



Cite this: DOI: 10.1039/d5sc08502k

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

Received 3rd November 2025  
Accepted 12th December 2025

DOI: 10.1039/d5sc08502k

rsc.li/chemical-science

# Se-catalyzed enantioselective lactamization enabled by a *N*-fluorosulfonyl group: total synthesis of diaporisoindole A

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Electrophilic cyclization of unsaturated carboxylic acids is a well-established strategy for the catalytic enantioselective synthesis of lactones. In contrast, analogous catalytic transformations of unsaturated carboxamides to access enantioenriched lactams remain underdeveloped due to the ambident nucleophilicity of carboxamides, favoring nucleophilic attack from their oxygen atom. Herein, we unveil *N*-(fluorosulfonyl)carboxamides as versatile and reactive substrates that overcome this limitation, as demonstrated by a selenium-catalyzed enantioselective lactamization, facile product derivatization, and application to the total synthesis of diaporisoindole A. To further showcase their synthetic potential, we briefly demonstrate their applicability in racemic iodo- and hydro-lactamizations.

## Introduction

Forging C–O and C–N bonds within the same synthetic platform not only simplifies synthesis planning but also enables facile access to both O- and N-analogs of target molecules. Such synthetic strategies have been widely and often implicitly practiced, from classical S<sub>N</sub>2 alkylation and acylation to modern metal-catalyzed cross-coupling reactions.<sup>1</sup> The intramolecular addition of a carboxylic acid to an electrophilically activated alkene is a common approach for the construction of a lactone ring *via* C–O bond formation,<sup>2,3</sup> as represented by halo-lactonization. To date, dozens of catalytic enantioselective electrophilic lactonizations have been developed for these transformations, driven by the advent of new chiral catalysts.<sup>4–16</sup>

While the direct application of electrophilic lactonization procedures to lactamization *via* C–N bond formation would be highly desirable, the intrinsically favored *O*-cyclization of carboxamides necessitates their modification prior to use (Fig. 1a).<sup>17–19</sup> In stoichiometric electrophilic lactamization, carboxamides are classically modified as *N,O*-bis-trimethylsilyl, *N*-Boc or *N*-tosyl carboxamides to harness the ambident reactivity for forging a C–N bond (Fig. 1b).<sup>20–24</sup> This prerequisite makes catalytic enantioselective electrophilic lactamization challenging, as it demands a fine balance between nucleophilic

activation of the carboxamide and electrophilic activation of the internal alkene. To date, only two examples have been reported (Fig. 1c): Yeung and co-workers achieved an enantioselective bromolactamization using an amino carbamate catalyst,<sup>25</sup> while the Denmark group reported a chiral Lewis base-catalyzed sulfenolactamization.<sup>26,27</sup> In both cases, *N*-tosyl carboxamides were predominantly used, despite the removal of the *N*-tosyl group from the lactams being challenging, if not impossible. This dearth of catalytic electrophilic lactamization led to the development of several, mechanistically distinct, transition metal-catalyzed lactamizations of unsaturated carboxamides.<sup>28–31</sup>

## Results and discussion

In our pursuit of chiral selenium  $\pi$ -acid catalysis,<sup>32–36</sup> we faced the difficulty of adapting our selenium-catalyzed enantioselective electrophilic lactonization,<sup>37</sup> that proceeds through an oxidative catalytic cycle,<sup>38</sup> to lactamization (Fig. 1d).<sup>39–41</sup> Our initial attempts using *N*-acyl and *N*-Boc carboxamides failed to yield the desired lactams. Even with the *N*-tosyl group, which has been validated not only in asymmetric lactamizations (Fig. 1c) but also selenium-catalyzed C–N bond formations,<sup>42–45</sup> no desired product was obtained. In one case using *N*-methoxy carboxamides, the corresponding iminolactones were obtained exclusively.<sup>46</sup> The breakthrough came from our separate project where *N*-(fluorosulfonyl)carbamates were identified as highly effective nitrogen nucleophiles.<sup>47–50</sup> Incorporating the *N*-fluorosulfonyl group into unsaturated carboxamides led to clean conversion, producing the desired lactams with high enantioselectivities (Fig. 1d, this work). Herein, we detail the outcomes of this study, including its application to the enantioselective total synthesis of diaporisoindole A,<sup>51</sup> and two other representative electrophilic lactamizations.

