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Synthesis and applications of highly functionalized 1-halo-3-substituted bicyclo[1.1.1]pentanes†

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Bicyclo[1.1.1]pentanes (BCPs) are important bioisosteres of 1,4-disubstituted arenes, *tert*-butyl and acetylenic groups that can impart physicochemical benefits on drug candidates. Here we describe the synthesis of BCPs bearing carbon and halogen substituents under exceptionally mild reaction conditions, *via* triethylborane-initiated atom-transfer radical addition ring-opening of tricyclo[1.1.1.0^{1,3}]pentane (TCP) with alkyl halides. This chemistry displays broad substrate scope and functional group tolerance, enabling application to BCP analogues of biologically-relevant targets such as peptides, nucleosides, and pharmaceuticals. The BCP halide products can be converted to the parent phenyl/*tert*-butyl surrogates through triethylborane-promoted dehalogenation, or to other derivatives including carbonyls, alcohols, and heterocycles.

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Introduction

Bioisosteres are important motifs in drug design,¹ enabling the expansion of chemical and intellectual property space² while also improving biological profiles relative to the parent functionality. Bicyclo[1.1.1]pentanes (BCPs) have shown particularly impressive results in this field as surrogates for 1,4-disubstituted arenes, *tert*-butyl, and alkyne groups, imparting desirable properties such as membrane permeability, solubility and metabolic stability.^{3,4} Examples include BCP analogues of the γ -secretase inhibitor BMS-708, 163 (**1**, Fig. 1a)⁵ and resveratrol (**2**),⁶ in which the BCP serves as a bioisostere for a *p*-substituted arene due to its comparable positioning of substituents; and of the pulmonary arterial hypertension agent bosentan (**3**, Fig. 1b), where a mono-substituted BCP replaces a *tert*-butyl group.^{7,8} Despite these favourable attributes, access to carbon-substituted BCPs remains a challenge. Their synthesis typically relies on multistep sequences from a small number of available precursors,⁹ or requires relatively harsh conditions,¹⁰ necessitating early-stage introduction of the BCP and limiting functional group tolerance.

The BCP system is most commonly accessed through 'strain release'¹¹ reactions of tricyclo[1.1.1.0^{1,3}]pentane **4** (TCP),¹² where in addition to classical approaches,^{3,13} recent work has

seen the development of elegant methods for the synthesis of heteroatom-substituted BCPs.^{11,14} The insertion of TCP into C–X bonds would offer an appealing entry to carbon-substituted BCPs, as well as the opportunity for further functionalization of the halide product (**5**, Fig. 1c).¹⁵ To date however, this process has only been achieved using highly activated reagents (*e.g.* CF₃I),¹³ or by mercury lamp irradiation of an alkyl or aryl halide,^{10*d,e*} or methyl lithium-promoted alkyl halide addition.^{10*b*} Despite the importance of these contributions, all display limitations in substrate scope, functional group compatibility, or scalability, due to the constraints of the reaction conditions.

Atom transfer radical addition (ATRA) reactions using chemical initiators are an attractive alternative for TCP ring

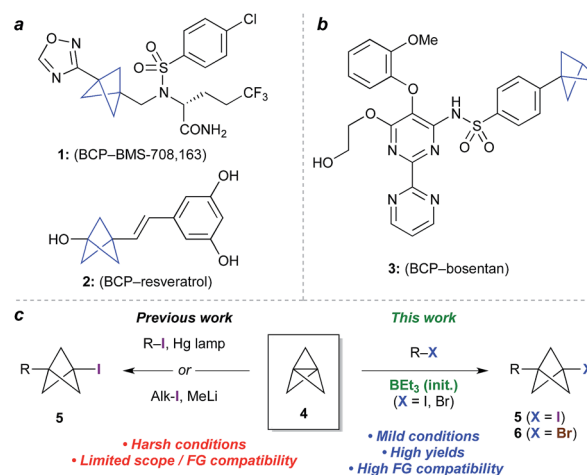


Fig. 1 (a) Bicyclo[1.1.1]pentanes (BCPs) as 1,4-disubstituted arene bioisosteres. (b) BCP as a *tert*-butyl bioisostere. (c) Approaches to 1-halo-3-substituted BCPs in previous and current work.

