# Chemical Science



### **EDGE ARTICLE**

View Article Online
View Journal | View Issue



Cite this: Chem. Sci., 2019, 10, 3765

dall publication charges for this article have been paid for by the Royal Society of Chemistry

Received 14th December 2018 Accepted 6th February 2019

DOI: 10.1039/c8sc05581e

rsc.li/chemical-science

# Chiral phosphoric acid-catalyzed enantioselective construction of structurally diverse benzothiazolopyrimidines†

Lucie Jarrige,<sup>a</sup> Danijel Glavač,<sup>ab</sup> Guillaume Levitre,<sup>a</sup> Pascal Retailleau,<sup>a</sup> Guillaume Bernadat,<sup>c</sup> Luc Neuville <sup>©</sup> and Géraldine Masson <sup>©</sup>\*

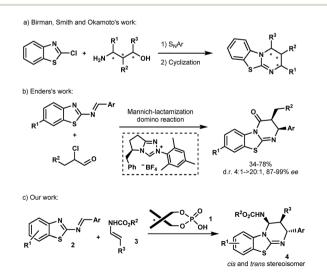
A highly efficient catalytic enantioselective [4+2] cycloaddition was developed between 2-benzothiazolimines and enecarbamates. A wide range of benzothiazolopyrimidines bearing three contiguous stereogenic centers was obtained in high to excellent yields and with excellent diastereo-and enantioselectivities (d.r. > 98:2 and up to >99% ee). Furthermore, this chiral phosphoric acid-catalyzed strategy was scalable and enabled access to a new class of optically pure Lewis base isothiourea derivatives.

#### Introduction

Isothioureas constitute an important class of organocatalysts due to their highly nucleophilic nature.1 Recent efforts have been directed to the development of chiral derivatives capable of promoting enantioselective transformations, which are difficult to control using traditional nucleophilic catalysts. Among the several promising examples, benzothiazolopyrimidine, better known as homobenzotetramisole (HBTM), has assumed a leading position as a chiral catalyst. Chiral HBTMs have been successfully used in a number of asymmetric transformations<sup>1a</sup> including the kinetic resolution of secondary alcohols and carboxylic acids,2 desymmetrization,3 acyl transfer,4 annulation processes,5 and lactonisation,6 among others.7 Nevertheless, despite the interest in optically active HBTMs, <sup>1a</sup> there have been only limited reports on their enantioselective synthesis.8 In 2008, Birman et al. disclosed the first asymmetric synthesis of HBTMs following a seven-step sequence involving a key  $S_NAr$  reaction between 2-chlorobenzo[d]thiazole and amino-alcohols prepared from the chiral pool of amino acids (Scheme 1a).2f Chiral amino-alcohols, produced by catalytic enantioselective procedures, were later engaged in similar routes as reported by the same group as well as by Smith et al. and Okamoto et al. (Scheme 1a). 4a,b,9 More recently, Enders et al.

established an efficient asymmetric synthesis of precursors of HBTMs by the *N*-heterocyclic carbene-catalyzed Mannich/lactamization domino reaction of 2-benzothiazolimines and α-chloroaldehydes (Scheme 1b).<sup>10</sup> Despite these examples, no catalytic asymmetric approach has been reported so far to directly construct these heterocycles. An attractive direct access relying on the aza-Diels–Alder approach has been reported by Mellor *et al.* for the synthesis of homobenzotetramisole in an achiral stoichiometric context.<sup>9,11</sup> The requirement of a full equivalent of the acid to perform the reaction is self-evident due to the presence of the basic isothiourea that might capture the acid during the formation of HBTMs.<sup>2f,12</sup> Turning this process into a catalytic enantioselective approach would be highly

 $<sup>\</sup>dagger$  Electronic supplementary information (ESI) available. CCDC 1815484 and 1851627. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c8sc05581e



Scheme 1 Previous studies and our strategy.

