Published in final edited form as:

J Am Chem Soc. 2010 October 13; 132(40): 14076–14078. doi:10.1021/ja107481a.

Palladium-Catalyzed Conversion of Aryl and Vinyl Triflates to Bromides and Chlorides

Xiaoqiang Shen, Alan M. Hyde, and Stephen L. Buchwald*

Department of Chemistry, Massachusetts Institute of Technology, Cambridge, Massachusetts 02139

Abstract

The palladium-catalyzed conversion of aryl and vinyl triflates to aryl and vinyl halides (bromides and chlorides) has been developed using dialkylbiaryl phosphine ligands. A variety of aryl, heteroaryl and vinyl halides can be prepared via this method in good to excellent yields.

Aryl halides are ubiquitous synthetic intermediates in organic chemistry that are used for numerous transformations.¹ They are also present in a wide variety of natural products, pharmaceuticals and agrochemicals.² Therefore, the development of general and regioselective methods for the preparation of functionalized aryl halides is of great importance. Sulfonate esters, often referred to as pseudo-halides, can be employed as alternatives to aryl halides in many cross-coupling reactions.³ Such pseudo-halides, however, cannot be used as precursors for the generation of free radicals⁴ or organometallic reagents.¹ Traditional methods for the synthesis of aryl halides (bromides or chlorides) from phenols require either forcing conditions⁵ or multi-step procedures.⁶ Herein, we report the first example of palladium-catalyzed direct conversion of readily available aryl sulfonate esters to aryl bromides and chlorides.

Although a catalytic method has not yet been reported, several mechanistically related metal-mediated carbon–halogen bond-forming processes have been published.⁷ For example, the reductive elimination of aryl halides from Pt(IV),⁸ Ni(III),⁹ Pd(IV),¹⁰ and Pd(III)¹¹ has been demonstrated. In addition, the reductive elimination of aryl halides from arylpalladium(II) halide complexes in the presence of a large excess of P(*t*-Bu)₃ has been reported.^{12,13} A number of copper¹⁴- and nickel¹⁵-mediated halide exchange reactions have also been developed in which carbon–halogen reductive elimination processes are proposed to take place.

Despite these advances in aryl–halide bond formation, to the best of our knowledge, there are few examples of the catalytic conversion of an aryl sulfonate to an aryl halide¹⁶ including the ruthenium-catalyzed conversion of 2-naphthyl triflate to 2-bromonaphthalene^{16b} and our recent report of the palladium-catalyzed conversion of aryl triflates to aryl fluorides.¹⁷ This prompted us to query whether we could uncover a catalytic process to convert aryl triflates into corresponding aryl bromides and chlorides; and results are disclosed herein.

Catalysts based on dialkylbiaryl monophosphines **1–4**, which have been successfully utilized as ligands in numerous transformations,^{17,18} were initially examined for the conversion of

sbuchwal@mit.edu.

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

4-*n*-butyl phenyltriflate into 4-*n*-butylbromobenzene (**5**). In initial experiments, these reactions were carried out in the presence of 2.0 equivalents of Bu₄NBr, 1 mol% Pd₂(dba)₃ and 3 mol% ligand in toluene at 100 °C. This set of conditions afforded only small amounts of aryl bromide products (1–6%, entries 1–4). Other bromide sources were tested but the results were unimpressive, with KBr providing the highest yield (11%, entry 5). Since potassium bromide has a low solubility in toluene, we reasoned that the use of a phase transfer catalyst (PTC) might be beneficial. Adding polyethylene glycols improved the yield to 32% (entry 6). All attempts to increase the yield of this reaction with more forcing conditions or higher catalyst loading failed. We were initially puzzled by this and wondered whether product inhibition was occurring. Therefore, we repeated the 4-*n*-butyl phenyltriflate to 4-*n*-butylbromobenzene reaction (eq. 1) in the presence of 1 equivalent of potassium triflate and found that the reaction was completely inhibited (entry 7). This was surprising since the triflate anion is known to be a poor nucleophile.¹⁹ In order to probe whether this result was due to the existence of an unfavorable equilibrium, **5** was heated under similar catalytic conditions with KOTf as the coupling partner. However, no conversion to aryl triflate was observed, ruling out this possibility. Our next efforts focused on the sequestration of KOTf as it was formed. After survey a broad range of Lewis acids, we found that the yield of the reaction could be increased to 50% when Et₃B was used as additive (entry 8). Full conversions were observed when commercially available *i*-Bu₂AlF or *i*-Bu₃Al was used; however, about 50–60% of the C–C coupling product, 1-*n*-butyl-4-*i*-butylbenzene, was formed in both cases (entries 9 and 10).²⁰ The use of the bulkier *i*-Pr₃Al additive prevented the formation of the C–C coupling product, although the desired aryl bromide was only formed in 25% yield with the reduction product, *n*-butylbenzene, now formed in 75% yield (entry 12). Fortunately, in situ formation of dialkylaluminum alkoxides by the addition of ketones or alcohols suppressed both C–C coupling and reduction byproducts; the yield was increased to 92% when 2-butanone was used (entry 13).

