BuOLi as the base in DMF solution (Table 1, entry 1). However, employing CuCl as catalyst precursor gave rise to compound **3a** in 10% yield (Table 1, entry 2). The introduction of Cu(OAc)₂ as the catalyst precursor improved the yield to 18% (Table 1, entry 3). However, large amounts of diyne byproduct **4** and perfluorobiphenyl compound **5** were produced.

We speculated that strong bases, such as tBuOLi, might accelerate the formation of 5. Under otherwise identical conditions, replacing the strong base with K₃PO₄ effectively inhibited the homocoupling of pentafluorophenylBpin (Table 1, entry 4). To our surprise, the addition of 2,3-dichloro-5,6-dicyanobenzoquinone (DDQ) significantly improved the yield to 58% and suppressed the formation of **4** (Table 1, entry 5). It is possible that DDQ serves as an electron-transfer mediator.^[12,21] To optimize the reaction performance, we screened the reaction parameters, including the base and the solvent. Of the bases examined, K₂CO₃ proved to be the most effective (entry 7). Both KF and Cs₂CO₃ gave significantly lower yields (entries 6 and 8). In addition, reaction optimization also revealed that the solvent had a significant impact on this reaction. Lower yields were observed when reactions were performed in other solvents such as 1,2-dichloroethane (DCE), CH₃CN, THF, DMSO, MTBE, and toluene (entries 9-14). Notably, the replacement of Ag₂O with O₂ failed to give any desired product (entry17), indicating the unique roles of Ag₂O in promoting this reaction. Attempts to run the reaction in air resulted in a very low yield of the desired product (entry 15). Reducing the amount of K₂CO₃ and Ag₂O also diminished the yield (Table 1, entries 18 and 19).

$F \xrightarrow{F} Bpin + H \xrightarrow{Ph} Ph \frac{[Cu], Phen, base}{additive, Ag_2O, solvent} F \xrightarrow{F} F \xrightarrow{Ph + Ph} Ph + Ph \xrightarrow{Ph} + C_6F_5 - C_6F_5$							
F F 40 °C,12 h			F F		4	5	
Entry	[Cu] Base Additive		<u>u</u>	Vield(%) ^[b]			
	(15 mol%)	(2 equiv)	(40 mol%)	Solvent	3a	4	5
1	CuBr ₂	tBuOLi	-	DMF	0	65	36
2	CuCl	<i>t</i> BuOLi	-	DMF	10	55	35
3	Cu(OAc) ₂	<i>t</i> BuOLi	-	DMF	18	60	35
4	Cu(OAc) ₂	K ₃ PO ₄	-	DMF	35	52	8
5	Cu(OAc) ₂	K ₃ PO ₄	DDQ	DMF	58	5	8
6	Cu(OAc) ₂	Cs_2CO_3	DDQ	DMF	11	25	25
7	Cu(OAc) ₂	K ₂ CO ₃	DDQ	DMF	82	3	4
8	$Cu(OAc)_2$	KF	DDQ	DMF	15	10	45
9	$Cu(OAc)_2$	K_2CO_3	DDQ	MTBE	0	5	35
10	Cu(OAc) ₂	K_2CO_3	DDQ	DCE	0	5	0
11	$Cu(OAc)_2$	K_2CO_3	DDQ	toluene	0	10	10
12	Cu(OAc) ₂	K_2CO_3	DDQ	DMSO	25	15	20
13	$Cu(OAc)_2$	K_2CO_3	DDQ	CH ₃ CN	10	15	10
14	$Cu(OAc)_2$	K_2CO_3	DDQ	THF	5	10	15
15 ^{[c],}	Cu(OAc) ₂	K_2CO_3	DDQ	DMF	5	5	40
16 ^[d]	$Cu(OAc)_2$	K_2CO_3	DDQ	DMF	5	10	35
17 ^[e,f]	$Cu(OAc)_2$	K_2CO_3	DDQ	DMF	0	6	10
18 ^[g]	$Cu(OAc)_2$	K_2CO_3	DDQ	DMF	35	5	30
19 ^[h]	$Cu(OAc)_2$	K_2CO_3	DDQ	DMF	25	28	30

Table 1. Optimization of Reaction Conditions^[a]

[a] Reaction conditions: 1a (0.4 mmol), 2a (0.45 mmol), Cu(OAc)₂ (15 mol%), phenanthroline (Phen, 15 mol%), Ag₂O (1.8 equiv), DDQ (40 mol%), base (2.0 equiv), anhydrous and degassed solvent (5 mL). The mixture was stirred at 40 °C under argon, in a sealed tube for 12 h. [b] **3a**: isolated yield, **4**: isolated yield, **5**: the yields were determined by GC-MS analysis vs. a calibrated internal standard (n-dodecane) and are averages of two runs; [c] The reaction was performed in air. [d] Room temperature. [e] In the absence of Ag₂O. [f] Under O₂. [g] Ag₂O (1.2 equiv.). [h] Base (1.0 equiv.).

