



Showcasing the Highlight article of Miss Namrata Kotwal and Professor Pankaj Chauhan, Department of Chemistry, Indian Institute of Technology Jammu, India

Evolution in the asymmetric synthesis of biaryl ethers and related atropisomers

This article highlights the seminal contributions towards the catalytic enantioselective synthesis of dual axial atropisomeric biaryl ethers and related amines.

As featured in:



See Namrata Kotwal and Pankaj Chauhan, *Chem. Commun.*, 2024, **60**, 6837.



Cite this: *Chem. Commun.*, 2024, 60, 6837

Received 9th April 2024,
Accepted 14th May 2024
DOI: 10.1039/d4cc01655f

rsc.li/chemcomm

Evolution in the asymmetric synthesis of biaryl ethers and related atropisomers

Namrata Kotwal and Pankaj Chauhan *

Axially chiral biaryl ethers and related compounds hold valuable potential in natural products, medicinal chemistry, and catalysis; however, their asymmetric syntheses have always been overlooked compared to other biaryl/hetero-biaryl atropisomers. Unlike the later class molecules bearing a single chiral axis, the former category possesses a unique type of atropisomerism bearing two potential axes. Due to their great importance in diverse research domains, catalytic atropselective biaryl ether synthesis has recently witnessed an upsurge. This highlight article provides an elaborated view on the developments of catalytic synthetic methods that have been explored to achieve dual axial chirality in biaryl ethers and related scaffolds.

1. Introduction

Biaryls/hetero-biaryls represent an essential class of organic molecules, frequently resolved as atropisomers due to the restricted rotation around the pivotal bond, specifically around

the carbon–carbon, carbon–nitrogen, and nitrogen–nitrogen bonds. The significance of atropisomeric biaryls and related hetero-biaryls has long been recognised due to their widespread presence in natural products,¹ drug molecules,² privileged organocatalysts³ and chiral ligands in metal catalysis.⁴ Considering the diverse applications and interesting properties, substantial progress has been witnessed in the atropselective synthesis of these biaryls/hetero-biaryls bearing a C2 axis of symmetry.⁵ Besides, another essential and fascinating class of

Department of Chemistry, Indian Institute of Technology Jammu, Jagti, NH-44, Nagrota Bypass, Jammu, 181221 J&K, India.
E-mail: pankaj.chauhan@iitjammu.ac.in; Web: <https://www.pankajchauhanresearch.com>



Namrata Kotwal

Namrata Kotwal was born in 1997 in Jammu, India. She obtained her BSc in 2017 and MSc Chemistry in 2019 from the University of Jammu. Since August 2020, she has been working as a PhD student in the research group of Dr Pankaj Chauhan at IIT Jammu. Her research interests include organocatalytic asymmetric synthesis of medium-sized carbo- and heterocyclic bridged biaryls, and the generation of stereogenic centres at the remote position of the unsaturated acceptors.



Pankaj Chauhan

Pankaj Chauhan obtained his BSc from Himachal Pradesh University in 2004 and MSc Chemistry from Guru Nanak Dev University, Amritsar, in 2007. He completed his PhD under the supervision of Prof. S. S. Chimni at Guru Nanak Dev University in 2012. In April 2013, he joined the research group of Professor Dieter Enders as a postdoctoral researcher at RWTH Aachen University, Germany. He worked as a sub-group leader in the same research group from April 2014 to September 2017. In October 2017, he started his independent research career as an Assistant Professor in the Department of Chemistry, Indian Institute of Technology Jammu, where he is currently working as an Associate Professor. His research interests include asymmetric synthesis, organocatalysis, photocatalysis and direct photoactivation, electroorganic synthesis and synergistic catalysis.

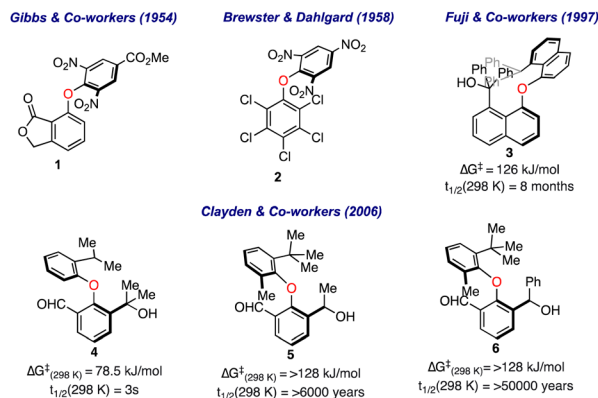
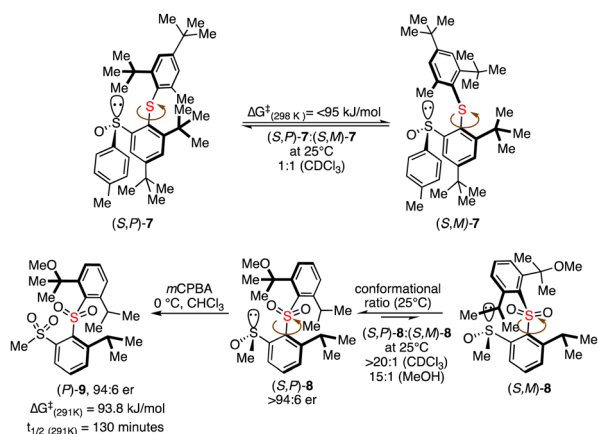