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opening, but previously required a large excess of the radical precursor, or suffered from the formation of oligomeric 'staffane' byproducts due to multiple insertions into TCP. ^{10d,13} Building on our studies of radical-mediated nucleoside alkylation, ¹⁶ we questioned whether triethylborane could serve as an effective initiator ¹⁷ for the synthesis of 1-halo-3-substituted bicyclopentanes (Fig. 1c). Here we report the development of this method as an efficient and highly functional group tolerant route to halogenated BCPs, the utility of which are illustrated through various derivatizations. Importantly, the mild conditions of this chemistry allow the synthesis of bicyclopentanes that could not be accessed using other methods, and open up opportunities for the late-stage functionalization of more complex molecules.

Results and discussion

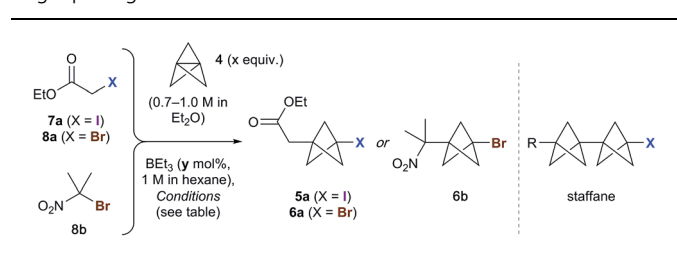
We began our studies using ethyl iodoacetate **7a**, and were delighted to observe complete reaction in just 15 min at room temperature using 1.3 equivalents of TCP and 10 mol% triethylborane (1 M in hexane), with the product iodide **5a** isolated in 83% yield (Table 1, entry 1). A mild exotherm was observed on addition of triethylborane to the reaction, which could be avoided by reducing the temperature (entry 2). The amount of initiator could be decreased to 1 mol% without detriment (entries 2–5), and the quantity of TCP could also be lowered (to

1.1 equiv.), affording **5a** in 92% yield (entry 6). Successful reaction was similarly observed using TCP as a solution in dibutyl ether (0.19 M), which is significant for industrial applications (entry 7). The presence of triethylborane proved crucial, as reactions run in its absence failed to reach completion in the dark or light (entries 8 and 9). ¹⁸ Importantly, no staffane byproducts were observed under the optimized conditions (entry 6).

Use of the equivalent bromoacetate **8a** led to incomplete conversion, or significant staffane formation (entry 10). ¹⁹ However, we were pleased to find that 2-bromo-2-nitropropane **8b** underwent rapid reaction with 1.3 equivalents of TCP to afford **6b** in 67% yield (entry 11). ²⁰ As with **7a**, full conversion was maintained on cooling the reaction to 0 °C (73%, entry 12); however, lowering the quantity of triethylborane resulted in incomplete reaction (entry 13). The success of substrate **8b** presumably reflects a more efficient bromine atom abstraction by the bicyclopentyl radical in the propagation step compared to ester **8a**, due to the greater stability of the tertiary nitro-substituted radical.

The scope of the triethylborane-initiated ATRA was next explored using organohalides featuring a variety of functional groups (Fig. 2). ²¹ While the optimized conditions (Table 1, entry

Table 1 Optimization of triethylborane-promoted tricyclopentane ring-opening



Entry	Substrate	x (equiv.)	y (mol%)	T	t	Yield ^a (%)
1	7a	1.3	10	rt	15 min	83
2	7a	1.3	10	0 °C	15 min	87
3	7a	1.3	5	0 °C	15 min	89
4	7a	1.3	1	0 °C	15 min	95
5	7a	1.3	0.5	0 °C	15 min	(67 : 33)
6	7a	1.1	1	0 °C	15 min	92
7	7a	2.0	10	rt	16 h	98 ^b
8	7a	2.0	0	rt	20 h	(60 : 40) ^c
9	7a	2.0	0	rt	20 h	(33 : 67)
10	8a	1.3	10	rt	20 h	47 ^d
11	8b	1.3	10	rt	15 min	67
12	8b	1.3	10	0 °C	15 min	73
13	8b	1.3	1	0 °C	15 min	(60 : 40)

^a Isolated yields. Figures in parentheses indicate incomplete reactions, and the ratio of starting material to product as judged by ¹H NMR spectroscopic analysis of the crude reaction mixture. ^b Reaction carried out in Bu₂O (0.19 M). ^c Reaction carried out in the dark. ^d Isolated as a 6 : 1 mixture of **6a** : staffane.

