



Cite this: *Org. Biomol. Chem.*, 2018, **16**, 7561

Received 24th September 2018,  
Accepted 1st October 2018

DOI: 10.1039/c8ob02375a

rsc.li/obc

## Transition metal-free coupling of terminal alkynes and hypervalent iodine-based alkyne-transfer reagents to access unsymmetrical 1,3-diynes†

J. Schörghumer and M. Waser \*

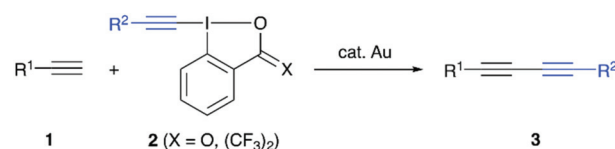
**A variety of unsymmetrical 1,3-diynes can easily be accessed in good yields under catalyst- and transition metal-free conditions by reacting terminal alkynes with hypervalent iodine-based electrophilic alkyne-transfer reagents.**

The 1,3-diyne motif is a frequently found structural element that either serves as a valuable synthon for further manipulations, or can be found as such in biologically active compounds as well as functional materials.<sup>1</sup> Thus, it comes as no surprise that the development of efficient synthesis strategies has attracted considerable interest.<sup>2</sup> The most classical strategy to access symmetric 1,3-diynes is the Cu-mediated oxidative homo-coupling of terminal alkynes, also known as Glaser coupling, which was first reported almost 150 years ago.<sup>3</sup> A lot of efforts have been made since that to introduce different and complementary strategies to carry out the transition metal-catalysed homo-coupling of alkynes.<sup>4</sup> In addition, the syntheses of unsymmetrically substituted 1,3-diynes became a very important topic recently.<sup>2,5–7</sup> The groups of Liu<sup>7a</sup> and Patil<sup>7b</sup> independently described very efficient protocols for the gold-catalysed coupling of terminal alkynes with hypervalent iodine-based alkyne transfer reagents in 2017. The use of these electrophilic alkyne transfer reagents for organic syntheses in general is well-established,<sup>8</sup> however their use to access 1,3-diynes has so far received less attention.<sup>7,9</sup> Besides the aforementioned very recent hetero-coupling relying on gold-catalysis,<sup>7</sup> the groups of Lee and Kitamura reported early examples for the coupling of alkynylcuprates with alkynylidonium salts in 1997 already.<sup>9</sup> We have recently reported the transition metal-free homocoupling of terminal alkynes in the presence of hypervalent iodine reagents.<sup>10</sup> Control experiments suggested the presence of *in situ* formed alkynylidonium species, which then undergo the C(sp)–C(sp) coupling upon

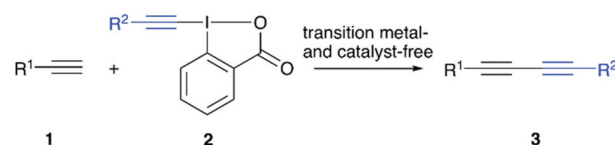
reaction with an *in situ* formed Li-acetylide. Based on the emerging interest in unsymmetrical 1,3-diynes we now investigated the development of an operationally simple and generally applicable strategy to access a variety of diynes **3** by reacting simple terminal alkynes **1** with preformed hypervalent iodine reagents **2** under transition metal- and catalyst-free conditions (Scheme 1).

We started our investigations by optimizing the reaction of *n*-hexyne (**1a**) with the phenylacetylene-based benziodoxolone **2a** (Table 1 gives an overview of the most relevant screening results). All reactions were carried out by deprotonating the terminal alkyne **1a** first, followed by subsequent addition of the iodine reagent **2a**. Dry THF was used as a solvent for all screening reactions (toluene or CH<sub>2</sub>Cl<sub>2</sub> were very low yielding only), and *n*-BuLi (1.6 M in hexanes) was the base of choice for the initial experiments. We immediately realized that both steps, deprotonation and electrophile addition, are best carried out at –78 °C (entries 1–3). Furthermore, *n*-BuLi should be used only in a very subtle excess, as addition of 1.5 equiv. of *n*-BuLi lead to lower yields accompanied by the

a) Gold-catalysed coupling by Liu et al.<sup>7a</sup> and Patil et al.<sup>7b</sup>



b) Transition metal-free uncatalysed 1,3-diyne synthesis (this work)

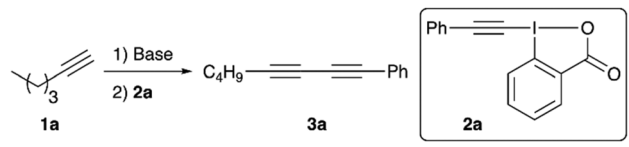


**Scheme 1** Previous gold-catalysed reports for the synthesis of dissymmetric 1,3-diynes **3** (a) and targeted heterocoupling of alkynes **1** and alkyne-transfer reagents **2** under transition metal-free non-catalysed conditions.