Fig. 2 Early studies on the restricted rotation in biaryl ethers.

Dahlgard performed IR and UV spectral studies on more than three dozen differently substituted biaryl ethers (including compound 2) and found a significant weakening in the absorption bands in IR and UV regions for the larger substituent at the *ortho* positions.¹⁴ Other precedents also appeared later, where the variable temperature NMR investigations were carried out to determine the rotation barriers in biaryl ethers.

In 1998, Fuji and co-workers reported the synthesis of optically active 8,8'-disubstituted 1,1'-binaphthyl ethers and calculated the rotation barrier based on the time-dependent alteration in the ee values.¹⁵ It was realized that the sterically crowded biaryl ether 3 has a half-life for racemization ($t_{1/2}$) at 25 °C of 8 months and sufficient activation barrier ($\Delta G_{\text{rot}} = 126 \text{ kJ mol}^{-1}$) to be resolved as an atropisomer. Later, Clayden's group further investigated the phenomenon of atropisomerism among diphenyl ethers with unsymmetrical substitution.¹⁶ It was observed that the free energy barrier was enhanced with an increase in steric bulk around the aromatic rings, as shown for compounds 4 to 6, where a significant atropisomerism is only apparent when the *tert*-butyl group is present.

Clayden and co-workers also successfully established the phenomenon of atropisomerism in biaryl sulfides and biaryl sulphones (Scheme 1).¹⁷ A separable diastereoisomeric sulfinyl sulfides 7 were detected in ¹H NMR at 20 °C with a 1 : 1 ratio;



Scheme 1 Atropisomerism in biaryl sulfides and biaryl sulphones.

however, a rapid interconversion was observed due to a lower energy barrier at ambient temperature. The conformationally and enantio-pure sulfinyl sulfone 8 could be converted to bis sulfone 9, possessing only biaryl axes without central chirality by maintaining the high enantiomeric excess.

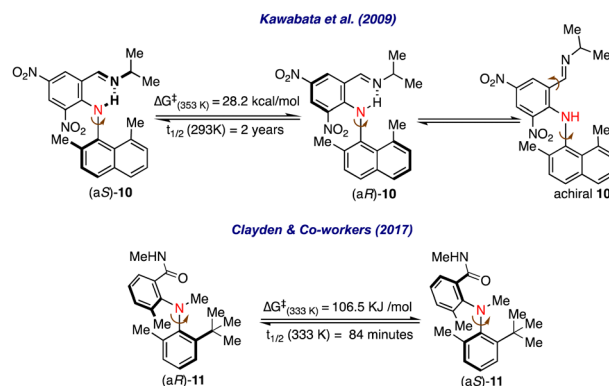
A similar atropisomerism could be witnessed in biaryl amines.¹⁸ In this direction, Kawabata *et al.* investigated atropisomerism in a series of biaryl amines having the possibility of intramolecular H-bonding.¹⁹ It was revealed that in biaryl-substituted amine 10, a stable axial chirality arises due to intramolecular N–H–N hydrogen-bonding interactions, and the hydrogen bond gets stronger by substituting donor groups on the imine nitrogen and electron-withdrawing groups on the naphthyl ring (Scheme 2). Further, it was realized that substituting the naphthyl ring with sterically demanding groups effectively increased the rotational barrier around the C–N pivotal bond, thus enabling stable axial chirality among these molecules.

Clayden and co-workers were able to resolve the atropisomers in biaryl amines without intramolecular H-bonding on HPLC by using a chiral stationary phase.²⁰ A significant rotation barrier and half-life of racemization were calculated by observing first-order decay in enantioselectivity for the biaryl amines 11.