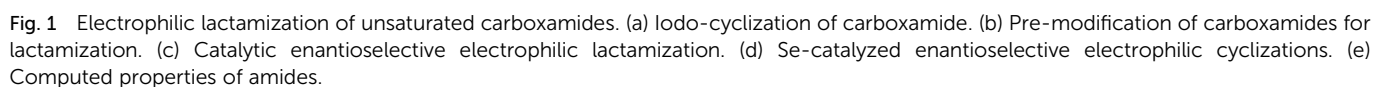
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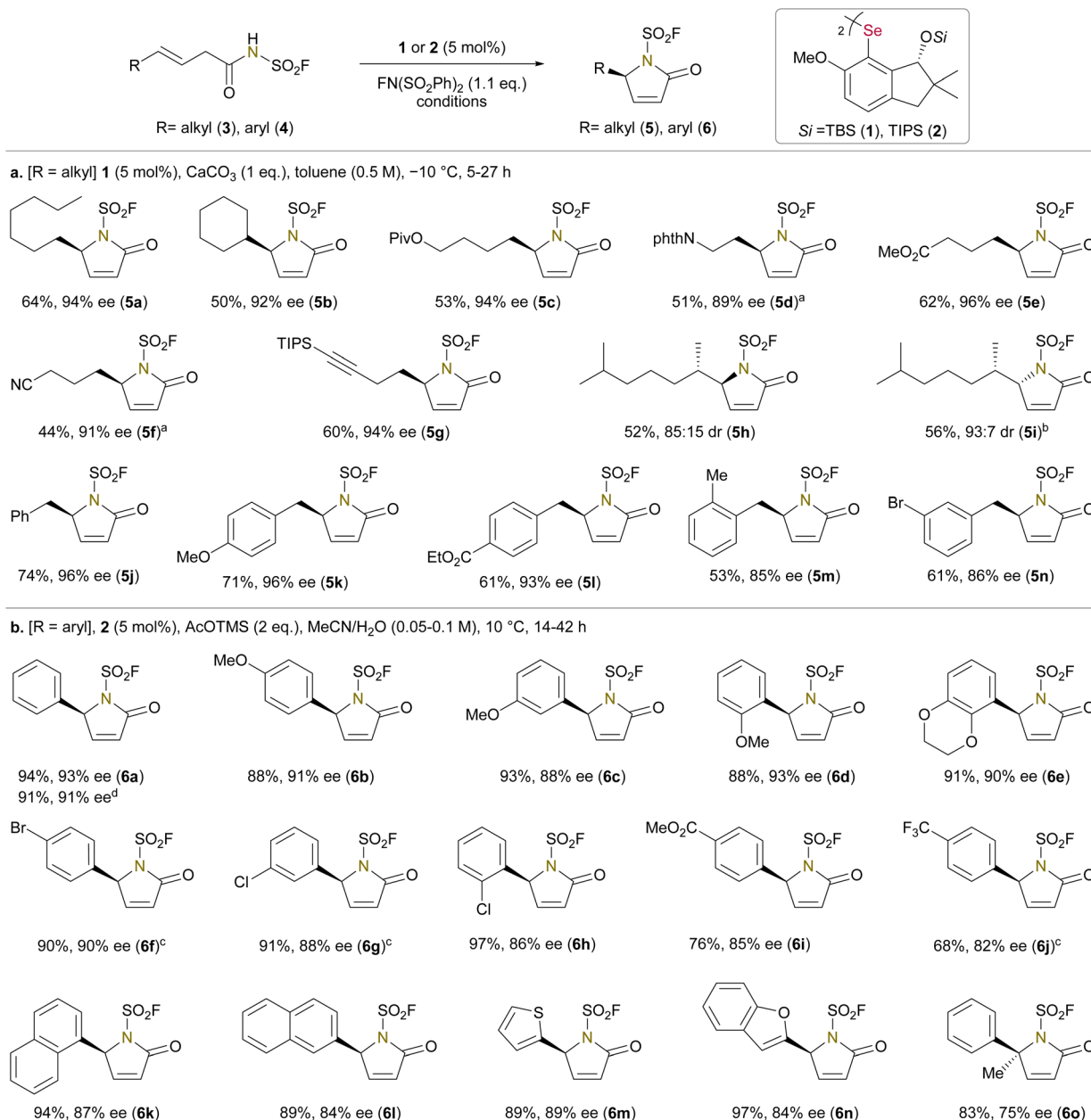
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In the further substrate evaluation, we encountered lower enantioselectivities with  $\gamma$ -aryl-substituted carboxamides **4**. It was concluded that the poorer enantioselectivity was caused by two factors: first, the formation of a desilylated catalyst which is still catalytically active but favors the opposite enantiomer (see



