

# One-Pot, Three-Component Arylalkynyl Sulfone Synthesis

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**S** Supporting Information

**ABSTRACT:** A one-pot three-component protocol for the preparation of arylsulfonyl alkynes through the reaction of ethynyl-benziodoxolone (EBX) reagents, DABSO (DABCO·SO<sub>2</sub>), and either organomagnesium reagents or aryl iodides with a palladium catalyst is reported. A broad range of aryl and heteroarylalkynyl sulfones were obtained in 46–85% overall yield.



Aryl sulfone-containing compounds display a variety of biological activities. Several of them are marketed drugs for treatment of human diseases. Examples include the antimigraine Vioxx (1),<sup>1a</sup> the antibacterial Dapsone (2)<sup>1b</sup> and the antiandrogen Casodex (3)<sup>1c</sup> (Figure 1).

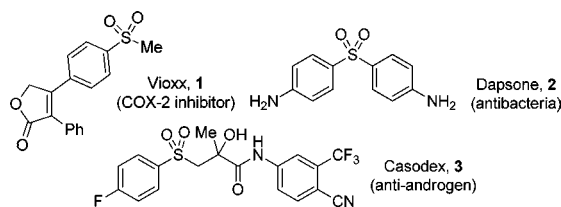


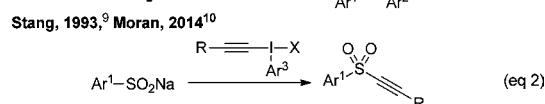
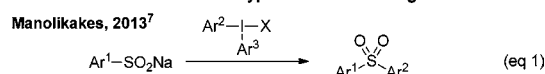
Figure 1. Biologically active aryl sulfones.

Among all sulfones, (aryl)alkynyl sulfones constitute an important class of building blocks due to their versatile reactivity. The sulfonyl unit has a strong electron-withdrawing character that enhances the reactivity of the triple bond.<sup>2</sup> Therefore, sulfonyl acetylenes are widely utilized in cyclo-additions and conjugate additions reactions. They were also found to react in certain cases with organometallic reagents or radicals through 1,3-addition to generate anions or radical intermediates, followed by an elimination of the sulfonyl group to give disubstituted alkynes.<sup>3</sup>

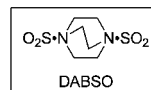
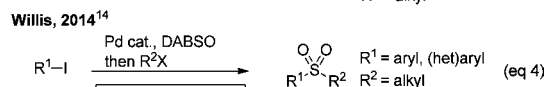
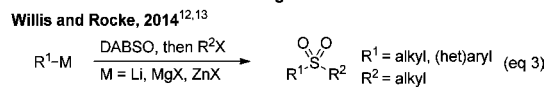
Up to now, the preparation of sulfones could be achieved by numerous approaches, the most frequent one being the oxidation of the corresponding thiols.<sup>4</sup> Another convenient method is based on the reaction of sulfinate salts with electrophiles.<sup>5</sup> If the introduction of an aryl or an alkynyl group is desired on the sulfinate, hypervalent iodine reagents have emerged as reagents of choice, due to their exceptional reactivity.<sup>6</sup> For example, Manolikakes and co-workers reported that diaryl sulfones could be synthesized from arylsulfinate and diaryliodonium salts (Scheme 1A, eq 1).<sup>7</sup> Electrophilic alkynyl iodonium salts are also widely used in alkynylation reactions,<sup>8</sup> and their utilization for the preparation of alkynyl sulfones from sulfinate salts were reported by the groups of Stang<sup>9</sup> and Moran (Scheme 1A, eq 2).<sup>10</sup> However, the scope of arylsulfinate salts