With the optimized conditions in hand, we focused our attention on investigating the scope and limitations of the oxidative cross-coupling reaction. As shown in Scheme 2, various fluorophenylboronate esters 1 containing 1-4 fluorine atoms were tested. Under the standard different tetrafluorophenylboronate conditions (Table 1, entry 7), esters and trifluorophenylboronate esters smoothly underwent alkynylation giving good to excellent

yields (Scheme 2, **3b-3f**). However, these reaction conditions were not suitable for Ar_FBpin substrates containing di- or mono-fluorinated arylboronates such as 2,5- or 2,3-difluorophenyl-Bpin (**1g** and **1i**) and 3-fluorophenyl-Bpin (**1h**), perhaps due to the lower Lewis acidity of the boronates which is impacted by the number fluorine and, especially, *ortho*-fluorine substituents. We speculated that increasing the temperature might be crucial for overcoming the barrier to C-B bond activation and thus to obtaining efficient catalysis. When reactions were performed at 80 °C, the corresponding products **3g** and **3i** were formed in good yields. It also noteworthy that replacement of the weak base with a stronger base afforded the corresponding product in good yield (**3h**).

Scheme 2. Scope of the Reaction with Respect to the Different Polyfluorophenyl Boronate Substrates 1 ^[a,b]



[a] Reaction conditions: **1** (0.4 mmol), **2a** (0.45 mmol), Cu(OAc)₂ (15 mol %), Phen (15 mol %), Ag₂O (1.2 equiv.), DDQ (40 mol %), K₂CO₃ (2.0 equiv.), DMF (4 mL), 40 °C, 12 h, Ar. [b] **3**: isolated yield. [c] Ag₂O (1.8 equiv.). [d] 80 °C. [e] *t*BuOLi.

Scheme 3. Scope of the Reaction with Respect to the Different Terminal Alkyne Substrates 2^[a,b]



[a] Reaction conditions: **1a** (0.4 mmol), **2** (0.45 mmol), Cu(OAc)₂ (15 mol %), Phen (15 mol %), Ag₂O (1.8 equiv), DDQ (40 mol %), K₂CO₃ (2.0 equiv.), DMF (4 mL), 40 °C, 12 h, Ar. [b] **6**: isolated yield. [c] 24 h.

The substituents of alkynes 2 were then varied, in order to explore further the scope of the reaction. As shown in Scheme 3, a series of alkynes 2 with different electron-withdrawing and electron-donating substituents on the aromatic ring were subjected to the optimal conditions. The experimental results showed that a broad range of substituents on the arylalkynes 2, including methyl, methoxy, chloro, bromo, and fluoro groups at the *ortho-*, *meta-*, and *para-*positions of the aromatic ring were well tolerated, providing the desired compounds in moderate to excellent yields (Scheme 3, **6a-6h**). Furthermore, the structures of compounds **6a** and **6g** were unambiguously confirmed *via* single crystal X-ray diffraction (*vide infra*). An ester group, which may not be tolerated in reactions employing organozinc reagents, is also compatible with this reaction (**6i**). Importantly, aliphatic alkynes proceeded to give the desired products in moderate to good yields (**6j** and **6k**). With a highly electron-withdrawing CF₃-substituent, only moderate yields were observed (**6l** and **6m**). Unfortunately, less reactive 4-nitro-phenyl and 4-cyano-phenyl alkynes were not suitable for the reaction under the standard conditions.

To examine the feasibility of scaling up the reaction, a gram-scale coupling of C_6F_5 -Bpin with phenylacetylene was employed (Scheme 4). The desired coupling product was obtained with minimal loss of yield (72%).