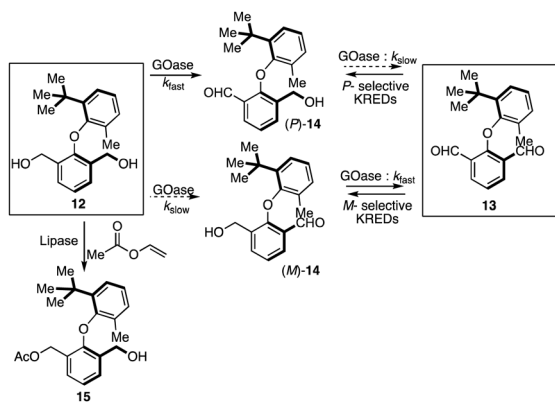
3. Catalytic atropselective synthesis of biaryl ethers

3.1 Biocatalytic approach

Clayden's group provided the first catalytic pathway for accessing enantiopure biaryl ethers either *via* the biocatalytic oxidative or reductive desymmetrization of symmetrical diol 12 or dialdehyde 13, respectively (Scheme 3).²¹ With a variant of galactose oxidase (GOase), a benzylic diol 12 provided the biaryl ether (*P*)-14 with 88% ee in an hour. The ee value was further enriched to a maximum of 94% due to converting the minor enantiomer (*M*)-14 to the dialdehyde 13. This enantio-enrichment was attributed to combining both enantioselective desymmetrization and kinetic resolution (KR). Further, an initial attempt for the selective acylation of diol 12 with a lipase and vinyl acetate provided moderate enantioselectivity (60% ee)



Scheme 2 Atropisomerism in biaryl amines.



Scheme 3 Biocatalytic oxidative or reductive desymmetrization of symmetrical biaryl ether diol or dialdehyde.

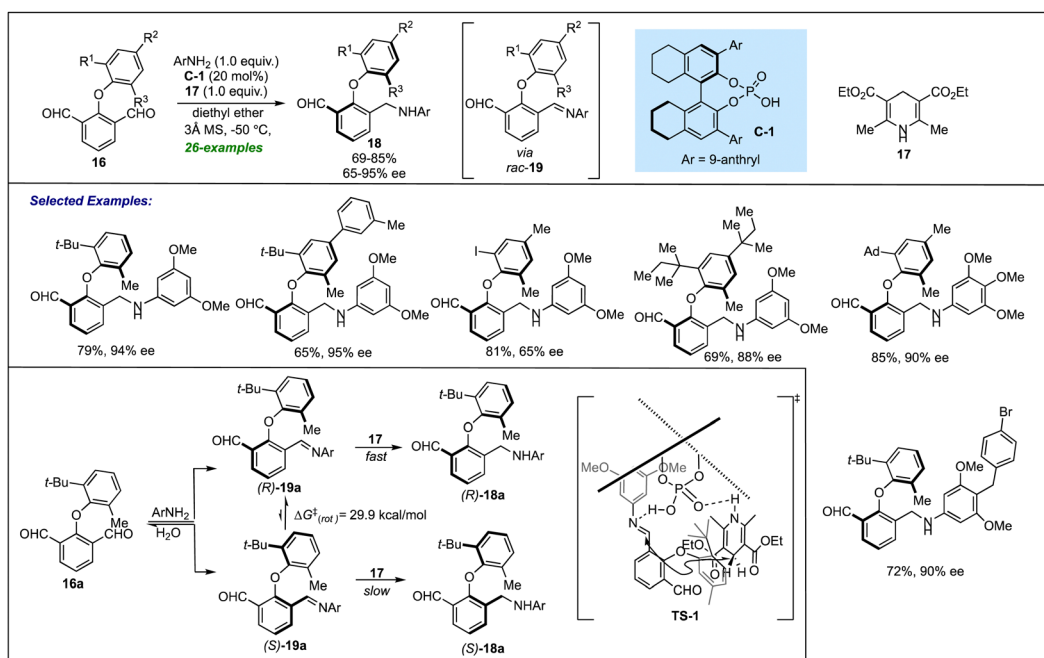
of **15** with approximately 50% conversion in 24 hours. Alternatively, a series of ketoreductases (KREDs) were screened for the reduction of dialdehyde **13** to reach the enantiomer of (*P*)-**14** selectively. In the enzymatic reduction, most of the screened KREDs were found to be active; however, the one showing perfect enantio-induction resulted in poor conversion.