## Scheme 1. Recent Reports on Sulfone Synthesis

### A. Sulfones from sulfonates and hypervalent iodine reagents



### B. In situ formation of sulfonates using DABSO



reported was limited (only PhSO<sub>2</sub>Na and *p*-TolSO<sub>2</sub>Na), probably due to the low number of commercially available sulfinate salts. Recently, Willis and co-workers showed that DABSO (DABCO·SO<sub>2</sub>, the combination of DABCO and sulfur dioxide) can serve as a surrogate of SO<sub>2</sub> for the in situ formation of sulfinate salts in the synthesis of sulfonamides.<sup>11</sup> More importantly, Willis and co-workers<sup>12</sup> and Rocke and co-workers<sup>13</sup> independently demonstrated a one-pot synthesis of sulfones from organolithium/magnesium and organozinc reagents, DABSO, and alkyl halides (Scheme 1B, eq 3). In addition, transition metal-catalyzed, e.g. palladium<sup>14</sup> (eq 4) and gold,<sup>15</sup> as well as metal-free<sup>16</sup> sulfinate salt formation with DABSO were also reported for sulfone synthesis. These recent breakthroughs have greatly simplified the synthesis of sulfones. However, a one-pot protocol for the preparation of arylalkynyl sulfones has still not been reported. The synthesis of this class

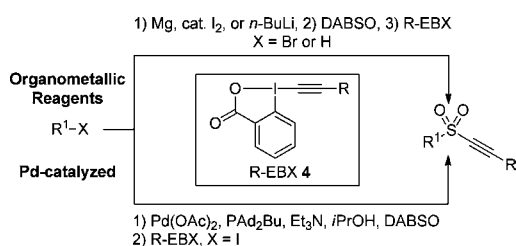
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of compounds is especially challenging, as the products themselves are also highly reactive.

Our group is interested in alkyne synthesis using electrophilic alkylation reagents, in particular ethynyl-benziodoxolones (EBX, **4**).<sup>17,18</sup> These cyclic hypervalent iodine reagents display high reactivity, but are more stable than the traditionally used alkynylidonium salts. We therefore thought they would be well-suited to develop the first one-pot three-component synthesis of alkynyl sulfones. Herein we would like to present the successful implementation of this approach using EBX reagents **4**, DABSO, and organomagnesium/lithium reagents or aryl iodides with a palladium catalyst (Scheme 2).

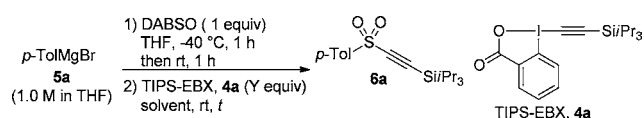
## Scheme 2. Our Approach toward the Synthesis of Alkynyl Sulfones



The preparation of sulfonyl alkynes starting from tolyl magnesium bromide (**5a**) with TIPS-EBX (**4a**) and DABSO was examined first. Initially, the protocol developed by Willis for preparation of sulfonamides<sup>11a</sup> was utilized. Unfortunately, the desired product was not obtained when using THF only as solvent. To our delight, 16% yield of **6a** was observed when THF was removed and replaced by DMF (Table 1, entry 1).

Longer reaction time (14 h), and DMF/H<sub>2</sub>O (*v/v* = 5/1) as a solvent system for the alkylation did not give better results (Table 1, entries 2–4). On the other hand, we were pleased to observe an increased yield of **6a** (65%) when THF was not removed before adding DMF for the alkylation step (Table