Scheme 4: Gram-Scale Synthesis of 3a



Based on previous reports,^[22] and the aforementioned observations, a plausible catalytic cycle for our oxidative cross-coupling reaction is shown in Scheme 5. The first step would involve the addition of alkynyl anion leading to the formation of alkynylcopper(II) species **B**. Subsequent transmetalation between Ar_FBpin and intermediate **B** occurs to form intermediate

C. The desired product **3a** would be generated by C-C reductive elimination. The Cu(0) species formed is reoxidized by DDQ, *vide supra*, ^[12,21] to regenerate **A**, completing the catalytic cycle.

Scheme 5: Proposed Mechanism



Molecular and Crystal Structures: Intermolecular $\pi \cdots \pi$ Stacking Interactions

The crystal structures of the cross-coupling products **6a** and **6g** were analyzed using single-crystal X-ray diffraction. The molecular geometries of these compounds in their crystal structures are shown in Figure 1. The central C=C bond lengths are 1.195(2) and 1.1996(6) Å (Table 2) and, hence, typical of C=C triple bonds (1.192 Å).^[23] The sp-sp² C–C single bonds between the alkyne and the fully fluorinated phenyl rings are slightly shorter (1.4265(7) and 1.427(2) Å) than the corresponding bonds to the mesityl ring of **6a** (1.4350(7) Å) or the para mono-fluorinated phenyl ring of 6g (1.437(2) Å). The sp-sp² C-C bonds to tetra- or penta-fluorinated phenyl rings are also shorter than those to the fully or mostly hydrogen-containing phenyl rings of mixed compounds in other partially fluorinated tolans,^[24] rod-like 1,4-bis(phenylethynyl)benzenes,^[25] and phenyl and perfluorophenyl end-capped polyynes.^[26] The shortening is due to the strong electron-withdrawing nature of the fluorine atoms, and the length difference is also observed in the co-crystals of fully hydrogen-containing fully fluorinated tolans.[24] rod-like and 1,4-bis(phenylethynyl)benzenes,^[27] and phenyl end-capped polyynes.^[26] The molecules of **6a**

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and **6g** are nearly planar with a very small twist between the aryl moieties (2.959(3)) and $3.04(5)^{\circ}$, Table 2). A small twist angle of between 0 and 6° is also typical of the hydrogenated and fluorinated tolans, rod-like 1,4-bis(phenylethynyl)benzenes, and phenyl endcapped polyynes.^[24-27] Larger twist angles were reported for compounds related to **6g** in which the fluorine atom at the *para*-position of the phenyl ring is substituted by iodine $(9.4(2)^{\circ})$, bromine $(15.69(8)^{\circ})$, and NO₂ $(9.90(7)^{\circ})$.^[28] This may be related to the prevalence of different intermolecular interactions in these compounds (see below).



Figure 1. Solid-state molecular structures of **6a** and **6g** determined by single-crystal X-ray diffraction at 100 K. Ellipsoids are drawn at the 50% probability level, and H atoms are omitted for clarity. Colors: white (carbon), green (fluorine).

Table 2. Selected bond lengths (Å) and angles (°) of **6a** and **6g**, and $\pi \cdots \pi$ stacking distances (Å).

	6a	6g	
C≡C triple bond	1.1996(6)	1.195(2)	
$C_{Aryl(H/F)} - C_{triple}$	1.4350(7)	1.437(2)	
$C_{Aryl(F)} - C_{triple}$	1.4265(7)	1.427(2)	
\angle Aryl(F)-Aryl(H/F)	2.959(3)	3.04(5)	
anntusid sentusid distance	3.586(3)	3.705(3)	
centroid-centroid distance	3.629(3)	3.913(3)	
internloner concretion	3.361(3) / 3.424(3)	3.349(2) / 3.415(2)	
interplanar separation	3.325(3) / 3.376(3)	3.379(2) / 3.438(2)	
offoot obifula	1.248(3) / 1.064(2)	1.586(3) / 1.439(3)	
onset smilt ¹⁴	1.455(2) / 1.332(2)	1.975(3) / 1.868(3)	

[a] The offset shift, also called inter-centroid shift, is the distance within a plane of an aryl ring between the centroid of the respective aryl ring and the intersection point with the normal to the plane through the centroid of the other aryl ring.