With the optimal conditions for this new transformation in hand, we next explored the reaction scope (Table 2). Both electron-poor and electron-rich aryl triflates are suitable substrates and provide aryl bromides in good yield. Significantly, heteroaryl triflates, such as quinolines, indoles, carbazoles and benzothiazoles can be efficiently converted to corresponding aryl bromides. Ester groups were well tolerated but can, in some cases, undergo unwanted transesterification. With 4-trifluoromethylsulfonyl ethylbenzoate as substrate, this undesired side reaction could be avoided by using EtOH as an additive in place of 2-butanone (Table 2, compound **8**). Aromatic amines are also tolerated in these transformations as exemplified by the formation of 7-bromo-1-naphthyl amine in moderate yield from the corresponding triflate (Table 2, compound **11**). In this case, the 2-butanone imine was initially formed, which was then cleaved after work up and flash chromatography. The bromination of 3-bromophenyl triflate and 4-chloro-1-naphthyl triflate demonstrates that polyhalogenated products may be efficiently formed. Vinyl triflates are also effective substrates for the bromination reaction. For example, a vinyl triflate derived from estrone is easily converted to the corresponding vinyl bromide in 92% yield (Table 2, compound **20**).

As highlighted in Table 3, aryl and vinyl triflates can also be converted to aryl and vinyl chlorides under similar reaction condition using KCl as the chloride source in good to excellent yields.²¹

In conclusion, a new method for the direct conversion of aryl and vinyl triflates to aryl and vinyl halides has been achieved using palladium catalysis. The use of sterically hindered dialkylbiaryl monophosphines is key for success in achieving this new transformation.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

Generous financial support from the National Institutes of Health (GM46059) is gratefully acknowledged.