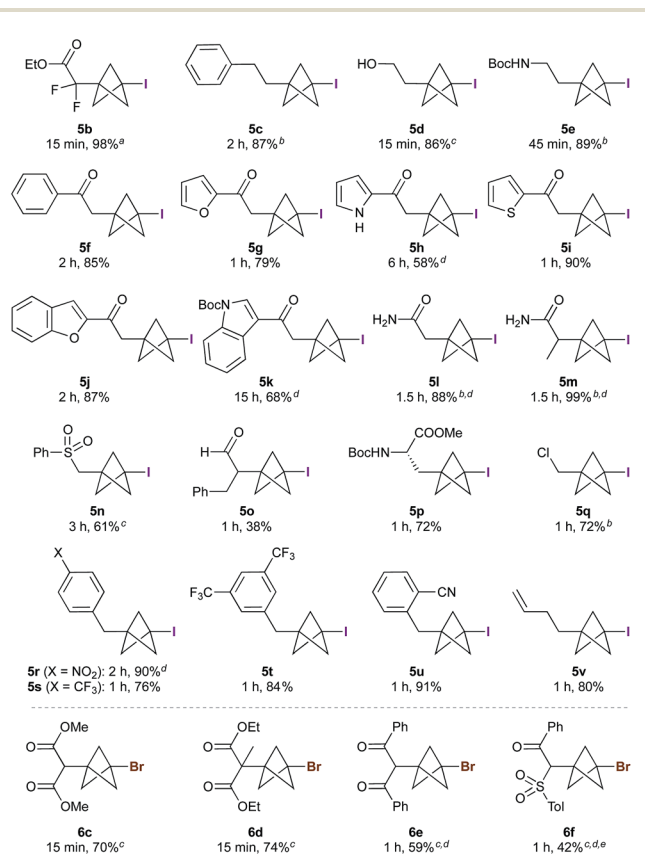


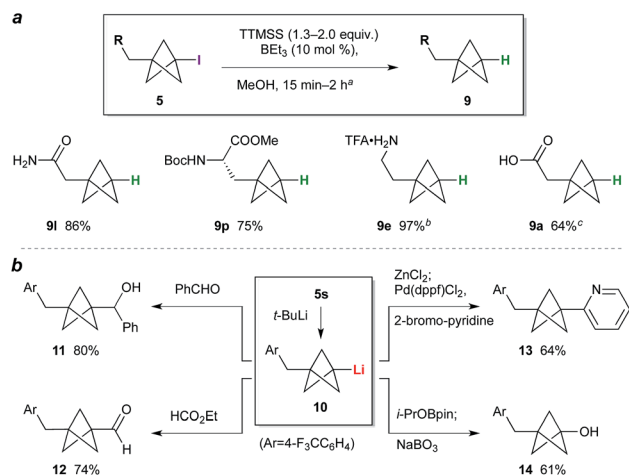
Fig. 2 Synthesis of 1-iodo- and 1-bromo-3-substituted BCPs. All reactions performed using 2 equiv. tricyclo[1.1.1.0^{1,3}]pentane (TCP) and 10 mol% BEt₃ (1 M in hexane) at room temperature, unless indicated otherwise. ^a1.1 equiv. TCP, 1 mol% BEt₃, 0 °C. ^b1.3 equiv. TCP, rt. ^c1.3 equiv. TCP, 0 °C. ^dCo-solvent added to solubilize the substrate: MeOH for **5m**, **n**; CH₂Cl₂ for **5h**, **k**, **r** and **6e**, **f**. ^e5% staffane observed.