**Fig. 2** Selenium-catalyzed enantioselective lactamization. Yields are isolated yields. Reactions are performed on 0.1 mmol scale. (a) Aliphatic substrate scope. (b) Aromatic substrate scope. <sup>a</sup>Acetonitrile as solvent. <sup>b</sup>Performed with 5 mol% ent-1. <sup>c</sup>CF<sub>3</sub>CO<sub>2</sub>TMS (2.0 eq.) as additive. <sup>d</sup>Performed on 1.0 mmol scale with 3 mol% 2.

the SI), and second, the gradual racemization of the more acidic product. The longer reaction times required for the full conversion of aromatic substrates exacerbated these issues. In light of these challenges, we re-optimized the reaction conditions (see the SI), making them applicable to various aromatic substrates (Fig. 2b). Key modifications were the use of more desilylation-resistant, TIPS-substituted catalyst 2, TMSOAc as the non-basic fluoride scavenger and CH<sub>3</sub>CN/H<sub>2</sub>O as the solvent. Under these optimized reaction conditions, unsaturated lactams **6a–6n** were obtained in high yields and enantioselectivities, regardless of their substituent patterns or electronic properties. In case of aromatic substrates, *O*-cyclized products were not detected in the

crude NMR, contributing to higher product yields. Notably, a tri-substituted alkene also participated in this lactamization, forming a tetra-substituted stereocenter with 78% ee (**6o**). Finally, the reaction was scaled up to 1 mmol, yielding **6a** in 91% yield and 91% ee under 3 mol% of diselenide 2.

With the sufficient supply of enantioenriched unsaturated lactam **6a**, we next explored its derivatization. A 1,4-addition of an organocuprate reagent proceeded smoothly, giving 4,5-trans-disubstituted  $\gamma$ -lactam **7** in 95% yield (Fig. 3a). Although the C5 stereocenter of  $\gamma$ -lactam **6a** is prone to base-mediated racemization, its enantioselectivity was retained throughout the transformation. A Giese radical alkylation was also applicable to