References

1. For general reviews on halogen–metal exchange, see: (a) Knochel P, Dohle W, Gommermann N, Kneisel FF, Kopp F, Korn T, Sapountzis I, Vu VA. *Angew Chem Int Ed.* 2003; 42:4302. (b) Parham WE, Bradsher CK. *Acc Chem Res.* 1982; 15:300.
2. (a) Gribble GW. *Acc Chem Res.* 1998; 31:141. (b) Gribble GW. *Chem Soc Rev.* 1999; 28:335.
3. For a selected review, see: Ritter K. *Synthesis.* 1993:735.
4. Bunnett JF. *Acc Chem Res.* 1992; 25:2.
5. (a) Wiley GA, Hershkowitz RL, Rein BM, Chung BC. *J Am Chem Soc.* 1964; 86:964. (b) Bay E, Bak DA, Timony PE, Leone-Bay A. *J Org Chem.* 1990; 55:3415.
6. (a) Wulff WD, Peterson GA, Bauta WE, Chan K, Faron KL, Gilbertson SR, Kaesler RW, Yang DC, Murray CK. *J Org Chem.* 1986; 51:277. (b) Thompson ALS, Kabalka GW, Akula MR, Huffman JW. *Synthesis.* 2005:547. (c) Kang H, Facchetti A, Stern CL, Rheingold AL, Kassel WS, Marks TJ. *Org Lett.* 2005; 7:3721. [PubMed: 16092859] (d) Grunewald GL, Seim MR, Regier RC, Criscione KR. *Bioorg Med Chem.* 2007; 15:1298. [PubMed: 17126018]
7. For general reviews on metal-mediated carbon–halogen bond formation, see: (a) Vigalok A. *Chem—Eur J.* 2008; 14:5102. (b) Sheppard TD. *Org Biomol Chem.* 2009; 7:1043. [PubMed: 19262919]
8. (a) Yahav-Levi A, Goldberg I, Vigalok A. *J Am Chem Soc.* 2006; 128:8710. [PubMed: 16819847] (b) Yahav-Levi A, Goldberg I, Vigalok A, Vedernikov AN. *Chem Commun.* 2010; 46:3324.
9. Higgs AT, Zinn PJ, Simmons SJ, Sanford MS. *Organometallics.* 2009; 28:6142.
10. (a) Whitfield SR, Sanford MS. *J Am Chem Soc.* 2007; 129:15142. [PubMed: 18004863] (b) Furuya T, Ritter T. *J Am Chem Soc.* 2008; 130:10060. [PubMed: 18616246] (c) Ball ND, Sanford MS. *J Am Chem Soc.* 2009; 131:3796. [PubMed: 19249867] (d) Arnold PL, Sanford MS, Pearson SM. *J Am Chem Soc.* 2009; 131:13912. [PubMed: 19788324]
11. Power DC, Ritter T. *Nature Chem.* 2009; 1:302. [PubMed: 21500602]
12. (a) Roy AH, Hartwig JF. *J Am Chem Soc.* 2001; 123:1232. [PubMed: 11456679] (b) Roy AH, Hartwig JF. *J Am Chem Soc.* 2003; 125:13944. [PubMed: 14611215]
13. An example of reversible oxidative addition of palladium into 2-bromoindoles has appeared: Newman SG, Lautens M. *J Am Chem Soc.* 2010; 132:11416. [PubMed: 20681622]
14. (a) Bacon RGR, Hill HAO. *J Chem Soc.* 1964:1097. (b) Klapars A, Buchwald SL. *J Am Chem Soc.* 2002; 124:14844. [PubMed: 12475315]
15. (a) Takagi K, Hayama N, Okamoto T. *Chem Lett.* 1978; 7:191. (b) Tsou TT, Kochi JK. *J Org Chem.* 1980; 45:1930. (c) Arvela RK, Leadbetter NE. *Synlett.* 2003:1145.
16. For the Ni-catalyzed conversion of pyranil triflates to pyranil bromides, see: (a) Milne JE, Jarowicki K, Kocienski PJ. *Synlett.* 2002:607. For the Ru-catalyzed conversion of alkenyl triflates to alkenyl halides, see: (b) Shirakawa E, Imazaki Y, Hayashi T. *Chem Commun.* 2009:5088.
17. Watson DA, Su M, Teverovskiy G, Zhang Y, Garcia-Fortanet J, Kinzel T, Buchwald SL. *Science.* 2009; 325:1661. [PubMed: 19679769]
18. (a) Cho E, Senecal TD, Kinzel T, Zhang Y, Watson DA, Buchwald SL. *Science.* 2010; 328:1679. [PubMed: 20576888] (b) Fors BP, Watson DA, Biscoe MR, Buchwald SL. *J Am Chem Soc.* 2008; 130:13552. [PubMed: 18798626] (c) Ikawa T, Barder TE, Biscoe MR, Buchwald SL. *J Am Chem Soc.* 2007; 129:13001. [PubMed: 17918833] (d) Anderson KW, Ikawa T, Tundel RE, Buchwald SL. *J Am Chem Soc.* 2007; 128:10694. [PubMed: 16910660] (e) Surry DS, Buchwald SL. *Angew Chem Int Ed.* 2008; 47:6338. (f) Martin R, Buchwald SL. *Acc Chem Res.* 2008; 41:1461. [PubMed: 18620434]
19. (a) Lawrance GA. *Chem Rev.* 1986; 86:17. (b) Howells RD, McCown JD. *Chem Rev.* 1977; 77:69.