Table 1. Optimization of the Arylalkynyl Sulfone Synthesis



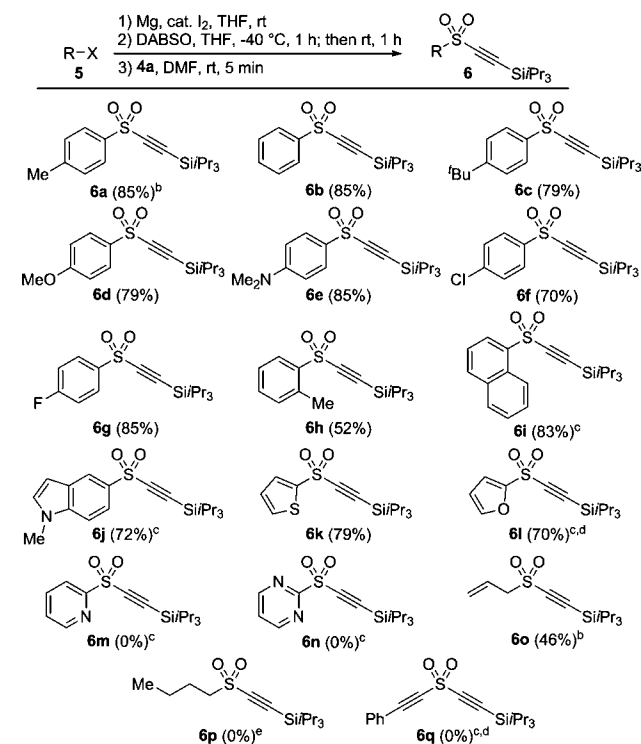
entry <sup>a</sup>	equiv of <b>4a</b>	solvent	t	yield (%) <sup>d</sup>
1	1.2	DMF <sup>b</sup>	2 h	16
2	1.2	DMF <sup>b</sup>	14 h	0
3	1.2	DMF/H <sub>2</sub> O <sup>b,c</sup>	2 h	15
4	1.2	DMF/H <sub>2</sub> O <sup>b,c</sup>	14 h	0
5	1.2	DMF	2 h	65
6	1.2	DMSO	2 h	0
7	1.5	DMF	2 h	50
8	1.5	DMF	1 h	50
9	1.5	DMF	30 min	75
10	1.5	DMF	5 min	78
11	1.2	DMF	5 min	80
12	1.1	DMF	5 min	75

<sup>a</sup>0.06 mmol *p*-tolylmagnesium bromide (**5a**) was used in 0.2 mL of THF. 0.2 mL of solvent was added for the second step (final concentration: 0.13 M). <sup>b</sup>THF was removed before adding 0.2 mL of solvent (final concentration: 0.3 M). <sup>c</sup>Ratio (*v/v*) = 5/1. <sup>d</sup>The yield was obtained based on <sup>1</sup>H NMR analysis using 1,3,5-trimethoxybenzene as internal reference.

1, entry 5). In contrast, the addition of DMSO was not successful (Table 1, entry 6). A fast examination of TIPS-EBX **4a** loading and reaction time (Table 1, entries 6–12) showed that 1.2 equiv of **4a** and 5 min reaction time resulted in the production of **6a** in 80% yield (Table 1, entry 11). The lower yields observed with longer reaction times are probably due to the high reactivity of the formed alkynyl sulfone **6a**.

With optimized conditions in hand we examined the scope of the one-pot sulfonylation alkylation from organometallic reagents with TIPS-EBX **4a** (Scheme 3). Acetylene **6a** was

## Scheme 3. Scope of the One-Pot Sulfonylation Alkylation<sup>a</sup>



<sup>a</sup>Reaction conditions: 0.20 mmol of **5** (1.0 equiv), 0.20 mmol of Mg (1.0 equiv), 0.20 mmol of DABSO (1.0 equiv), and THF (0.65 mL) were used for the first step. 0.24 mmol of TIPS-EBX **4a** (1.2 equiv) and DMF (0.65 mL) were added for the second step. Isolated yield after purification on column chromatography is given. <sup>b</sup>Commercial Grignard reagent was used. <sup>c</sup>0.5 M organometallic reagent in THF. <sup>d</sup>R-Li was generated from the corresponding C–H bond with *n*-BuLi. <sup>e</sup>Commercial *n*-BuLi was used.