6) translated smoothly to ethyl difluoroiodoacetate, giving bench-stable BCP **5b** in 98% yield,²² incomplete reactions were observed in other cases. To ensure consistency, the substrate screen was therefore conducted using 10 mol% BEt_3 and 1.3–2.0 equivalents of TCP at either 0 °C or room temperature, according to the requirements of the substrate. Alkyl iodides showed excellent reactivity, tolerating groups such as hydroxyls and carbamates (**5c–e**). α -Iodoketones gave high yields across a range of substrates, including heteroaromatics (**5f–k**). Other electron-withdrawing groups such as amides and sulfones also proved viable (**5l–5n**),²⁰ including the sensitive but potentially valuable aldehyde **5o**. The formation of **5p** is particularly notable due to the potential utility of BCP analogues of amino acids;²³ no racemization was observed in this reaction. Even chloriodomethane was found to be a suitable substrate, delivering chloromethyl BCP **5q** in good yield. Benzyl halides also underwent smooth reaction, providing electron-withdrawing substituents were present on the arene (which presumably accelerates the halide abstraction step from the intermediate BCP radical, **5r–5u**).²⁰ The involvement of radical intermediates was supported by the use of (iodomethyl)cyclopropane, which exclusively gave the ring-opened product **5v**. Finally, other electron-deficient bromides proved suitable partners, affording products **6c–f** in short reaction times; for these substrates, the presence of two electron-withdrawing groups proved essential for efficient bromine atom abstraction.

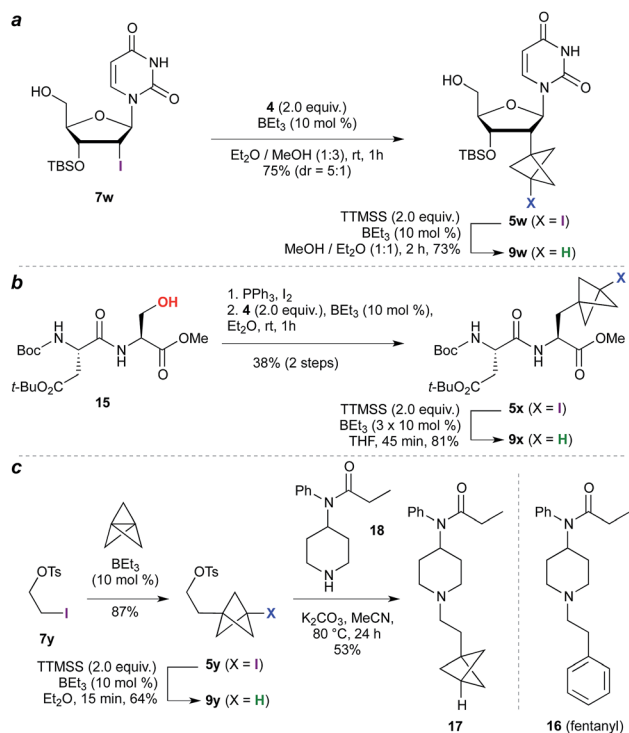
While the BCP iodide products are stable towards storage at –20 °C (particularly if crystalline), in some cases gradual coloration was observed at ambient temperature. Deiodination was expected to enhance BCP stability, and would simultaneously access the parent phenyl/*tert*-butyl bioisostere. We questioned whether triethylborane could again be used as initiator for this C–I bond reduction; pleasingly, the use of 1.3 equivalents of tris(trimethylsilyl)silane (TTMSS) as a non-toxic hydrogen atom source²⁴ and 10 mol% BEt_3 effected smooth deiodination of various BCP iodides (Scheme 1a), giving amide **9l**, a BCP analogue of phenylalanine (**9p**), and trifluoroacetate salt **9e**, the free base of the latter being a BCP equivalent of the monoaminergic neuromodulator phenethylamine. Given the use of the same initiator for both radical processes, a one-pot reaction sequence was also performed in which BCP **9a** was prepared directly from **7a** in a single operation (64%). Transformation of the iodide into substituents other than hydrogen was also explored (Scheme 1b): lithiation of **5s**,^{10b} followed by reaction of the resulting organolithium **10** with benzaldehyde or ethyl formate, gave alcohol **11** and aldehyde **12** respectively. Alternatively, transmetalation to the organozinc and cross-coupling with 2-bromopyridine afforded pyridyl BCP **13** (64%).^{10a,b} Pleasingly, the synthesis of a phenol bioisostere⁶ (**14**) could also be accomplished by a borylation quench, followed by oxidation with sodium perborate.