In compounds **6a** and **6g**, the nearly planar molecules are related by inversion symmetry and are oriented offset face-to-face in a head-to-tail fashion forming infinite π -stacks (Figure 2). The interplanar separations between the aromatic rings (3.325(3) – 3.438(2) Å, Table 2) are in the normal range of π - π stacking interactions, which are typical of molecules for which the packing is dominated by arene-perfluoroarene interactions. The differences in electronegativity of hydrogen and fluorine atoms with respect to the carbon atoms leads to the formation of opposite multipoles for fully fluorinated and nonfluorinated aryl groups and, hence, to attractive multipole forces between these groups.^[29] Head-to-tail stacking *via* arene-perfluoroarene interactions, analogously to that observed in **6a** and **6g**, is commonly found in self-complementary compounds that contain both fluorinated and fluorinated tolans^[24] nonfluorinated aryl groups. Examples are partially and phenyl-endcapped polyynes,^[26] but also co-crystals of bis(phenylethynyl)benzenes with inversely alternating fluorinated and nonfluorinated phenyl rings.^[25] We conclude that methylation at the 2-, 4-, and 6-positions of the phenyl ring in 6a does not alter this common stacking motif and, hence, the influence of arene-perfluoroarene interaction on the molecular packing. Arene-perfluoroarene π -stacking was also observed in the 1:1 co-crystal of mesitylene and hexafluorobenzene.^[30] Weak intermolecular C-H...F, C...F, and F...F interactions exist between adjacent stacks in **6a** and **6g** (Figure 2, Table S2 in the Supporting Information). Mono-fluorination at the *para*-position of the phenyl ring in **6g** does not have a significant influence on the arene-perfluoroarene packing, which is very similar to that of 1-pentafluorophenyl-2-phenylacetylene.^[24] This was expected as the mono-chlorination of partially fluorinated tolan at the same *para* position did not alter the packing motif.^[28a] The effect of halogenation with chlorine, bromine, and iodine atoms at the para-positions of partially fluorinated tolans on the presence of arene-perfluoroarene interaction, studied earlier by Marder and co-workers,^[28a] revealed the absence of arene-perfluoroarene stacking only for the compounds substituted with the heavier halogens (Br, I). This was explained by the prevalence of Br...Br and I...I interactions determining the packing of the molecules.^[28a] Also note the larger twist angle between the phenyl rings in these compounds (15.69(8) and 9.4(2)°) when compared to those in arene-perfluoroarene π -stacked tolans (see discussion above). Similarly, the substitution of other strong electron-withdrawing groups such as NO_2

and CN at the *para*-position of the phenyl ring in partially fluorinated tolans showed the prevalence of $O \cdots O$ and $C-H \cdots N$ interactions and the absence of arene-perfluoroarene interactions in their crystal structures.^[28b]



Figure 2. Crystal structures of (left) **6a** and (right) **6g** projected along (top) the stacking direction of the molecules, and (bottom) the *b* and *a* axis, respectively, at 100 K. Molecules are π -stacked along the *a* axis (**6a**) and the *b* axis (**6g**), respectively, in alternating orientations. Four unit cells are shown in each projection. All ellipsoids are drawn at the 50% probability level, and H atoms are omitted for clarity. Colors: white (carbon), green (fluorine). Red dotted lines represent intermolecular contacts which are shorter than the sum of the Van der Waals radii.

In conclusion, we have developed a copper-catalyzed method for the direct alkynylation of electron-deficient polyfluorophenylboronate esters with terminal alkynes. This reaction features broad functional group tolerance, mild reaction conditions, and simple operation. From a synthetic point of view, the present reaction has the potential to be applied widely in organic synthesis because many shelf-stable aryl and alkyl boronate esters are commercially available. The partially fluorinated tolans also display interesting

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fluoroarene-arene π -stacking interactions in the solid-state as demonstrated by single-crystal X-ray diffraction in two cases.

Crystallographic details

Crystal data collection and processing parameters are given in the Supporting Information. CCDC-2000968 (**6a**), 2000970 (**6g**) contain the supplementary crystallographic data for this paper. These data are provided free of charge by The Cambridge Crystallographic Data Centre.

Author Contributions

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Conflict of interest

The authors declare no conflict of interest.

Keywords: boronate ester · coupling reaction · fluorine · fluoroarene · Sonogashira reaction

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Graphical Abstract

BFF: Boron Helps Make Fluorinated Alkynes: We report a mild procedure for the synthesis of alkynyl(fluoro)arenes *via* copper-catalyzed oxidative cross-coupling of polyfluorophenylboronate esters with terminal alkynes.

Bpin + H R [Cu], ligand, base additive, oxidant -R n=1, 2, 3, 4, 5