3.2. Organocatalytic approaches

3.2.1 Chiral Brønsted acid catalysis for atropselective synthesis of axially chiral biaryl ethers. After Clayden discovered the biocatalytic access to enantiopure ethers, it took more than a decade to see another catalytic approach leading to the biaxial biaryl atropisomers. In 2023, research groups of Zeng²² and Yang²³ independently published two works exploring the catalytic potential of chiral Brønsted acid to achieve the

atropselective synthesis of axially chiral biaryl ethers. Zeng and co-workers explored a dynamic kinetic resolution (DKR) approach for accessing these atropisomers, starting from symmetrical dicarbaldehydes **16** and aniline derivatives (Scheme 4).²² The chiral phosphoric acid **C-1** was identified as the best catalyst for the asymmetric transfer hydrogenation (ATH) of *in situ* generated imines **19** with Hantzsch ester **17** as a reductant to afford biaryl ethers **18**. During substrate scope investigation, it was observed that the aryl anilines with 3,5-dimethoxy substituents resulted in high enantioselectivity. Replacing one of these methoxy groups with hydrogen, methyl and chloro substituents did not affect the results in terms of yields; however, the enantioselectivities were slightly diminished. On the other hand, dicarbaldehydes bearing electron-donating groups and halogens were well tolerated for this protocol and afforded the desired products **18** in good yields and excellent enantioselectivities. The steric crowding at one of the *ortho* positions with alkyl groups such as *tert*-butyl, *tert*-pentyl and adamantyl was found to be crucial for high enantioselectivity, where an iodo substituent resulted in lower enantioselectivity of 65% ee.

A deep insight into the reaction mechanism revealed that in the absence of Hantzsch ester, the corresponding imine was generated in good yield but in <5% ee. This clearly indicated that even in the presence of a chiral catalyst, no enantio-induction was provided by the imine formation step. Further, computational studies combined with experimental observations led to the conclusion that *in situ* generated (*R*)-**19** undergoes a fast ATH reaction catalyzed by **C-1** to provide good enantioselectivity of the desired amine **18a**. Conversely, another conformer (*S*)-**19** with a slow reaction rate hydrolyzes to recover **16a** dynamically. A stereochemical model (**TS-1**) was also



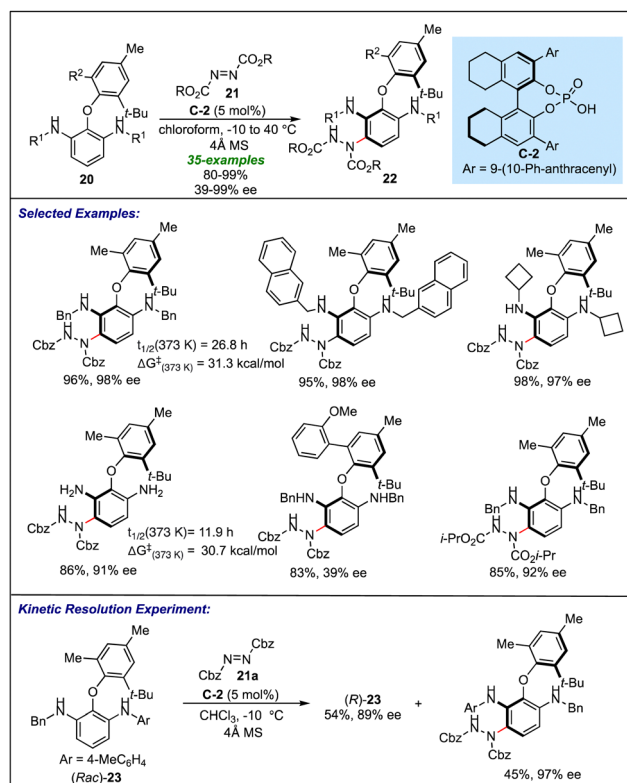
Scheme 4 Phosphoric acid catalyzed DKR via ATH of the *in situ* generated imines for accessing atropisomeric biaryl ethers.

proposed for the transformation, indicating the role of a chiral catalyst for the atropselective synthesis of biaryl ethers.