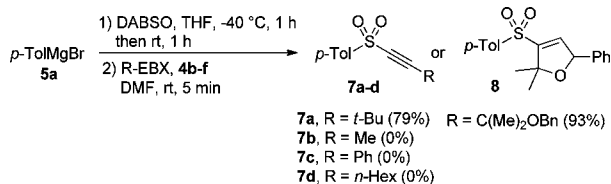
obtained in good yield on preparative scale (85%). We further expanded the utility of this protocol by preparing the organomagnesium/lithium reagents immediately before use. Phenylalkynyl sulfone **6b** could be synthesized in 85% overall yield starting from bromobenzene **5b**. Other electron-rich and electron-poor functional groups were well-tolerated. For example, *p*-*tert*-butyl, *p*-methoxy, and *p*-dimethylaminosulfonyl alkynes **6c**, **6d**, and **6e** were obtained in good yields (79, 79, and 85% respectively). *p*-Chloro and *p*-fluorophenylsulfonyl alkynes **6f**, and **6g** were also synthesized in 70 and 85% yield, respectively.

*o*-Tolyl sulfinate magnesium salt gave a lower yield of product **6h** (52%). In addition, alkyne **6i** was also obtained from 1-bromonaphthalene in 83% yield. The heteroaryl bromides 5-bromo-*N*-methylindole (**5j**) and 2-bromo-thiophene (**5k**) gave products **6j** and **6k** in 72 and 79% yield,

respectively. Sulfone **6l** was synthesized in 70% yield starting from furan **5l** using a selective lithiation at C2. However, 2-pyridinyl and 2-pyrimidinyl alkynyl sulfones **6m** and **6n** could not be synthesized using this one-pot protocol. Allylsulfonyl alkyne **6o** was obtained in 46% yield. Sulfones **6p** and **6q** could not be obtained when starting from the corresponding organolithium reagents.

Further extension of the scope of the one-pot sulfonylation alkylation was focused on *p*-tolyl Grignard (**5a**) with R-EBX reagents (Scheme 4). Alkyne **7a** was made in 79% yield using *t*-

#### Scheme 4. Iodine Reagent Scope of the One-Pot Sulfonylation Alkylation<sup>a</sup>



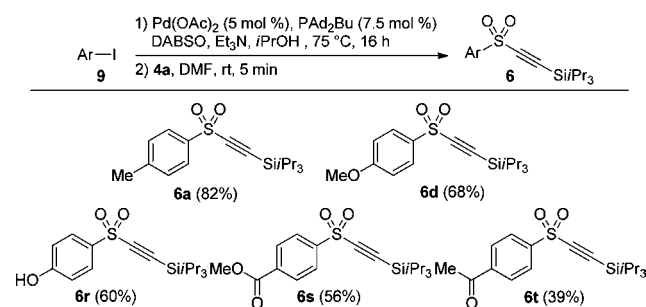
<sup>a</sup>Reaction conditions: 0.20 mmol of *p*-tolylMgBr (**5a**) (1.0 equiv), 0.20 mmol of DABSO (1.0 equiv) and THF (0.65 mL) were used for the first step. 0.24 mmol of R-EBX **4** (1.2 equiv) and DMF (0.65 mL) were added for the second step. Isolated yield after purification on column chromatography is given.

Bu-EBX **4b**.<sup>18f</sup> Unfortunately, Me-EBX **4c**,<sup>18k</sup> Ph-EBX **4d**,<sup>18f</sup> and *n*-Hex-EBX **4e**<sup>18f</sup> were not able to react with the in situ formed *p*-tolyl sulfinate salt to deliver the desired alkynes **7b**, **7c**, and **7d** respectively. Interestingly, dihydrofuran **8** was obtained in 93% yield when the EBX reagent **4f** was utilized in the alkylation.