The high functional group tolerance and mild reaction conditions suggested that this methodology could be exploited in the functionalization of more complex organic molecules. For example, 2'-iodouridine derivative **7w** (Scheme 2a) underwent successful ATRA reaction to generate **5w** (75%, 5 : 1 dr), with deiodination delivering the 2'-deoxy-2'-BCP nucleoside **9w**



Scheme 1 (a) BCP deiodination. ^aSee the ESI† for experimental conditions for each substrate. ^bDeiodination product from **5e** treated with $\text{CF}_3\text{CO}_2\text{H}$ (yield over two steps). ^cIodide **7a** processed through TCP ring opening, deiodination, and hydrolysis (NaOH , MeOH) without purification. (b) BCP iodide functionalization.

(73%). More generally, the availability of iodides from hydroxyl groups provides a multitude of opportunities for bioisostere installation: Appel reaction of the aspartate–serine dipeptide **15** (Scheme 2b) afforded iodide **5x**, which on ATRA reaction and subsequent TTMSS reduction generated BCP-functionalized dipeptide **9x**. This sequence corresponds to a three step conversion of a serine residue to a potential phenylalanine



Scheme 2 Synthesis of nucleoside, dipeptide, and pharmaceutical BCP analogues.



- J. McCarthy, P. G. Parsons, G. H. Walter, M. T. Smith, H. M. Cooper, S. K. Nilsson, J. Tsanaktsidis, G. P. Savage and C. M. Williams, *Angew. Chem., Int. Ed.*, 2016, **55**, 3580–3585; (f) J. A. Bull, R. A. Croft, O. A. Davis, R. Doran and K. F. Morgan, *Chem. Rev.*, 2016, **116**, 12150–12233.
- 5 A. F. Stepan, C. Subramanyam, I. V. Efremov, J. K. Dutra, T. J. O'Sullivan, K. J. DiRico, W. S. McDonald, A. Won, P. H. Dorff, C. E. Nolan, S. L. Becker, L. R. Pustilnik, D. R. Riddell, G. W. Kauffman, B. L. Kormos, L. Zhang, Y. Lu, S. H. Capetta, M. E. Green, K. Karki, E. Sibley, K. P. Atchison, A. J. Hallgren, C. E. Oborski, A. E. Robshaw, B. Sneed and C. J. O'Donnell, *J. Med. Chem.*, 2012, **55**, 3414–3424.
- 6 Y. L. Goh, Y. T. Cui, V. Pendharkar and V. A. Adsool, *ACS Med. Chem. Lett.*, 2017, **8**, 516–520.
- 7 M. V. Westphal, B. T. Wolfstädter, J.-M. Plancher, J. Gatfield and E. M. Carreira, *ChemMedChem*, 2015, **10**, 461–469.
- 8 (a) For other recent examples of BCPs in medicinal chemistry, see: N. D. Measom, K. D. Down, D. J. Hirst, C. Jamieson, E. S. Manas, V. K. Patel and D. O. Somers, *ACS Med. Chem. Lett.*, 2017, **8**, 43–48; (b) K. C. Nicolaou, J. Yin, D. Mandal, R. D. Erande, P. Klahn, M. Jin, M. Aujay, J. Sandoval, J. Gavrilyuk and D. Vourloumis, *J. Am. Chem. Soc.*, 2016, **138**, 1698–1708; (c) R. Pellicciari, R. Filosa, M. C. Fulco, M. Marinozzi, A. Macchiarulo, C. Novak, B. Natalini, M. B. Hermit, S. Nielsen, T. N. Sager, T. B. Stensbøl and C. Thomsen, *ChemMedChem*, 2006, **1**, 358–365.
- 9 (a) P. Kaszynski and J. Michl, *J. Org. Chem.*, 1988, **53**, 4593–4594; (b) 1-azido-3-iodo BCP: K. B. Wiberg and N. McMurdie, *J. Am. Chem. Soc.*, 1994, **116**, 11990–11998.
- 10 (a) I. S. Makarov, C. E. Brocklehurst, K. Karaghiosoff, G. Koch and P. Knochel, *Angew. Chem., Int. Ed.*, 2017, **56**, 12774–12777; (b) M. Messner, S. I. Kozhushkov and A. de Meijere, *Eur. J. Org. Chem.*, 2000, 1137; (c) E. W. Della and D. K. Taylor, *J. Org. Chem.*, 1994, **59**, 2986–2996; (d) P. Kaszynski, A. C. Friedli and J. Michl, *J. Am. Chem. Soc.*, 1992, **114**, 601–620; (e) P. Kaszynski, N. D. McMurdie and J. Michl, *J. Org. Chem.*, 1991, **56**, 307–316; to our knowledge only esters and a ketone have been shown to be tolerated under mercury lamp irradiation, and only tetrahydropyranyl ethers under MeLi promoted TCP ring opening.