Institute of Organic Chemistry, Johannes Kepler University Linz,  
Altenbergerstrasse 69, 4040 Linz, Austria. E-mail: mario.waser@jku.at

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8ob02375a



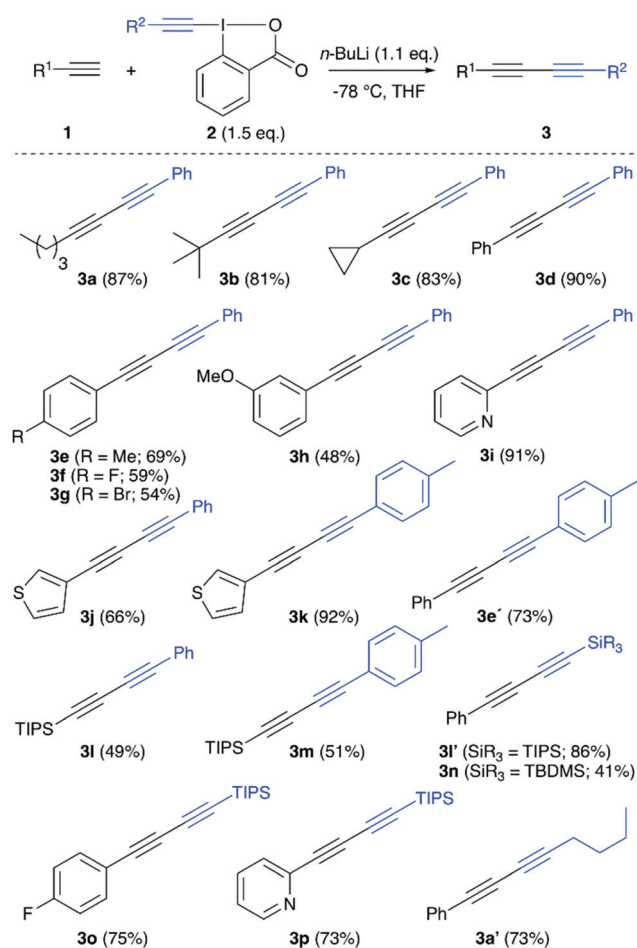
**Table 1** Identification of the best-suited conditions for the synthesis of 1,3-diyne **3a**


Entry <sup>a</sup>	Base (equiv.)	Deprotonation (T)	2a (equiv.)	Cond. (T, t)	Yield <sup>b</sup> (%)
1	<i>n</i> -BuLi (1.1)	-40 °C	1.1	-78 °C-r.t., 2 h	46
2	<i>n</i> -BuLi (1.1)	-40 °C	1.1	-78 °C-r.t., 5 h	47
3	<i>n</i> -BuLi (1.1)	-78 °C	1.1	-78 °C-r.t., 2 h	65
4	<i>n</i> -BuLi (1.5)	-78 °C	1.1	-78 °C-r.t., 3 h	53
5	<i>n</i> -BuLi (1.1)	-78 °C	1.5	-78 °C-r.t., 3 h	87
6	LDA (1.1)	-78 °C	1.5	-78 °C-r.t., 3 h	10
7	NaHMDS (1.1)	-78 °C	1.5	-78 °C-r.t., 3 h	21

<sup>a</sup> All reactions were carried out using 0.2 mmol **1a** in dry THF (1 mL) by first adding the base at the indicated temperature and stirring for 1–2 hours, followed by adding **2a** at -78 °C and slowly warming the reaction mixture to r.t. over the indicated period. <sup>b</sup> Isolated yields.