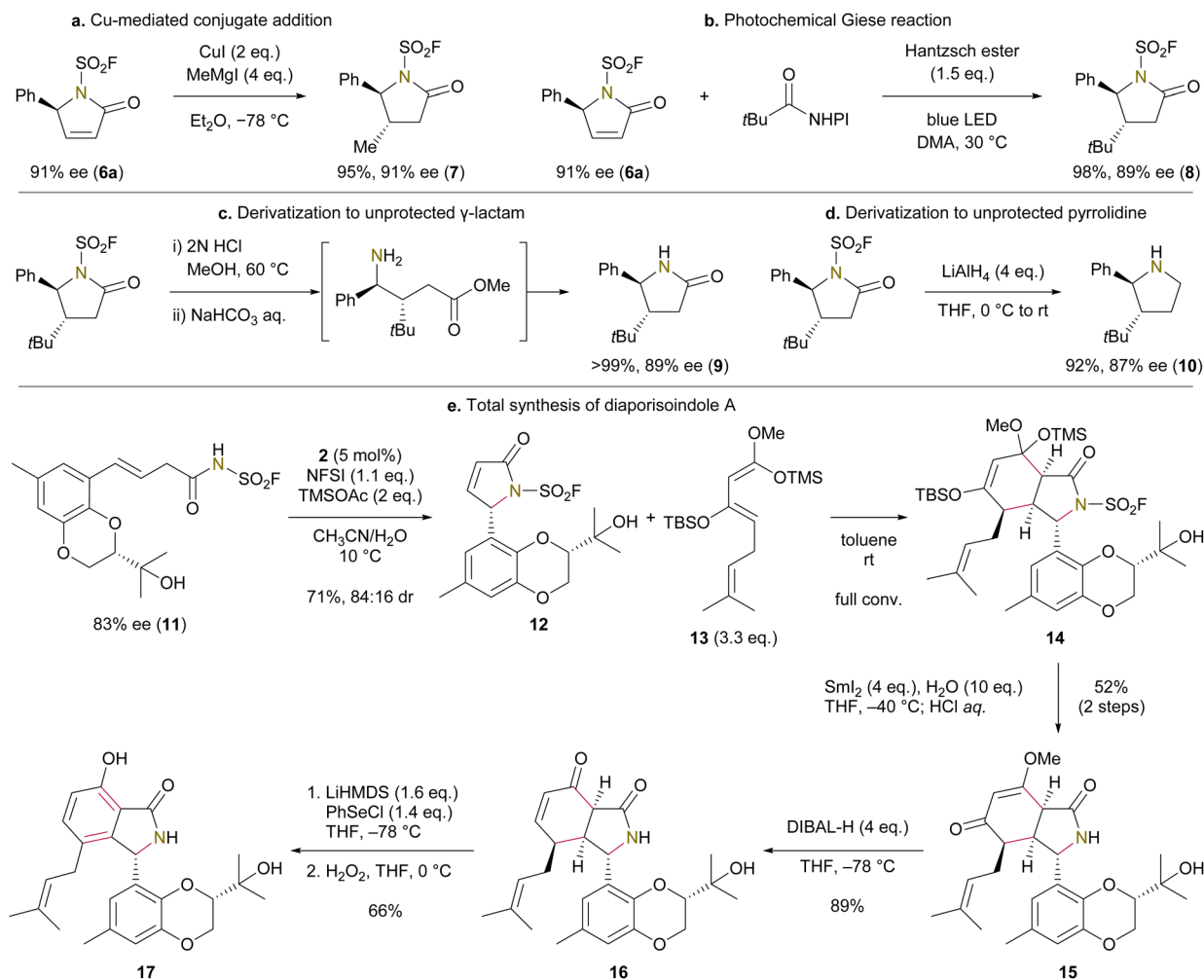


Fig. 3 Synthetic applications of selenium-catalyzed enantioselective lactamization. (a) Cu-mediated conjugate addition. (b) Photochemical Giese reaction. (c) Derivatization of unprotected  $\gamma$ -lactam. (d) Derivatization to unprotected pyrrolidine. (e) Total synthesis of diaporisoindole A.

lactam **6a**,<sup>63</sup> enabling the installation of a bulky *t*-butyl group on the  $\beta$ -position (Fig. 3b). The *N*-fluorosulfonyl group of **8** was readily removed by sequential treatment with acid and base (Fig. 3c). Moreover,  $\text{LiAlH}_4$ -mediated reduction of **8** furnished the corresponding *N*-H free pyrrolidine without opening the ring structure (Fig. 3d). These deprotection protocols further highlight the advantage of using the *N*-fluorosulfonyl group over the *N*-tosyl group.<sup>27</sup>

We next sought to demonstrate the feasibility of our selenium-catalyzed lactamization, and product derivatization and deprotection protocols in a more complex context—the enantioselective total synthesis of diaporisoindole A (Fig. 3e).<sup>51</sup> This natural product was chosen due to our particular interest in exploring a Diels–Alder reaction, capitalizing on the strongly electron-deficient nature of our unsaturated lactam products.<sup>64</sup> Literature reports indicate that successful Diels–Alder reactions of  $\alpha,\beta$ -unsaturated  $\gamma$ -lactams typically require a low-lying LUMO on the lactam, presenting a significant challenge for this transformation.<sup>65–69</sup> The lactamization precursor **11** was synthesized from cresol in enantioenriched form (see the SI), and was then subjected to our selenium catalysis, affording the key

unsaturated lactam **12** in 71% yield with 84:16 dr. We next investigated the Diels–Alder reaction, identifying electron-rich 1,3-bis-silyloxybutadiene **13** which reacted efficiently with **12** to furnish bicyclic compound **14** with the concurrent installation of the prenyl group. The relative stereochemical relationship of this Diels–Alder reaction was tentatively assigned as shown in the scheme, based on SCXRD of a model compound (see the SI). Our attempt to deprotect the *N*-fluorosulfonyl group as described in Fig. 3c did not result in the formation of the *N*-H free lactam in satisfactory yield. To address this issue, we employed a single-electron reduction of the fluorosulfonyl group by  $\text{SmI}_2$  which cleanly provided the *N*-H free lactam **15** without compromising the molecule's structural integrity.<sup>70</sup> Redox manipulation of this bicyclic intermediate led to diaporisoindole A **17**, which matched the reported NMR data and optical rotation. Notably, the synthesized quantity of diaporisoindole A allowed us to characterize a broad  $^{13}\text{C}$  peak at C8 that was unassigned in the original isolation research (see the SI).