- 11 (a) R. Gianatassio, J. M. Lopchuk, J. Wang, C.-M. Pan, L. R. Malins, L. Prieto, T. A. Brandt, M. R. Collins, G. M. Gallego, N. W. Sach, J. E. Spangler, H. Zhu, J. Zhu and P. S. Baran, *Science*, 2016, **351**, 241; (b) J. M. Lopchuk, K. Fjelbye, Y. Kawamata, L. R. Malins, C.-M. Pan, R. Gianatassio, J. Wang, L. Prieto, J. Bradow, T. A. Brandt, M. R. Collins, J. Elleraas, J. Ewanicki, W. Farrell, O. O. Fadeyi, G. M. Gallego, J. J. Mousseau, R. Oliver, N. W. Sach, J. K. Smith, J. E. Spangler, H. Zhu, J. Zhu and P. S. Baran, *J. Am. Chem. Soc.*, 2017, **139**, 3209–3226.
- 12 K. R. Mondanaro and W. P. Dailey, *Org. Synth.*, 1998, **75**, 98. For a more recent preparation using PhLi see ref. 11a. Appropriate precautions should be taken in the preparation of **4** due to the need for relatively large volumes of organolithium solutions.
- 13 K. B. Wiberg and S. T. Waddell, *J. Am. Chem. Soc.*, 1990, **112**, 2194–2216.
- 14 (a) R. M. Bär, S. Kirschner, M. Nieger and S. Bräse, *Chem.–Eur. J.*, 2018, **24**, 1373–1382; (b) J. Kanazawa, K. Maeda and M. Uchiyama, *J. Am. Chem. Soc.*, 2017, **139**, 17791–17794; (c) K. Bunker, C. Guo, M. Grier, C. Hopkins, J. Pinchman, D. Slee, P. Q. Huang and M. Kahraman, WO Pat., 2016/044331, 2016; (d) Y. L. Goh and V. A. Adsool, *Org. Biomol. Chem.*, 2015, **13**, 11597–11601; (e) N. T. Thirumoorthi, C. J. Shen and V. A. Adsool, *Chem. Commun.*, 2015, **51**, 3139–3142; (f) Y. L. Goh, E. K. W. Tam, P. H. Bernardo, C. B. Cheong, C. W. Johannes, A. D. William and V. A. Adsool, *Org. Lett.*, 2014, **16**, 1884–1887; (g) K. D. Bunker, N. W. Sach, Q. Huang and P. F. Richardson, *Org. Lett.*, 2011, **13**, 4746–4748.
- 15 (a) J. L. Adcock and A. A. Gakh, *J. Org. Chem.*, 1992, **57**, 6206–6210; see also ref. 9a, 10b and 14c; (b) C. Mazal, A. J. Paraskos and J. Michl, *J. Org. Chem.*, 1998, **63**, 2116–2119.
- 16 (a) M. M. Haugland, A. H. El-Sagheer, R. J. Porter, J. Peña, T. Brown, E. A. Anderson and J. E. Lovett, *J. Am. Chem. Soc.*, 2016, **138**, 9069–9072; (b) M. Sakeda, S. Ichikawa, A. Matsuda and S. Shuto, *J. Org. Chem.*, 2003, **68**, 3465–3475; (c) M. Sakeda, S. Ichikawa, A. Matsuda and S. Shuto, *Angew. Chem., Int. Ed.*, 2002, **41**, 4748–4750.
- 17 (a) D. P. Curran and T. R. McFadden, *J. Am. Chem. Soc.*, 2016, **138**, 7741–7752; (b) C. Ollivier and P. Renaud, *Chem. Rev.*, 2001, **101**, 3415–3434.
- 18 While the reaction of **7a** was not affected if conducted in degassed solvent under a nitrogen atmosphere (<15 min), the equivalent formation of **5s** (Table 1) was significantly retarded, requiring 20 h reaction time to reach completion. We believe this supports a standard oxygen-mediated initiation mechanism. See ref. 17, and the ESI.†
- 19 For Et₃B-promoted ATRA using **8a**, see: H. Yorimitsu, H. Shinokubo, S. Matsubara, K. Oshima, K. Omoto and H. Fujimoto, *J. Org. Chem.*, 2001, **66**, 7776–7785.
- 20 Low temperature single crystal X-ray diffraction data were 40 collected for **5m**, **5r**, **5s**, **6b** and **17** using a (Rigaku) Oxford Diffraction SuperNova A diffractometer. Raw frame data were collected and reduced using CrysAlisPro. The structures were solved using SuperFlip [L. Palatinus and G. Chapuis, *J. Appl. Crystallogr.*, 2007, **40**, 786–790] and refined using CRYSTALS [P. W. Betteridge, J. R. Carruthers, R. I. Cooper, K. Prout and D. J. Watkin, *J. Appl. Crystallogr.*, 2003, **36**, 1487; R. I. Cooper, A. L. Thompson and D. J. Watkin, *J. Appl. Crystallogr.*, 2010, **43**, 1100–1107]. See the ESI (CIF).† CCDC 1825056–1825060 contain the crystallographic data for this paper.†
- 21 See the ESI† for a discussion of substrates and functional groups that were not tolerated, or led to incomplete conversion, or significant staffane formation.
- 22 This reaction was also conducted on a 5.00 mmol scale, which afforded **5b** in 94% yield (1.50 g).
- 23 (a) For examples of BCP amino acids, see ref. 4c, 8c, and: S. O. Kokhan, A. V. Tymtsunik, S. L. Grage, S. Afonin,