<sup>&</sup>quot;Institut de Chimie des Substances Naturelles, CNRS UPR 2301, Université Paris-Sud, Université Paris-Saclay, 1, Avenue de la Terrasse, 91198 Gif-sur-Yvette Cedex, France. E-mail: geraldine.masson@cnrs.fr

<sup>&</sup>lt;sup>b</sup>Ruđer Bošković Institute, Division of Organic Chemistry and Biochemistry, Bijenička cesta 54, 10000 Zagreb, Croatia

LERMIT, 5, rue J. B. Clément, 92296 Châtenay Malabry, France

**Chemical Science Edge Article** 

appealing, not only to access new potent chiral Lewis bases but also for the generation of novel biologically active molecules.<sup>13</sup>

In the past few years, chiral phosphoric acids have been widely recognized as powerful catalysts for the synthesis of various heterocycles containing nitrogen atoms. 14 In this context, we have recently documented phosphoric acid-catalyzed enantioselective aza-Diels-Alder reactions14,15 of various azadienes with enecarbamates as dienophiles. 16 Based on these studies, we envisioned that this Brønsted acid 1 may be capable of promoting the [4+2] cycloaddition between 2-benzothiazolimines 2 and enecarbamates 3 (Scheme 1c) allowing the development of an enantioselective synthesis of benzothiazolopyrimidines 4.

Because of the formation of the basic isothiourea during the process, catalyst poisoning might lead to difficulties in developing a catalytic version. Nevertheless, we reasoned that tuning the catalyst or using additives to induce a hydrogen shuttle could help in overcoming the problem.17

#### Results and discussion

With this proposal in mind, we examined the aza-Diels-Alder reaction of 2-benzothiazolimine<sup>18</sup> 2a with (E)-N-benzyl propenylcarbamate (3a) in the presence of chiral phosphoric acids 1. Gratifyingly, in dichloromethane at room temperature, the reaction promoted by 1a afforded benzothiazolopyrimidine 4a as a single diastereomer, with an enantiomeric excess of 95% albeit with a low yield (Table 1, entry 1). Obviously, while yields remained low, we were delighted to observe that no full catalyst inhibition was occurring. Encouraged by this result, we screened various reaction conditions to search for the optimal reaction conditions. Performing the reaction at 50 °C in DCE

Table 1 Optimization of the enantioselective [4+2] cycloaddition<sup>a</sup>

MeO

Ar = 
$$\rho$$
-BrC<sub>6</sub>H<sub>4</sub>

NHCbz

3a

Cat (10 mol%)

Ar

MeO

Ar

Ar

Cat.

CbzHN

NN

NA

Ar

Cat.

Entry	Ar	<i>T</i> (°C)	Solvent	$Yield^{b}$ (%)	ee <sup>c</sup> (%)
1 <sup>d</sup> 2 2 4	Ph (1a) 1a 9-Phenantryl (1b) 2,4,6-TRIP (1c)	RT 50 50 50	DCM DCE DCE DCE	40 41 29 62	95 80 85 98
5 <sup>e</sup>	1c	50	DCE	90 (68) <sup>f</sup>	99

<sup>&</sup>lt;sup>a</sup> Reaction conditions: 2a (0.10 mmol), 3a (0.20 mmol) and 1 (0.01 mmol) in 1.0 mL of solvent for 16 h. b Yields refer to the chromatographically pure diastereomer 4a determined to be higher than 98:2 by <sup>1</sup>H NMR. <sup>c</sup> ee values were determined by HPLC with a chiral stationary phase. d For 24 h. e With 3 equiv. of 3a. f With 5 mol% of 1c.

slightly improved the kinetics (16 h instead of 24 h) even if overall yields were not superior and significant erosion of enantioselectivity was observed.19 Then, a series of phosphoric acids were screened in DCE at 50 °C (entries 2-4), and we found that the catalyst structure had a marked impact on the reaction efficiency. For instance, the use of chiral phosphoric acid 1c having two bulky 3,3'-(2,4,6-triisopropylphenyl) groups improved the yield, presumably by facilitating the catalyst turnover. A further survey of reaction conditions revealed that the optimal conditions consisted in performing the reaction with 3 equivalents of 3a in the presence of 10 mol% of 1c, delivering 4a in 90% yield with almost perfect diastereo- and enantioselectivity (entry 5).