Finally, to gain preliminary insight into the utility of *N*-fluorosulfonyl carboxamides in other electrophilic lactamizations, we examined racemic iodo- and hydro-lactamization using



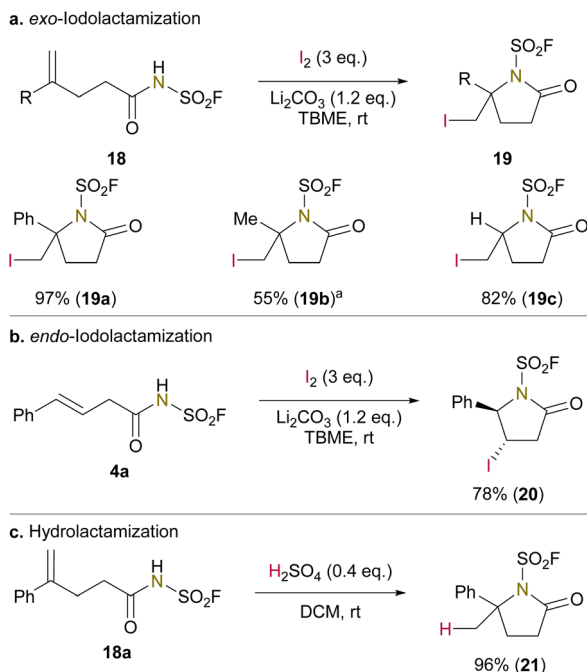


Fig. 4 Iodo- and hydro-lactamizations of unsaturated *N*-(fluorosulfonyl)carboxamides. (a) *exo*-lodolactamization. (b) *endo*-lodolactamization. (c) Hydrolactamization. <sup>a</sup>1,4-Dioxane as solvent.

selected *N*-fluorosulfonyl carboxamides (Fig. 4).<sup>71</sup> *Exo*- and *endo*-iodolactamizations using molecular iodine provided the corresponding lactams in moderate to good yields, along with minor amounts of iminolactones (Fig. 4a and b). Hydrolactamization catalyzed by  $H_2SO_4$  proceeded cleanly to give **21** in nearly quantitative yield (Fig. 4c). These results would provide a basis for developing catalytic enantioselective variants.<sup>72,73</sup> It should be noted that we observed the conversion of the initially formed iminolactone into the lactam in the hydrolactamization, but not in the seleno- or iodo-lactamizations, indicating the reversibility of this acid-catalyzed process and its thermodynamic preference for lactam formation.

## Conclusions

The synthetic community has recognized the difficulty of achieving electrophilic lactamization since the 1950s.<sup>74</sup> While pre-functionalization of amides has addressed this issue in stoichiometric contexts, the challenge continues to persist in asymmetric catalysis, widening the gap in progress between enantioselective electrophilic lactonization and lactamization. In this work, we demonstrated that *N*-(fluorosulfonyl)amides serve as effective substrates for chiral selenium-catalyzed electrophilic lactamization, a reactivity unattainable with other amides. The practicality of this method is underscored by the straightforward derivatization of the lactam products, including the enantioselective total synthesis of diaporisoindole A. This strategy also proved applicable to racemic iodo- and hydro-lactamizations, paving the way for the future development of enantioselective variants.

## Author contributions

D. Y. and S. Y. designed and conducted the experiments and prepared the SI. T. H. supervised the project and wrote the manuscript with input from D. Y. and S. Y.

## Conflicts of interest

There are no conflicts to declare.

## Data availability

CCDC 2482628–2482632 and 2484902 contain the supplementary crystallographic data for this paper.<sup>75a–f</sup>

The data that supports the findings of this study are available in the supplementary information (SI) of this article. Supplementary information is available. See DOI: <https://doi.org/10.1039/d5sc08502k>.