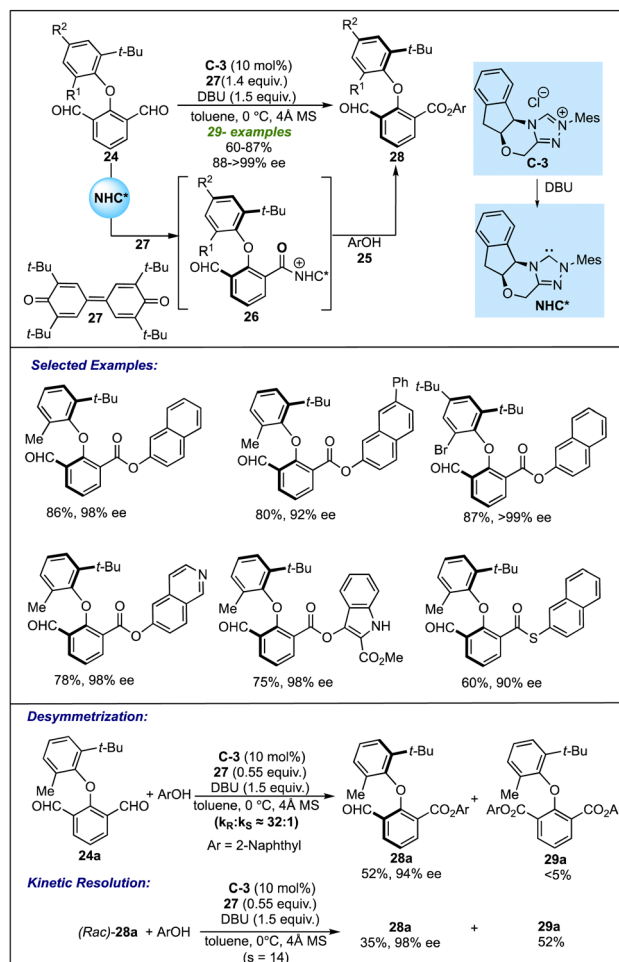
Yang's group explored the applications of chiral phosphoric acid catalysis to efficiently synthesise axially chiral biaryl ethers through a desymmetrization of symmetrical biaryl ether appended aryl amines **20** (Scheme 5).²³ Using C-2 as a catalyst, the asymmetric electrophilic amination of the 2-aryloxy-substituted 1,3-benzenedimaines **20** with azodicarboxylate **21** resulted in the broad range of enantiopure desymmetrized biaryl ethers **22**. It was found that the substrates with various *N*-benzyl and *N*-alkyl substituents (with both cyclic and acyclic systems) at the R¹ position resulted in excellent results in terms of yield and enantioselectivity. Interestingly, a free amine also undergoes amination reactions with high enantioselectivity (91% ee). Many alkyl and aryl groups, as well as a benzyl group on one of the aryl rings of biaryl ethers (at R²), were found to be tolerable, except the sterically crowded substrate (R² = 2-MeOC₆H₄), which resulted in only 39% ee. Apart from a good acceptance of different substituents on different positions of the biaryl ethers, many commercially available azodicarboxylates were also examined, where the atropisomers of biaryl ethers were accessed in the highly enantioselective fashion except for di-*tert* butyl azodicarboxylates, which proved to be the unsuitable substrate for this transformation due to the steric reasons. The configurational stability of the biaryl ether products was further confirmed by thermal racemization experiments, where products were found to have sufficient

racemization half-life and a high rotation barrier. Kinetic resolution of an unsymmetrical starting amine **23** occurs with high efficiency, suggesting the versatility of this amination process for delivering diverse atropisomeric biaryl ethers.

3.2.2 Chiral NHC-catalysis for atropselective synthesis of axially chiral biaryl ethers. Recently, four independent related works were published one after another on the NHC-catalysed desymmetrization reactions leading to atropselective synthesis of axially chiral biaryl ethers. Biju's group was the first to disclose a straightforward atropselective synthesis of axially chiral dialdehydes with an NHC-catalysed desymmetrization of the 2-aryloxyisophthalaldehydes **24** with various aryl alcohols **25** (Scheme 6).²⁴ This desymmetrization process involves the oxidative generation of acylazolium intermediate **26** from the substrates **24** with the assistance of NHC generated from triazolium salt C-3 and an external oxidant **27**. The resulting intermediate **26** is trapped by suitable alcohol to yield desymmetrized axially chiral biaryl ethers **28**. It was found that irrespective of the substitution pattern and nature of the group on the 2-naphthols, efficient esterification of the dialdehydes **24** produced the corresponding axially chiral biaryl ethers in



Scheme 5 Phosphoric acid catalyzed desymmetrization via electrophilic amination of aryl amines for accessing atropisomeric biaryl ethers.



Scheme 6 Biju's work on the NHC-catalyzed desymmetrization for atropselective synthesis of biaryl ethers.

Highlight

excellent ee-values. Not only 2-naphthols but also 1-naphthol and other alcohols bearing larger aryl rings, such as 9-phenanthrene; and heteroaryl alcohols (a 3-indanol derivative and quinolinol), sesamol and 2-naphthalene thiols proved to be suitable nucleophiles for this transformation without much impacting the yield and enantioselectivities of the corresponding products.