20. For examples of Pd-catalyzed cross-coupling of organoaluminum reagents, see: (a) Babu S, Negishi E. *J Am Chem Soc.* 1976; 98:6729. (b) Hirota K, Isobe Y, Maki Y. *J Chem Soc Perkin Trans.* 1989; 1:2513.
21. Note: About 5% of 1-*n*-butyl-4-iodobenzene was formed when 4-*n*-butyl phenyltriflate was subjected to similar catalytic condition using KI as the iodide source.

TABLE 1

Evaluation of Reaction Conditions for Bromination^a

(1)

Entry	MBr	Ligand	PTC	Additive	Yield (%) ^b
1	Bu ₄ NBr	1	---	---	1 ^c
2	Bu ₄ NBr	2	---	---	2 ^c
3	Bu ₄ NBr	3	---	---	3 ^c
4	Bu ₄ NBr	4	---	---	6 ^c
5	KBr	4	---	---	11 ^c
6	KBr	4	PEG	---	32 ^c
7	KBr	4	PEG	KOTf ^d	1 ^c
8	KBr	4	PEG	Et ₃ B	50 ^c
9	KBr	4	PEG	^t Bu ₂ AlF	54 ^e
10	KBr	4	PEG	^t Bu ₃ Al	41 ^e
11	KBr	4	PEG	(^t PrO) ₃ Al	1 ^c
12	KBr	4	PEG	^t Pr ₃ Al ^f	25 ^g
13	KBr	4	PEG	^t Bu ₃ Al/2-butanone ^h	92

R = Cy, BrettPhos (1)
R = ^tBu, ^tBuBrettPhos (4)

^aReaction conditions: 0.25 mmol aryl triflate, 2.0 equiv of MBr, 1.5 equiv of additive, 50 mg PEG3400: poly(ethylene glycol), $M_n \approx 3400$, toluene (1.0 mL).

^bYields determined by GC analysis with an internal standard of dodecane.

^cThe remaining material was the 4-*n*-butyl phenyltriflate.

^d1.0 equiv of KOTf.

^eThe remaining material was the 1-*n*-butyl-4-*i*-butylbenzene.

^fTriisopropylaluminum diethyletherate as additive.

^gThe remaining material was the *n*-butylbenzene.

^h1.5 equiv of *i*-Bu₃Al and 2-butanone as additives.

TABLE 2

Palladium-Catalyzed Conversion of Triflates to Bromides^a

^a Reaction conditions: 1.0 mmol triflate, 1.5–2.5 mol% of Pd₂(dba)₃, 3.75–6.25 mol% of **4** (**4** : Pd = 1.25 : 1), 1.5 mmol KBr, 1.5 mmol 2-butanone, 1.5 mL of *i*Bu₃Al (1M solution in toluene), 120 mg PEG3400, toluene (6.0–8.0 mL), 20–24 h; Isolated yields (average of two runs).

^b 1.25 equiv of *i*-Bu₃Al and EtOH as additives; contains ~5% of ethyl 4-*i*-butylbenzoate.

^c Contains 1,4-dichloronaphthalene and 1,4-dibromonaphthalene (4–10% total).

Table 3Palladium-Catalyzed Conversion of Triflates to Chlorides^a

^aReaction conditions: 1.0 mmol triflate, 1.5–2.5 mol% of Pd₂(dba)₃, 3.75–6.25 mol% of **4** (**4** : Pd = 1.25 : 1), 1.5 mmol KCl, 1.5 mmol 2-butanone, 1.5 mL of *i*Bu₃Al (1M solution in toluene), 120 mg PEG3400, toluene (6.0–8.0 mL), 20–24 h; Isolated yields (average of two runs).

^b1.25 equiv of *i*-Bu₃Al and EtOH as additives; contains ~5% of ethyl 4-*i*-butylbenzoate.