## Acknowledgements

This work was supported by JSPS KAKENHI Grant Numbers 19H02710 and 22H02072 (T. H.), and RIKEN Student Researcher (RSR) and Junior Research Associate (JRA) programs (D. Y.). The computations were performed using RIKEN's HOKUSAI BigWaterfall2.

## Notes and references

- B. Schlummer and U. Scholz, *Adv. Synth. Catal.*, 2004, **346**, 1599–1626.
- M. D. Dowle and D. I. Davies, *Chem. Soc. Rev.*, 1979, **8**, 171–197.
- S. Ranganathan, K. M. Muraleedharan, N. K. Vaish and N. Jayaraman, *Tetrahedron*, 2004, **60**, 5273–5308.
- C. K. Tan, L. Zhou and Y.-Y. Yeung, *Synlett*, 2011, **2011**, 1335–1339.
- S. E. Denmark, W. E. Kuester and M. T. Burk, *Angew. Chem., Int. Ed.*, 2012, **51**, 10938–10953.
- U. Hennecke, *Chem.-Asian J.*, 2012, **7**, 456–465.
- W. Zhang, N. Liu, C. M. Schienebeck, K. Decloux, S. Q. Zheng, J. B. Werness and W. P. Tang, *Chem.-Eur. J.*, 2012, **18**, 7296–7305.
- J. M. J. Nolsoe and T. V. Hansen, *Eur. J. Org. Chem.*, 2014, **2014**, 3051–3065.
- A. Sakakura and K. Ishihara, *Chem. Rec.*, 2015, **15**, 728–742.
- M. H. Gieuw, Z. Ke and Y.-Y. Yeung, *Chem. Rec.*, 2017, **17**, 287–311.
- R. Kristianslund, J. E. Tungen and T. V. Hansen, *Org. Biomol. Chem.*, 2019, **17**, 3079–3092.
- H. China, R. Kumar, K. Kikushima and T. Dohi, *Molecules*, 2020, **25**, 6007.
- S. De, A. Kumar Dan, R. Sahu and D. Das, *Eur. J. Org. Chem.*, 2022, **2022**, e202200817.
- S. Liu, B.-Q. Zhang, W.-Y. Xiao, Y.-L. Li and J. Deng, *Adv. Synth. Catal.*, 2022, **364**, 3974–4005.
- L. Liao and X. Zhao, *Acc. Chem. Res.*, 2022, **55**, 2439–2453.