formation of notable quantities of by-products already (compare entries 3 and 4). It should be noted that we generally observed two side-reactions that may potentially occur, depending on the reaction conditions: (a) homocoupling of *n*-hexyne **1a** and (b) homocoupling of phenylacetylene (which may be rationalized by cleavage off phenylacetylene from **2a** followed by coupling with a second molecule of **2a**). However, these side-reactions could be suppressed rather efficiently by using only 1.1 equivalent of *n*-BuLi combined with a slight excess of alkyne-transfer reagent **2a**, which gave the dissymmetric diyne **3a** in a satisfying and easily reproducible isolated yield of 87% (entry 5). Other bases were tested as well (see entries 6 and 7 for two examples), but neither of them was found to be satisfying and we thus used the conditions shown in entry 5 (Table 1) to investigate the application scope for this reaction by using different terminal alkynes **1** with a small variety of different hypervalent iodine reagents **2** (Scheme 2).

Addition of different aliphatic terminal alkynes to the phenylacetylene-based electrophilic reagent **2a** proceeded rather high yielding (see products **3a–3c**), and also aryl-substituted alkynes allowed for the synthesis of the diaryl-containing 1,3-diyne **3d–3j** in reasonable yields, although some influence of the aryl substituents was observed (*i.e.* for products **3g** and **3h**, in those cases notable amounts of homocoupling side-products were observed). By using a tolylacetylene-based iodine reagent instead, a similar, maybe even slightly higher, reactivity as for the phenyl-based one could be observed (see products **3k** and **3e'**). The use of simple TIPS-acetylene as the nucleophilic reaction partner was possible as well (giving products **3l** and **3m**) albeit it was found that the inverse approach by adding an arylacetylene nucleophile to the TIPS-acetylene-containing iodine reagent allows for a clearly higher yield (compare the results for **3l** and **3l'**). Surprisingly, the TBDMS-acetylene-based iodine reagent gave the diyne **3n** in significantly lower yield, and we hereby observed quite a large amount of the homocoupling of TBDMS-acetylene. Finally, also the hexyne-based iodine reagent could be successfully employed as demonstrated in the synthesis of **3a'**.



**Scheme 2** Application scope (all reactions were carried out by adding 1.1 eq. *n*-BuLi to 0.2 mmol **1** in dry THF at -78 °C, stirring for 2 hours, followed by adding 1.5 eq. of **2** at -78 °C and slowly warming the reaction mixture to r.t. over 3 h; TIPS = triisopropylsilyl; TBDMS = *t*-butyldimethylsilyl).



Summing up the observations made during the investigations of the scope, it becomes obvious that the nature of the employed terminal alkyne plays an important role (which supports the crucial role of the intermediate Li-acetylide). Thus this reaction works best for more electron-neutral terminal alkynes, while the presence of more polar groups or silyl groups unfortunately reduces the reactivity and increasing amounts of homocoupling products are observed in those cases.

In conclusion, we have found that dissymmetric 1,3-diynes **3** can be synthesized without the need of any (transition metal) catalysts by reacting terminal alkynes **1** (which are *in situ* deprotonated with *n*-BuLi) with hypervalent iodine-based electrophilic alkyne-transfer reagents **2**. This procedure works in reasonable yields for different terminal alkynes **1** as well as iodine reagents **2** and thus may provide a complementary protocol to the recently developed powerful gold-catalysed approaches.<sup>7</sup>

**General reaction procedure:** 140  $\mu$ L (0.22 mmol, 1.1 eq.) of a solution of *n*-BuLi (1.6 M in hexane) were added to a solution of the corresponding terminal alkyne **1** (0.20 mmol, 1.0 eq.) in dry THF (1 mL) at  $-78$  °C. After stirring for 2 h, the corresponding ethynyl-benziodoxolone **2** (0.30 mmol, 1.5 eq.) was added in one portion. The mixture was allowed to reach room temperature over 3 h while stirring rapidly. The resulting suspension was quenched with 2 mL of a saturated solution of NaHCO<sub>3</sub> and extracted three times with 5 mL dichloromethane. After evaporation of the solvent, the crude product was purified by column chromatography (silica gel) to afford the targeted diyne **3** in the reported yield.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The NMR spectrometers used were acquired in collaboration with the University of South Bohemia (CZ) with financial support from the European Union through the EFRE INTERREG IV ETC-AT-CZ program (project M00146, "RERI-uasb").