With optimized reaction conditions in hand, the scope of this chiral phosphoric acid-catalyzed enantioselective aza-Diels-Alder reaction was investigated and the results are summarized in Scheme 2. First, 2-benzothiazolimines bearing various substituents on the aromatic rings were evaluated. Pleasingly, the reaction was insensitive to the electronic nature of the substituents on the benzothiazole rings. For instance, the benzothiazole bearing either electron-donating or electronwithdrawing groups, such as methyl, halogen and trifluoromethyl groups, provided the corresponding benzothiazolopyrimidine 4b-f in high yields and excellent enantioselectivities (up to 99% ee). An unsubstituted benzothiazole ring proceeded uneventfully to give cycloadduct 4g with almost perfect enantioselectivity (up to 99% ee). It is worth noting that the reactions performed on a 1 mmol scale displayed the same results as the ones on a 0.10 mmol scale, further indicating the practicality of this protocol. On the other hand, the electronic properties of aromatic rings that came from aldehydes had no significant impact on the enantiomeric excess, though the reaction with the electron-rich imine gave a slightly lower yield (4h vs. 4a). When the reaction was conducted with imine, which had bromine at the *meta*-position, the desired cycloadduct 4k was obtained in 80% yield and 98% ee. Notably, 4k can be enriched to >99% ee by single recrystallization, which is very interesting in view of their use as organocatalysts. Moreover, a relative and absolute configuration was unambiguously determined by single crystal X-ray analysis (see the ESI†). Finally, the reaction with imine 21 derived from 2thiophenecarboxaldehyde proceeded smoothly to give the cycloadduct 4l in almost quantitative yields with 98% ee. This novel catalytic enantioselective cycloaddition was also applied to representative enecarbamates (Scheme 2). Notably, a variety of protecting groups bonded to the nitrogen atom of 3 (including Boc, Alloc and 4-(trifluoromethyl)benzylcarbamate) were well tolerated and the corresponding benzothiazolopyrimidines (4m-q) were, in all cases, isolated in excellent yields and excellent stereoselectivities. It is noteworthy that the catalyst loading was successfully reduced to 5.0 mol% (4q, entry 10). A comparable enantioselectivity (97%) was still achieved with a slightly lower chemical yield (70%).

In addition, β-substituted enecarbamates bearing different alkyl groups reacted efficiently to deliver expected trisubstituted-cycloadducts 4r-u in good to excellent yields with high selectivities (94 to 98% ee). The reaction with 3c or 3d

**Edge Article** 

Scheme 2 Scope of the enantioselective synthesis of benzothiazolopyrimidines 4.  $^{\rm a}$  General reaction conditions: 2 (0.1 mmol), 3 (0.3 mmol), 1c (10 mol%), 1,2-DCE (1 mL, 0.1 M), 50  $^{\rm c}$ C, and 24 h. Yields refer to chromatographically purified compounds. The d.r. values were determined by  $^{\rm 1}$ H NMR analysis to be higher than 98 : 2 in all cases. The ee values were determined by HPLC on a chiral stationary phase: see the ESI† for details.  $^{\rm a}$  From 1.0 mmol of 2g.  $^{\rm b}$  After recrystallization.  $^{\rm c}$  From 0.3 mmol of 2b.  $^{\rm d}$  With 5 mol% of 1c.

containing a cyclopropyl or silyl ether group formed products 4s or 4t with a slight decrease in the yield without affecting asymmetric induction.