- 16 J. H. Yan, Z. Q. Zhou, Q. Q. He, G. Z. Chen, H. B. Wei and W. Q. Xie, *Org. Chem. Front.*, 2022, **9**, 499–516.
- 17 S. Robin and G. r. Rousseau, *Tetrahedron*, 1998, **54**, 13681–13736.
- 18 M. Breugst, T. Tokuyasu and H. Mayr, *J. Org. Chem.*, 2010, **75**, 5250–5258.
- 19 H. Mayr, M. Breugst and A. R. Ofial, *Angew. Chem., Int. Ed.*, 2011, **50**, 6470–6505.
- 20 S. Knapp, K. E. Rodriques, A. T. Levorse and R. M. Ornaf, *Tetrahedron Lett.*, 1985, **26**, 1803–1806.
- 21 S. Knapp and A. T. Levorse, *J. Org. Chem.*, 1988, **53**, 4006–4014.
- 22 Y.-Y. Yeung and E. J. Corey, *Tetrahedron Lett.*, 2007, **48**, 7567–7570.
- 23 Y.-Y. Yeung, S. Hong and E. J. Corey, *J. Am. Chem. Soc.*, 2006, **128**, 6310–6311.
- 24 A. J. Biloski, R. D. Wood and B. Ganem, *J. Am. Chem. Soc.*, 1982, **104**, 3233–3235.
- 25 Y. A. Cheng, W. Z. Yu and Y. Y. Yeung, *Angew. Chem., Int. Ed.*, 2015, **54**, 12102–12106.
- 26 S. E. Denmark and H. M. Chi, *J. Am. Chem. Soc.*, 2014, **136**, 8915–8918.
- 27 J. L. Panger and S. E. Denmark, *Org. Lett.*, 2020, **22**, 2501–2505.
- 28 M. Hatano and T. Nishimura, *Angew. Chem., Int. Ed.*, 2015, **54**, 10949–10952.
- 29 M. Nagamoto, T. Yanagi, T. Nishimura and H. Yorimitsu, *Org. Lett.*, 2016, **18**, 4474–4477.
- 30 M. Freis, M. Balkenhohl, D. M. Fischer, T. Georgiev, R. C. Sarott and E. M. Carreira, *Org. Lett.*, 2023, **25**, 6380–6384.
- 31 X. Yang, P. Chen and G. Liu, *Angew. Chem., Int. Ed.*, 2024, e202408305.
- 32 F. V. Singh and T. Wirth, *Catal. Sci. Technol.*, 2019, **9**, 1073–1091.
- 33 S. Ortgies and A. Breder, *ACS Catal.*, 2017, **7**, 5828–5840.
- 34 L. Liao and X. Zhao, *Synlett*, 2021, **32**, 1262–1268.
- 35 L. Shao, Y. Li, J. Lu and X. Jiang, *Org. Chem. Front.*, 2019, **6**, 2999–3041.
- 36 J. T. Stadel and T. G. Back, *Chem.–Eur. J.*, 2024, **30**, e202304074.
- 37 Y. Kawamata, T. Hashimoto and K. Maruoka, *J. Am. Chem. Soc.*, 2016, **138**, 5206–5209.
- 38 D. M. Browne, O. Niyomura and T. Wirth, *Org. Lett.*, 2007, **9**, 3169–3171.
- 39 A. Toshimitsu, K. Terao and S. Uemura, *J. Org. Chem.*, 1987, **52**, 2018–2026.
- 40 M. Tiecco, L. Testaferri, M. Tingoli and F. Marini, *J. Chem. Soc., Chem. Commun.*, 1994, 221, DOI: [10.1039/c39940000221](https://doi.org/10.1039/c39940000221).
- 41 M. Tiecco, L. Testaferri, F. Marini, S. Sternativo, L. Bagnoli, C. Santi and A. Temperini, *Tetrahedron: Asymmetry*, 2001, **12**, 1493–1502.
- 42 F. Krättschmar, S. Ortgies, R. Y. N. Willing and A. Breder, *Catalysts*, 2019, **9**, 153.
- 43 T. Lei, S. Graf, C. Schöll, F. Krättschmar, B. Gregori, T. Appleson and A. Breder, *ACS Catal.*, 2023, **13**, 16240–16248.
- 44 Z. Tao, B. B. Gilbert and S. E. Denmark, *J. Am. Chem. Soc.*, 2019, **141**, 19161–19170.
- 45 E. M. Mumford, B. N. Hemric and S. E. Denmark, *J. Am. Chem. Soc.*, 2021, **143**, 13408–13417.
- 46 Y. Otsuka, Y. Shimazaki, H. Nagaoka, K. Maruoka and T. Hashimoto, *Synlett*, 2019, **30**, 1679–1682.
- 47 C. Wata and T. Hashimoto, *J. Am. Chem. Soc.*, 2021, **143**, 1745–1751.
- 48 C. Wata and T. Hashimoto, *Synthesis*, 2021, **53**, 2594–2601.
- 49 Y. Oe, R. Yoshida, A. Tanaka, A. Adachi, Y. Ishibashi, T. Okazoe, K. Aikawa and T. Hashimoto, *J. Am. Chem. Soc.*, 2022, **144**, 2107–2113.
- 50 H. Kimura, K. Aikawa and T. Hashimoto, *Asian J. Org. Chem.*, 2023, **13**, e202300522.
- 51 H. Cui, Y. Lin, M. Luo, Y. Lu, X. Huang and Z. She, *Org. Lett.*, 2017, **19**, 5621–5624.
- 52 A. Kütt, S. Tshepelevitsh, J. Saame, M. Lõkov, I. Kaljurand, S. Selberg and I. Leito, *Eur. J. Org. Chem.*, 2021, **2021**, 1407–1419.
- 53 R. G. Parr and W. Yang, *J. Am. Chem. Soc.*, 1984, **106**, 4049–4050.
- 54 W. Yang and W. J. Mortier, *J. Am. Chem. Soc.*, 1986, **108**, 5708–5711.
- 55 S. Liu, C. Rong and T. Lu, *J. Phys. Chem. A*, 2014, **118**, 3698–3704.
- 56 B. Wang, C. Rong, P. K. Chattaraj and S. Liu, *Theor. Chem. Acc.*, 2019, **138**, 124.
- 57 T. Lu and F. Chen, *J. Comput. Chem.*, 2012, **33**, 580–592.
- 58 T. Lu, *J. Chem. Phys.*, 2024, **161**, 082503.
- 59 Y. A. Cheng, W. Z. Yu and Y. Y. Yeung, *J. Org. Chem.*, 2016, **81**, 545–552.
- 60 T. Arai, O. Watanabe, S. Yabe and M. Yamanaka, *Angew. Chem., Int. Ed.*, 2015, **54**, 12767–12771.
- 61 T. Gensch, G. dos Passos Gomes, P. Friederich, E. Peters, T. Gaudin, R. Pollice, K. Jorner, A. Nigam, M. Lindner-D'Addario, M. S. Sigman and A. Aspuru-Guzik, *J. Am. Chem. Soc.*, 2022, **144**, 1205–1217.
- 62 C. H. Ng, E. R. Wearing, D. E. Blackmun, G. G. Terrones, J. Toigo, K.-M. Tong, K. C. Harper, J. C. Donovan, M. A. Gonley, E. A. Voight, B. D. Bergstrom, M. O. Wolf, H. J. Kulik and C. S. Schindler, *J. Am. Chem. Soc.*, 2025, **147**, 29722–29731.
- 63 C. Zheng, G.-Z. Wang and R. Shang, *Adv. Synth. Catal.*, 2019, **361**, 4500–4505.
- 64 N. Slavov, J. Cvengroš, J.-M. Neudörfl and H.-G. Schmalz, *Angew. Chem., Int. Ed.*, 2010, **49**, 7588–7591.
- 65 E. Merifield and E. J. Thomas, *J. Chem. Soc., Chem. Commun.*, 1990, 464, DOI: [10.1039/c39900000464](https://doi.org/10.1039/c39900000464).
- 66 M. J. Bamford, M. Beard, T. Cherry David and M. G. Moloney, *Tetrahedron: Asymmetry*, 1995, **6**, 337.
- 67 J. H. Bailey, D. T. Cherry, K. M. Crapnell, M. G. Moloney, S. B. Shim, M. J. Bamford and R. B. Lamont, *Tetrahedron*, 1997, **53**, 11731–11744.
- 68 K. Yoshida, T. Morikawa, N. Yokozuka, S. Harada and A. Nishida, *Tetrahedron Lett.*, 2014, **55**, 6907–6910.