- O. Babii, M. Berditsch, A. V. Strizhak, D. Bandak, M. O. Platonov, I. V. Komarov, A. S. Ulrich and P. K. Mykhailiuk, *Angew. Chem., Int. Ed.*, 2016, **55**, 14788–14792. (b) S. Pritz, M. Pätz, G. Szeimies, M. Dathe and M. Bienert, *Org. Biomol. Chem.*, 2007, **5**, 1789–1794; (c) For additional examples, see: R. Pellicciari, M. Raimondo, M. Marinozzi, B. Natalini, G. Costantino and C. Thomsen, *J. Med. Chem.*, 1996, **39**, 2874–2876.
- 24 (a) TTMSS has been employed in the reduction of BCP halides using AIBN as an initiator, see ref. 14c and e. Tributyltin hydride has also been used as the reducing agent, see ref. 10a; (b) N. T. Thirumoorthi and V. A. Adsool, *Org. Biomol. Chem.*, 2016, **14**, 9485–9489.
- 25 (a) F. Weigend and R. Ahlrichs, *Phys. Chem. Chem. Phys.*, 2005, **7**, 3297–3305; (b) Y. Zhao and D. G. Truhlar, *Theor. Chem. Acc.*, 2008, **120**, 215–241.
- 26 For appropriateness of the method chosen, see ref. 14b, and the ESI.†