The configurational stability of the product was depicted with thermal stability experiments and DFT calculations. Both investigations revealed that compound **28a** has a significant rotation barrier to be resolved as an atropisomer. Further insight into the mechanism revealed that lowering the equivalents of oxidant under optimized conditions led to the formation of the desired product in 52% yield and 94% ee, along with the formation of diester **29** as a by-product in traces. On the other hand, the reaction of racemic product **28a** with 0.55 equivalent of oxidant resulted in kinetic resolution leading to enantiopure **28a** in 35% yield and diester **29** in 52% yield. These observations suggested that desymmetrization plays a major role in controlling enantioselectivity; however, the possibility of further enantio-enrichment by a subsequent KR process could not be eliminated completely.²⁵

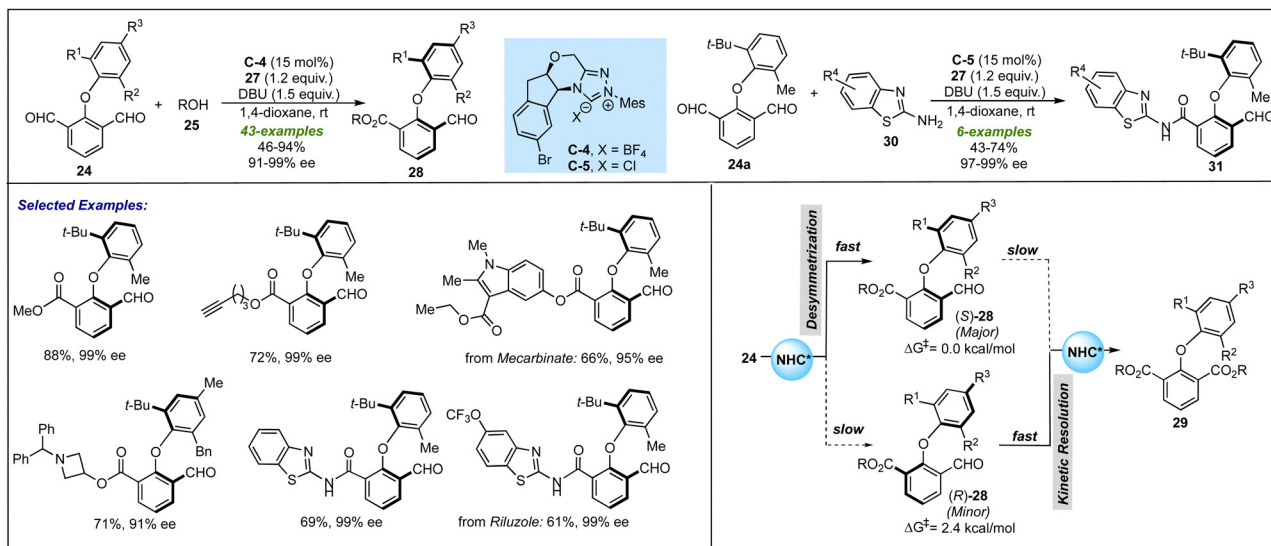
Parallely, Ye and co-workers disclosed similar NHC-catalyzed desymmetrization of 2-aryloxyisophthalaldehydes **24** with alcohols **25** and amines **30** (Scheme 7).²⁶ It was proposed that the NHC generated from **C-4** or **C-5** oxidatively provide azolium intermediate **26** from **24**, which undergoes esterification or amidation to furnish asymmetrical biaryl ethers **28** and amides **31** with excellent enantioselectivities. By using CD₃OD as a nucleophile, it was observed that deuterium incorporation did not occur in the remaining formyl group, which suggests the non-reversible formation of the Breslow intermediate. Based on similar controlled experiments as performed by Biju's group, the enantio-enrichment was proposed to be further assisted by a downstream kinetic resolution process in this desymmetrization

process. The key features of this protocol include broad functional group tolerance and late-stage functionalizations under ambient reaction conditions. Apart from phenol derivatives as nucleophiles, the successful applications of aliphatic alcohols and amine nucleophiles, *i.e.* 2-aminobenzothiols **30**, could be seen as further advancement in the NHC-catalyzed desymmetrization leading to biaxial biaryl ethers.