At least three different mechanisms could be considered for the alkylation step in this one-pot protocol. A first possibility frequently occurring with hypervalent iodine reagents is direct nucleophilic attack of sulfinate onto the iodine atom of the benziodoxolone, followed by a C–S bond formation via a reductive elimination step.<sup>6</sup> Our group has recently discovered by computation an alternative mechanism for the alkylation of thiols involving a concerted three-atom transition state including the iodine, the sulfur and the  $\alpha$ -carbon atom of the alkyne.<sup>18k</sup> However, these two mechanisms are less likely to be involved in the alkylation of sulfonates because (i) dihydrofuran **8** was obtained and this product most probably results from 1,5-C–H insertion of a carbene intermediate, and (ii) EBX reagents **4c–f** did not give the desired alkyne products, in contrast to the high yields observed with thiols.<sup>18k</sup> These facts suggest that the reaction mechanism is different for sulfonates and most probably involves a third alternative: a conjugate addition of the sulfinate onto the  $\beta$ -alkynyl carbon of ethynyl-benziodoxolone, followed by an  $\alpha$ -elimination of the aryl iodide to give a carbene intermediate, and finally a 1,2-shift to form the alkyne.<sup>10,19</sup>

One disadvantage of the developed sulfonylation-alkynylation protocol involving organo-magnesium or -lithium reagents is that it cannot be applied to substrates sensitive to strong bases or nucleophiles. In order to further enhance the generality of the one-pot approach for the synthesis of alkynyl sulfones, we then examined a Pd-catalyzed ammonium sulfinate salt formation starting from aryl iodides which can proceed under much milder conditions (Scheme 1B, eq 4).<sup>14</sup> Alkynes **6a** (82%) and **6d** (68%) could be synthesized in comparable yields using the Pd-catalyzed sulfonylation in the first step (Scheme 5).

#### Scheme 5. Scope of the Pd-Catalyzed One-Pot Sulfonylation Alkylation<sup>a</sup>



<sup>a</sup>Reaction conditions: 0.20 mmol of ArI **9** (1.0 equiv), 10  $\mu$ mol of Pd(OAc)<sub>2</sub> (5 mol %), 15  $\mu$ mol of Pd<sub>2</sub>Bu (7.5 mol %), 0.20 mmol DABSO (1.0 equiv), Et<sub>3</sub>N (3.0 equiv) and *i*PrOH (1.7 mL) were used for the first step. 0.24 mmol of TIPS-EBX **4a** (1.2 equiv) and DMF (0.65 mL) were used for the second step. Isolated yield after purification on column chromatography is given.

Gratifyingly, alkynes **6r**, **6s** and **6t** bearing potentially base- and nucleophile sensitive hydroxy, methyl ester, and methyl ketone groups were synthesized successfully in 39–60% yield.

In conclusion, we report a simple one-pot protocol for sulfonylation-alkynylation starting either from organomagnesium/lithium reagents or from aryl iodides with a palladium catalyst, DABSO, and EBX reagents, in up to 85% overall yield. The method from organomagnesium/lithium reagents gives an unprecedented efficient access to aryl-alkynyl sulfones bearing benzene rings with electron-poor or electron-rich substituents, as well as heterocycles. The complementary Pd-catalyzed protocol can be used for substrates sensitive to the use of strongly basic or nucleophilic organometallic reagents. Extension of the scope and investigations on the mechanism of the sulfonyl alkylation are currently underway and will be reported in due course.

#### ■ ASSOCIATED CONTENT

##### Supporting Information

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

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##### Notes

The authors declare no competing financial interest.

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#### ■ REFERENCES

- (1) (a) Prasiat, P.; Wang, Z.; Brideau, C.; Chan, C. C.; Charleson, S.; Gauthier, J. Y.; Gordon, R.; Guay, J.; Gresser, M.; Kargman, S.; Kennedy, B.; Leblanc, Y.; Leger, S.; Mancini, J.; O'Neill, G. P.; Ouellet, M.; Percival, M. D.; Perrier, H.; Riendeau, D.; Rodger, I.; Tagari, P.; Therien, M.; Vickers, P.; Wong, E.; Xu, L. J.; Young, R. N.; Zamboni, R. *Bioorg. Med. Chem. Lett.* **1999**, *9*, 1773. (b) Artico, M.; Silvestri, R.; Pagnozzi, E.; Bruno, B.; Novellino, E.; Greco, G.; Massa, S.; Ettorre,

A.; Giulia Loi, A.; Scintu, F.; La Colla, P. *J. Med. Chem.* **2000**, *43*, 1886.  
(c) Lopez de Compadre, R. L.; Pearlstein, R. A.; Hopfinger, A. J.; Seyde, J. K. *J. Med. Chem.* **1987**, *30*, 900.