- 69 K. R. Owens, S. V. McCowen, K. A. Blackford, S. Ueno, Y. Hirooka, M. Weber and R. Sarpong, *J. Am. Chem. Soc.*, 2019, **141**, 13713–13717.
- 70 M. Szostak, M. Spain and D. J. Procter, *Chem. Soc. Rev.*, 2013, **42**, 9155–9183.
- 71 T.-H. Chou, B.-H. Yu and R.-J. Chein, *Chem. Commun.*, 2019, **55**, 13522–13525.
- 72 R. Maji, S. Ghosh, O. Grossmann, P. Zhang, M. Leutzsch, N. Tsuji and B. List, *J. Am. Chem. Soc.*, 2023, **145**, 8788–8793.
- 73 N. Luo, M. Turberg, M. Leutzsch, B. Mitschke, S. Brunen, V. N. Wakchaure, N. Nöthling, M. Schelwies, R. Pelzer and B. List, *Nature*, 2024, **632**, 795–801.
- 74 P. N. Craig, *J. Am. Chem. Soc.*, 1952, **74**, 129–131.
- 75 (a) CCDC 2482628: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pbc6](https://doi.org/10.5517/ccdc.csd.cc2pbc6); (b) CCDC 2482629: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pbct7](https://doi.org/10.5517/ccdc.csd.cc2pbct7); (c) CCDC 2482630: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pbcv8](https://doi.org/10.5517/ccdc.csd.cc2pbcv8); (d) CCDC 2482631: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pbcw9](https://doi.org/10.5517/ccdc.csd.cc2pbcw9); (e) CCDC 2482632: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pbcbx](https://doi.org/10.5517/ccdc.csd.cc2pbcbx); (f) CCDC 2484902: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pdr40](https://doi.org/10.5517/ccdc.csd.cc2pdr40).