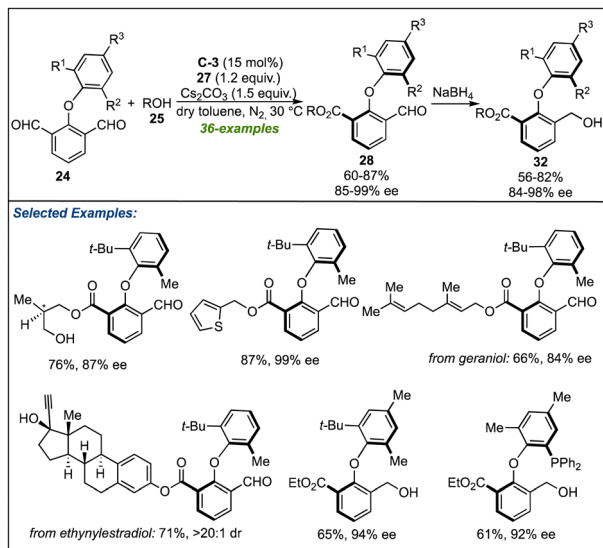
Soon after the works of Biju's and Ye's lab, the research groups of Zeng²⁷ and Gao²⁸ designed similar NHC-catalysed strategies to synthesize the biaxial biaryl ethers. Using **C-3** as NHC-pre-catalyst and Cs₂CO₃ as a base assisted the above-described desymmetrization of isophthalaldehyde derivatives **24** with a broad range of alcohols (Scheme 8).²⁷ Again, an excellent level of enantioselectivity of esters **28** or alcohols **32** (formed by reduction of **28**) was observed. With the triazolium salt **C-6** as an NHC precursor, desymmetrization of isophthalaldehyde derivatives **24** with a wide variety of aliphatic alcohols and some phenol derivatives yielded the desired biaryl ethers in good to excellent enantioselectivities (Scheme 9).²⁸ The last two strategies are important not only in the context of synthetic methods leading to atropisomeric biaryl ethers but also in the utilization of corresponding biaxial ethers in chiral catalysis.

3.3. Transition-metal catalyzed approach

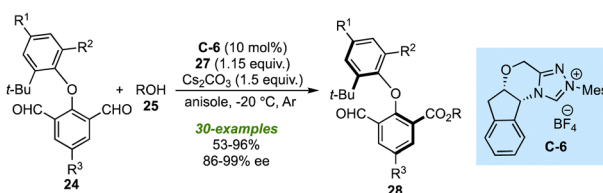
Recently, Yao and co-workers applied transition-metal catalysis to provide direct access to the dual axial chirality among biaryl ether scaffolds. A new class of bisalkyne-based biaryl ethers **33** has been developed for Cu-catalyzed azide-alkyne Huisgen cycloaddition (CuAAC), leading to desymmetrization of the substrate (Scheme 10).²⁹ An indane fused BOX ligand **L-1**, with the assistance of copper metal, catalyses the cycloaddition reaction between bis alkynes and azides to give a diverse array of axially chiral C–O atropisomers **34** in excellent yields and enantioselectivities. The developed methodology accommodates diverse substitution patterns irrespective of the nature



Scheme 7 Ye's work on the NHC-catalyzed desymmetrization for atropselective synthesis of biaryl ethers.



Scheme 8 Zeng's work on the NHC-catalyzed desymmetrization for atropselective synthesis of axially chiral biaryl ethers.

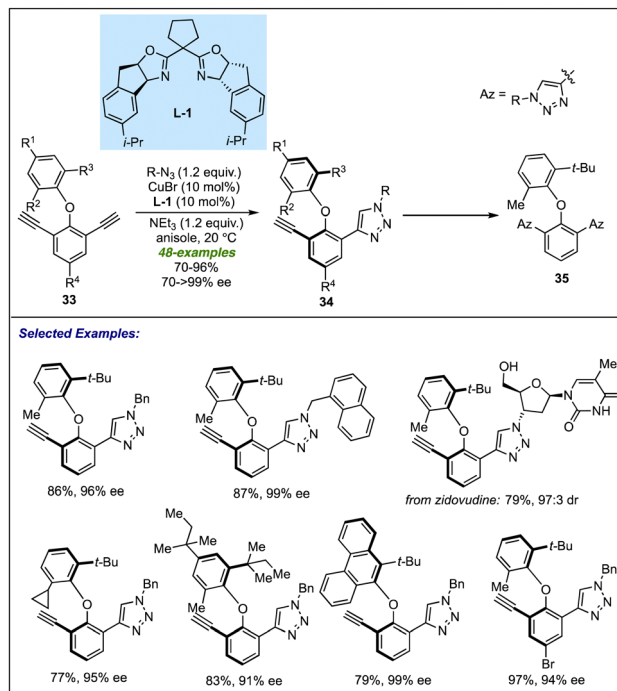


Scheme 9 Gao's work on the NHC-catalyzed desymmetrization for atropselective synthesis of axially chiral biaryl ethers.