(2) For selected reviews, see: (a) Back, T. G.; Clary, K. N.; Gao, D. *Chem. Rev.* **2010**, *110*, 4498. (b) Garcia Ruano, J. L.; Aleman, J.; Parra, J.; Marzo, L. *Eur. J. Org. Chem.* **2014**, 1577.

(3) Selected examples for the anionic pathway, see: (a) Smorada, R. L.; Truce, W. E. *J. Org. Chem.* **1979**, *44*, 3444. (b) de Lucchi, O.; Licini, G.; Pasquato, L.; Senta, M. *J. Chem. Soc., Chem. Commun.* **1985**, 1597. (c) Garcia Ruano, J. L.; Aleman, J.; Marzo, L.; Alvarado, C.; Tortosa, M. *Angew. Chem., Int. Ed.* **2012**, *51*, 2712. (d) Garcia Ruano, J. L.; Aleman, J.; Marzo, L.; Alvarado, C.; Tortosa, M.; Diaz-Tendero, S.; Fraile, A. *Chem.—Eur. J.* **2012**, *18*, 8414. (e) Marzo, L.; Perez, L.; Yuste, F.; Aleman, J.; Garcia Ruano, J. L. *Chem. Commun.* **2015**, *51*, 346. Selected examples for the radical pathway, see: (f) Schaffner, A. P.; Darmency, V.; Renaud, P. *Angew. Chem., Int. Ed.* **2006**, *45*, 5847. (g) Hoshikawa, T.; Kamijo, S.; Inoue, M. *Org. Biomol. Chem.* **2013**, *11*, 164. (h) Todoroki, H.; Iwatsu, M.; Urabe, D.; Inoue, M. *J. Org. Chem.* **2014**, *79*, 8835.

(4) (a) Simpkins, N. S. *Sulphones in Organic Synthesis*; Pergamon Press: Oxford, U.K., 1993. (b) Back, T. G. *Tetrahedron* **2001**, *57*, 5263.

(5) Aziz, J.; Messaoudi, S.; Alami, M.; Hamze, A. *Org. Biomol. Chem.* **2014**, *12*, 9743.

(6) Zhdankin, V. V. *Hypervalent Iodine Chemistry: Preparation, Structure, and Synthetic Applications of Polyvalent Iodine Compounds*; Wiley: Chichester, U.K., 2014.

(7) (a) Manolikakes, G.; Umierski, N. *Org. Lett.* **2013**, *15*, 188. (b) Manolikakes, G.; Umierski, N. *Org. Lett.* **2013**, *15*, 4972.

(8) (a) Zhdankin, V. V.; Stang, P. J. *Tetrahedron* **1998**, *54*, 10927. (b) Brand, J. P.; Waser, J. *Chem. Soc. Rev.* **2012**, *41*, 4165.

(9) Tykwinski, R. R.; Williamson, B. L.; Fischer, D. R.; Stang, P. J.; Arif, A. M. *J. Org. Chem.* **1993**, *58*, 5235.

(10) Hamnett, D. J.; Moran, W. J. *Org. Biomol. Chem.* **2014**, *12*, 4156.

(11) (a) Woolven, H.; Gonzalez-Rodriguez, C.; Marco, L.; Thompson, A. L.; Willis, M. C. *Org. Lett.* **2011**, *13*, 4876. For a recent review on the use of DABSO, see: (b) Deeming, A. S.; Emmett, E. J.; Richards-Taylor, S. R.; Willis, M. C. *Synthesis* **2014**, 46, 2701.