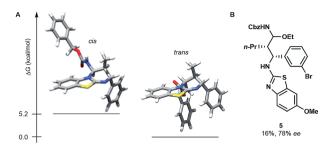
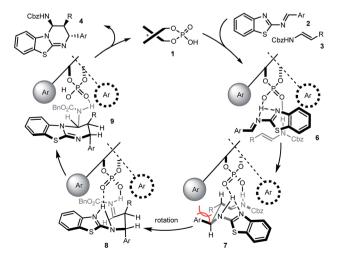


Fig. 1 (A) Most stable *cis* and *trans* conformers of 4g found at the B3LYP/6-31G(d) level. (B) Structure of intermediate 5 obtained in the presence of 3.0 equiv. of ethanol on the way to 4k.

The stereoselective issue of the reaction ((E)-enecarbamates to the cis product) raises questions about the mechanism. In order to get insight, we computationally evaluated the free energy of both diastereoisomers of the product 4g (Fig. 1A). The trans form was found favoured by more than 5.2 kcal  $mol^{-1}$  over its cis sibling (see the ESI†). This result together with the fact that the cis isomer is actually observed suggests that the cycloaddition occurs under kinetic control. A stepwise process was established by performing a control experiment in the presence of 3 equivalents of ethanol, which evidenced the formation of an N-Cbz-iminium intermediate (5, Fig. 1B).

Based on this, as well as our previous work and other published reports, 14,16,20 a possible catalytic mechanism was proposed in Scheme 3. We hypothesized that 1 would form a hydrogen network with the nitrogen Lewis basic sites of 2-benzothiazolimine 2 and the acidic site of enecarbamate 3, leading to complex 6. Mediated by the phosphoric acid double activation, subsequent bond formation would deliver complex 7. The aromatic and R residue would be positioned in a *trans* situation as found in the product, but the pseudo equatorial arrangement would create substantial strain. Maintained in the chiral cavity by the double hydrogen anchors, rotation of the backbone would furnish a sterically less-congested conformer 8. Finally the intramolecular nucleophilic addition of benzothiazole to iminium will give the observed *cis-trans* substituted



Scheme 3 Plausible mechanism and transition states

**Chemical Science Edge Article** 

Scheme 4 Derivatization of the benzothiazolopyrimidine product 4q.

benzothiazolopyrimidine 4.21 Heating, rapid formation of hydrogen-bonded complex 6 or use of excess 3 may all facilitate the decomplexation of 9 and ensure the catalyst turnover.

As a further illustration of the utility of this novel catalytic cycloaddition, we developed conditions to convert the carbamate group into other functional groups. The N-Boc group of 4q was removed by mild acidic hydrolysis to produce free alcohol 10 (a key intermediate to install additional molecular diversity) with no erosion of enantiopurity (Scheme 4). The absolute configuration was confirmed without ambiguity by X-ray diffraction analysis (see the ESI†).

#### Conclusions

We have developed an efficient enantioselective [4+2] cycloaddition organocatalyzed by chiral phosphoric-acid for the synthesis of benzothiazolopyrimidines bearing three contiguous stereogenic centers, which were unreported before. Involving 2-benzothioazolimines and enecarbamates substrates, this scalable strategy furnished a wide range of compounds with very high yields and enantioselectivity in a diastereospecific manner. The obtained cycloadducts were successfully converted into corresponding free alcohol heterocycles without any loss of enantioselectivity. Further investigations are in progress concerning the catalytic activity of these new compounds as chiral nucleophilic organocatalysts.

#### Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

We thank ICSN and CNRS for financial support; L. J. acknowledges Saclay University for the doctoral fellowship. D. J. acknowledges the French Embassy and an RBI fellowship for financial support. The IT department of Paris-Sud University is acknowledged for providing computing resources.