## Notes and references

- (a) F. Bohlmann, T. Burkhardt and C. Zdero, *Naturally Occurring Acetylenes*, Academic Press, London, 1973;
- (b) L. Hansen and P. M. Boll, *Phytochemistry*, 1986, **25**, 285–293; (c) H. Matsunaga, M. Katano, H. Yamamoto, H. Fujito, M. Mori and K. Takata, *Chem. Pharm. Bull.*, 1990, **38**, 3480–3482; (d) J. M. Tour, *Chem. Rev.*, 1996, **96**, 537–553; (e) R. E. Martin and F. Diederich, *Angew. Chem., Int. Ed.*, 1999, **38**, 1350–1377; (f) A. L. K. S. Shun and R. R. Tykwinski, *Angew. Chem., Int. Ed.*, 2006, **45**, 1034–1057.
- W. Shi and A. Lei, *Tetrahedron Lett.*, 2014, **55**, 2763–2772.
- C. Glaser, *Ber. Dtsch. Chem. Ges.*, 1869, **2**, 422–424.
- For some recent transition metal-catalysed approaches: (a) J.-H. Li, Y. Liang and X.-D. Zhang, *Tetrahedron*, 2005, **61**, 1903–1907; (b) J.-H. Li, Y. Liang and Y.-X. Xie, *J. Org. Chem.*, 2005, **70**, 4393–4396; (c) J. Yan, J. Wu and H. Jin, *J. Organomet. Chem.*, 2007, **692**, 3636–3639; (d) T. Kurita, M. Abe, M. Tomohiro, Y. Monguchi and H. Sajiki, *Synlett*, 2007, 2521–2524; (e) K. Kamata, S. Yamaguchi, M. Kotani, K. Yamaguchi and N. Mizuno, *Angew. Chem., Int. Ed.*, 2008, **47**, 2407–2410; (f) S. Chen, W. Wu and F. Tsai, *Green Chem.*, 2009, **11**, 269–274; (g) S. Zhang, X. Liu and T. Wang, *Adv. Synth. Catal.*, 2011, **353**, 1463–1466; (h) A. Leyva-Perez, A. Domenech, S. I. Al-Resayes and A. Corma, *ACS Catal.*, 2012, **2**, 121–126; (i) M. G. Leeming, G. N. Khairallah, S. Osburn, K. Vikse and R. A. J. O'Hair, *Aust. J. Org. Chem.*, 2014, **67**, 701–710; (j) Z.-J. Wang, P.-H. Wang, J.-J. Lv, J.-J. Feng, X. Xu, A.-J. Wang, C.-T. Au and R. Qiu, *RSC Adv.*, 2015, **5**, 96372–96376; (k) X.-L. Shi, Q.-Q. Hu, F. Wang, W.-Q. Zhang and P.-G. Duan, *J. Catal.*, 2016, **337**, 233–239.
- A. L. K. Shi Shun, E. T. Chernick, S. Eisler and R. R. Tykwinski, *J. Org. Chem.*, 2003, **68**, 1339–1347.
- (a) H. Peng, Y. Xi, N. Ronaghi, B. Dong, N. G. Akhmedov and X. Shi, *J. Am. Chem. Soc.*, 2014, **136**, 13174–13177; (b) J. D. Myrtle, A. M. Beekman and R. A. Barrow, *Org. Biomol. Chem.*, 2016, **14**, 8253–8260; (c) S. Ghorai and D. Lee, *Tetrahedron*, 2017, **73**, 4062–4069; (d) E. Godin, A.-C. Bedard, M. Raymond and S. K. Collins, *J. Org. Chem.*, 2017, **82**, 7576–7582; (e) B. S. Chinta and B. Baire, *Org. Biomol. Chem.*, 2017, **15**, 5908–5911.
- (a) X. Li, X. Xie, N. Sun and Y. Liu, *Angew. Chem., Int. Ed.*, 2017, **56**, 6994–6998; (b) S. Banerjee and N. T. Patil, *Chem. Commun.*, 2017, **53**, 7937–7940.
- For a detailed review see: J. P. Brand and J. Waser, *Chem. Soc. Rev.*, 2012, **41**, 4165–4179.
- C.-H. Lee, N.-H. Jeong, S.-W. Lee and T. Kitamura, *J. Ind. Eng. Chem.*, 1997, **3**, 155–159.
- J. Schörgenhuber and M. Waser, *Tetrahedron Lett.*, 2016, **57**, 1678–1680.