(12) Deeming, A. S.; Russell, C. J.; Hennessy, A. J.; Willis, M. C. *Org. Lett.* **2014**, *16*, 150.

(13) Rocke, B. N.; Bahnck, K. B.; Herr, M.; Lavergne, S.; Mascitti, V.; Perreault, C.; Polivkova, J.; Shavnaya, A. *Org. Lett.* **2014**, *16*, 154.

(14) Emmett, E. J.; Hayter, B. R.; Willis, M. C. *Angew. Chem., Int. Ed.* **2014**, *53*, 10204.

(15) (a) Johnson, M. W.; Bagley, S. W.; Mankad, N. P.; Bergman, R. G.; Mascitti, V.; Toste, F. D. *Angew. Chem., Int. Ed.* **2014**, *53*, 4404. (b) Ye, S.; Wang, H.; Xiao, Q.; Ding, Q.; Wu, J. *Adv. Synth. Catal.* **2014**, *356*, 3225.

(16) Zhang, D.; An, Y.; Li, Z.; Wu, J. *Angew. Chem., Int. Ed.* **2014**, *53*, 2451. (b) Zhang, D.; Li, Y.; An, Y.; Wu, J. *Chem. Commun.* **2014**, 50, 8886.

(17) (a) Ochiai, M.; Masaki, Y.; Shiro, M. *J. Org. Chem.* **1991**, *56*, 5511. (b) Zhdankin, V. V.; Kuehl, C. J.; Krasutsky, A. P.; Bolz, J. T.; Simonsen, A. J. *J. Org. Chem.* **1996**, *61*, 6547.

(18) Selected examples: (a) Brand, J. P.; Charpentier, J.; Waser, J. *Angew. Chem., Int. Ed.* **2009**, *48*, 9346. (b) Brand, J. P.; Waser, J. *Angew. Chem., Int. Ed.* **2010**, *49*, 7304. (c) Fernandez Gonzalez, D.; Brand, J. P.; Waser, J. *Chem.—Eur. J.* **2010**, *16*, 9457. (d) Nicolai, S.; Erard, S.; Fernandez Gonzalez, D.; Waser, J. *Org. Lett.* **2010**, *12*, 384. (e) Nicolai, S.; Piemontesi, C.; Waser, J. *Angew. Chem., Int. Ed.* **2011**, *50*, 4680. (f) Brand, J. P.; Chevalley, C.; Scopelliti, R.; Waser, J. *Chem.—Eur. J.* **2012**, *18*, 5655. (g) Fernández González, D.; Brand, J. P.; Mondière, R.; Waser, J. *Adv. Synth. Catal.* **2013**, *355*, 1631. (h) Frei, R.; Waser, J. *J. Am. Chem. Soc.* **2013**, *135*, 9620. (i) Li, Y.; Brand, J. P.; Waser, J. *Angew. Chem., Int. Ed.* **2013**, *52*, 6743. (j) Chen, C. C.; Waser, J. *Chem. Commun.* **2014**, 50, 12923. (k) Frei, R.; Wodrich, M. D.; Hari, D. P.; Borin, P.-A.; Chauvier, C.; Waser, J. *J. Am. Chem. Soc.* **2014**, *136*, 16563. For a review, see: (l) Brand, J. P.; Fernandez Gonzalez, D.; Nicolai, S.; Waser, J. *Chem. Commun.* **2011**, 47, 102.

(19) (a) Ochiai, M.; Kunishima, M.; Nagao, Y.; Fuji, K.; Shiro, M.; Fujita, E. *J. Am. Chem. Soc.* **1986**, *108*, 8281. (b) Ochiai, M.; Ito, T.; Takaoka, Y.; Masaki, Y.; Kunishima, M.; Tani, S.; Nagao, Y. *J. Chem. Soc., Chem. Commun.* **1990**, 118.