#### Notes and references

- 1 (a) J. Merad, J.-M. Pons, O. Chuzel and C. Bressy, Eur. J. Org. Chem., 2016, 5589; (b) S. Vellalath and D. Romo, Angew. Chem., Int. Ed., 2016, 55, 13934; (c) J. E. Taylor, S. D. Bull and J. M. J. Williams, Chem. Soc. Rev., 2012, 41, 2109; (d) N. De Rycke, F. Couty and O. R. P. David, Chem.-Eur. J., 2011, 17, 12852.
- 2 For selected examples, see: (a) M. D. Greenhalgh, S. M. Smith, D. M. Walden, J. E. Taylor, Z. Brice,

E. R. T. Robinson, C. Fallan, D. B. Cordes, A. M. Z. Slawin, H. C. Richardson, M. A. Grove, P. H.-Y. Cheong and A. D. Smith, Angew. Chem., Int. Ed., 2018, 57, 3200; Angew. Chem., 2018, 130, 3254; (b) K. Nakata, K. Gotoh, K. Ono, K. Futami and I. Shiina, Org. Lett., 2013, 15, 1170; (c) I. Shiina, K. Nakata, K. Ono, L. Y.-S. Onda and M. Itagaki, J. Am. Chem. Soc., 2010, 132, 11629; (d) I. Shiina, K. Nakata, K. Ono, M. Sugimoto and A. Sekiguchi, Chem.-Eur. J., 2010, 16, 167; (e) X. Yang and V. B. Birman, Adv. Synth. Catal., 2009, 351, 2301; (f) V. B. Birman and X. Li, Org. Lett., 2008, 10, 1115.

- 3 For selected recent examples, see: (a) J. Merad, P. Borkar, F. Caijo, J.-M. Pons, J.-L. Parrain, O. Chuzel and C. Bressy, Angew. Chem., Int. Ed., 2017, 56, 16052; Angew. Chem., 2017, 129, 16268; (b) J. Merad, P. Borkar, T. B. Yenda, C. Roux, J.-M. Pons, J.-L. Parrain, O. Chuzel and C. Bressy, Org. Lett., 2015, 17, 2118; (c) V. B. Birman, H. Jiang and X. Li, Org. Lett., 2007, 9, 3237.
- 4 For selected examples, see: (a) C. Joannesse, C. P. Johnson, C. Concellon, C. Simal, D. Philp and A. D. Smith, Angew. Chem., Int. Ed., 2009, 48, 8914; Angew. Chem., 2009, 121, 9076; (b) Y. Zhang and V. B. Birman, Adv. Synth. Catal., 2009, 351, 2525; (c) V. B. Birman, E. W. Uffman, H. Jiang, X. Li and C. J. Kilbane, J. Am. Chem. Soc., 2004, 126, 12226.
- 5 For representative examples, see: (a) S. Wang, C. Rodríguez-Escrich and M. A. Pericas, Angew. Chem., Int. Ed., 2017, 56, 15068; Angew. Chem., 2017, 129, 15264; (b) S. L. Wang, J. Izquierdo, C. Rodriguez-Escrich and M. A. Pericàs, ACS Catal., 2017, 7, 2780; (c) E. R. T. Robinson, D. M. Walden, C. Fallan, M. D. Greenhalgh, P. H.-Y. Cheong and Sci., 2016, 7, 6919; Smith, Chem. E. R. T. Robinson, C. Fallan, C. Simal, A. M. Z. Slawin and A. D. Smith, Chem. Sci., 2013, 4, 2193.
- 6 For selected examples, see: (a) W. Kong and D. Romo, J. Org. Chem., 2017, 82, 13161; (b) L. C. Morrill, J. Douglas, T. Lebl, A. M. Z. Slawin, D. J. Fox and A. D. Smith, Chem. Sci., 2013, 4, 4146; (c) C. Simal, T. Lebl, A. M. Z. Slawin and A. D. Smith, Angew. Chem., Int. Ed., 2012, 51, 3653; Angew. Chem., 2012, **124**, 3713; (*d*) D. Belmessieri, L. C. Morrill, C. Simal, A. M. Z. Slawin and A. D. Smith, J. Am. Chem. Soc., 2011, 133, 2714; (e) C. A. Leverett, V. C. Purohit and D. Romo, Angew. Chem., Int. Ed., 2010, 49, 9479; Angew. Chem., 2010, 122, 9669; (f) V. C. Purohit, A. S. Matla and D. Romo, J. Am. Chem. Soc., 2008, 130, 10478.
- 7 For further examples, see: (a) M. E. Abbasov, B. M. Hudson, D. J. Tantillo and D. Romo, Chem. Sci., 2017, 8, 1511; (b) J. N. Arokianathar, A. B. Frost, A. M. Z. Slawin, D. Stead and A. D. Smith, ACS Catal., 2018, 8, 1153; (c) M. E. Abbasov, B. M. Hudson, D. J. Tantillo and D. Romo, J. Am. Chem. Soc., 2014, 136, 4492; (d) P.-P. Yeh, D. S. B. Daniels, D. B. Cordes, A. M. Z. Slawin and A. D. Smith, Org. Lett., 2014, 16, 964; (e) L. C. Morrill, L. A. Ledingham, J.-P. Couturier, J. Bickel, A. D. Harper, C. Fallan and A. D. Smith, Org. Biomol. Chem., 2014, 12, 624; (f) D. G. Stark, L. C. Morrill, P.-P. Yeh, A. M. Z. Slawin, T. J. C. O'Riordan and A. D. Smith, Angew. Chem., Int. Ed., 2013, 52, 11642; Angew. Chem., 2013, 125, 11856.

**Edge Article** 

8 Examples of racemic synthesis: (a) T. Raj, H. Sharma, Mayank, A. Singh, T. Aree, N. Kaur, N. Singh and D. O. Jang, ACS Sustainable Chem. Eng., 2017, 5, 1468; (b) S. Okamoto, Y. Sakai, S. Watanabe, S. Nishi, A. Yoneyama, H. Katsumata, Y. Kosaki, R. Sato, M. Shiratori, M. Shibuno and T. Shishido, Tetrahedron Lett., 2014, 55, 1909; (c) B. Ranieri, O. Robles and D. Romo, J. Org. Chem., 2013, 79, 6291.

- 9 B. Viswambharan, T. Okimura, S. Suzuki and S. Okamoto, *J. Org. Chem.*, 2011, 76, 6678.
- 10 Q. Ni, X. Song, J. Xiong, G. Raabe and D. Enders, *Chem. Commun.*, 2015, 51, 1263.
- 11 J. M. Mellor and H. Rataj, Tetrahedron Lett., 1996, 37, 2619.
- 12 (a) B. Maji, C. Joannesse, T. A. Nigst, A. D. Smith and H. Mayr, *J. Org. Chem.*, 2011, **76**, 5104; (b) M. Kobayashi and S. Okamato, *Tetrahedron Lett.*, 2006, **47**, 4347.
- 13 (a) S. Gupta, N. Ajmera, N. Gautam, R. Sharma and D. Gautam, Indian J. Chem., 2009, 48B, 853; (b) G. Shukla, A. K. Tiwari, V. K. Singh, A. Bajpai, H. Chandra and A. K. Mishra, Chem. Biol. Drug Des., 2008, 72, 533; (c) R. A. Glennon, S. M. Tejani-Butt, W. Padgett and J. W. Daly, J. Med. Chem., 1984, 27, 1364; (d) J. Yevich, D. Temple, R. Covington, D. Owens, R. Seidehamel and K. Dungan, J. Med. Chem., 1982, 25, 864.
- 14 Recent reviews on chiral phosphoric acid catalysis: (a)
  J. Merad, C. Lalli, G. Bernadat, J. Maury and G. Masson, Chem.-Eur. J., 2018, 24, 3925; (b) D. Parmar, E. Sugiono,
  S. Raja and M. Rueping, Chem. Rev., 2017, 117, 10608; Chem. Rev., 2014, 114, 9047; (c) S. Schenker, A. Zamfir,
  M. Freund and S. B. Tsogoeva, Eur. J. Org. Chem., 2011, 2209; (d) M. Rueping, A. Kuenkel and I. Atodiresei, Chem. Soc. Rev., 2011, 40, 4539; (e) D. Kampen, C. M. Reisinger and B. List, Top. Curr. Chem., 2010, 291, 395; (f) M. Terada, Bull. Chem. Soc. Jpn., 2010, 83, 101; (g) M. Terada, Synthesis, 2010, 1929; (h) M. Terada, Chem. Commun., 2008, 4097; (i) T. Akiyama, Chem. Rev., 2007, 107, 5744; (f) T. Akiyama, J. Itoh and K. Fuchibe, Adv. Synth. Catal., 2006, 348, 999.
- 15 (a) G. Masson, C. Lalli, M. Benohoud and G. Dagousset, *Chem. Soc. Rev.*, 2013, **42**, 902; (b) X. Jiang and R. Wang, *Chem. Rev.*, 2013, **113**, 5515; (c) D. Bello, R. Ramón and

- R. Lavilla, *Curr. Org. Chem.*, 2010, **14**, 332; (*d*) V. V. Kouznetsov, *Tetrahedron*, 2009, **65**, 2721; (*e*) P. Buonora, J.-C. Olsen and T. Oh, *Tetrahedron*, 2001, 57, 6099.
- 16 (a) H. Liu, G. Dagousset, G. Masson, P. Retailleau and J. Zhu, J. Am. Chem. Soc., 2009, 131, 4598; (b) G. Dagousset, J. Zhu and G. Masson, J. Am. Chem. Soc., 2011, 133, 14804; (c) G. Dagousset, P. Retailleau, G. Masson and J. Zhu, Chem.-Eur. J., 2012, 18, 5869; (d) L. He, M. Bekkaye, P. Retailleau and G. Masson, Org. Lett., 2012, 14, 3158; (e) L. He, G. Laurent, P. Retailleau, B. Folléas, J.-L. Brayer and G. Masson, Angew. Chem., Int. Ed., 2013, 52, 11088; Angew. Chem., 2013, 125, 11294; (f) J. Brioche, T. Courant, L. Alcaraz, M. Stocks, M. Furber, J. Zhu and G. Masson, Adv. Synth. Catal., 2014, 356, 1719; (g) J. Pous, T. Courant, G. Bernadat, B. I. Iorga, F. Blanchard and G. Masson, J. Am. Chem. Soc., 2015, 137, 11950; (h) A. Dumoulin and G. Masson, J. Org. Chem., 2016, 81, 10154; (i) L. Jarrige, F. Blanchard and G. Masson, Angew. Chem., Int. Ed., 2017, 56, 10573; Angew. Chem., 2017, 129, 10709.
- 17 For selected examples, see: (a) M. Rueping and C. Azap, Angew. Chem., Int. Ed., 2006, 45, 7832; Angew. Chem., 2006, 118, 7996; (b) G. Li and J. C. Antilla, Org. Lett., 2009, 11, 1075; (c) S. Müller, M. J. Webber and B. List, J. Am. Chem. Soc., 2011, 133, 1853; (d) L. Hong, W. Sun, D. Yang, G. Li and R. Wang, Chem. Rev., 2016, 116, 4006.
- 18 Selected examples using 2-benzothiazolimines in catalytic enantioselective reaction: (a) M. Frías, A. C. Carrasco, A. Fraile and J. Alemán, Chem.-Eur. J., 2018, 24, 3117; (b) H.-X. He, W. Yang and D.-M. Du, Adv. Synth. Catal., 2013, 355, 1137; (c) L. Li, B. Song, P. S. Bhadury, Y. Zhang, D. Hu and S. Yang, Eur. J. Org. Chem., 2011, 4743.
- 19 Addition of AcOH to improve catalyst turnover (see ref. 17) leads to substantial degradation.
- 20 (a) J. P. Reid and J. M. Goodman, Chem.-Eur. J., 2017, 23, 14248; (b) J. P. Reid, L. Simón and J. M. Goodman, Acc. Chem. Res., 2016, 49, 1029; (c) L. Simón and J. M. Goodman, J. Org. Chem., 2011, 76, 1775.
- 21 G. Fraenkel, J. Foos, C. C. Ho, Y. Liang, S. Q. A. Rizvi, H. Stucki and F. Steel, *Tetrahedron Lett.*, 1978, **19**, 327.