and position of groups on the bisalkynes. The cycloaddition reaction proceeded smoothly with bio-active molecules such as zidovudine to furnish the desired desymmetrized product in 79% yield and 97 : 3 dr. It was observed that steric bulk at the C2 position is necessary to provide satisfactory results as replacing the bulky group (*tert*-Bu, *tert*-pentyl, *etc.*) with ethyl or isopropyl greatly affected the enantioselectivity of the corresponding products (18% ee with isopropyl and racemic with ethyl). The control experiment revealed that a prolonged reaction time of 24 hours resulted in the formation of di-triazole product 35 of 21% *via* a double CuAAC with a slight enantioselectivity enhancement (96% ee to 99% ee) of 34. This result suggests that the enantio-induction is primarily governed by the desymmetrization step, with little role of the KR.

4. Catalytic atropselective synthesis of related ethers

Although a significant upsurge has been seen in the last year for procuring biaryl ethers, it was in 2018 when Gustafson's group laid a foundation for the catalytic atropselective synthesis of such biaxially chiral molecules. Their work was focused on the enantioselective synthesis of naphthoquinone-based aryl

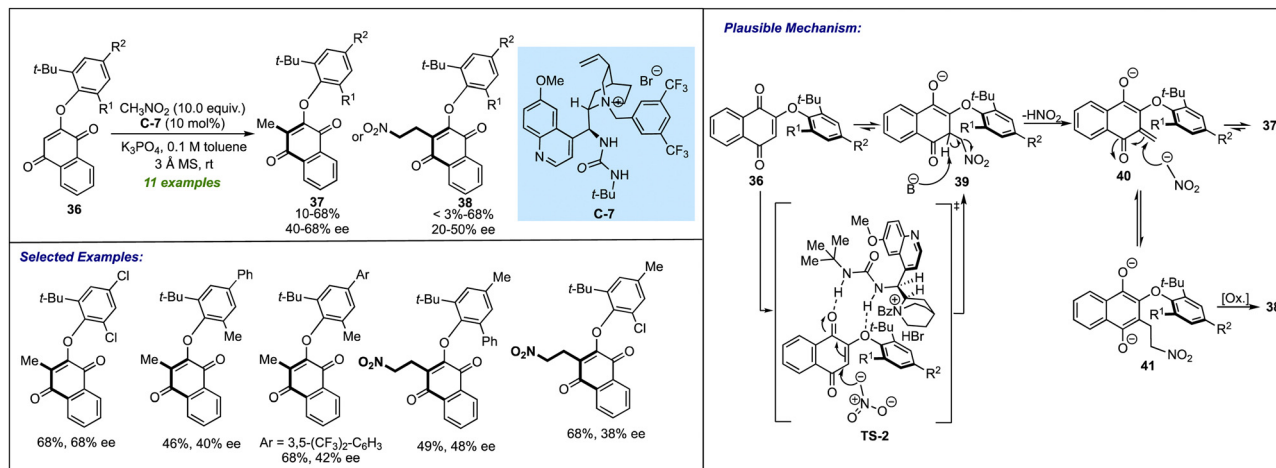


Scheme 10 CuAAC for atropselective synthesis of axially chiral biaryl ethers.

ethers (Scheme 11).³⁰ By using cinchona-derived urea-bearing quaternary ammonium salt C-7, direct access to atropisomeric biaryl ethers 37 or 38 has been achieved through C(sp²)-H alkylation of substrate 36 with nitromethane or by a similar C(sp²)-H alkylation followed by second nitromethane addition. The C-7 assisted 1,4-nucleophilic addition of nitromethane to quinones 36, followed by the elimination of nitrous acid, resulting in atropselective synthesis of quinone-based biaryl ethers 37. Among the screened substrates, in some instances, the nitroethylated products 38 were isolated as a major product in decent yields and moderate enantioselectivities. In these cases, after eliminating nitrous acid from intermediate 39, the enone 40 (if having a sufficiently longer lifetime) undergoes another Michael-type addition to provide 41, which, after oxidation, results in the corresponding nitroethylated product 38. Notably, after one round of titration, the enantioselectivity of both products has been enriched to an excellent level. The stereochemical model for the formation of the desired product revealed that there was hydrogen bonding between the urea moiety of the catalyst with the ether oxygen and one of the carbonyl carbons of the low-energy conformer of quinone substrate. As a result of this, the quinone-based biaxial biaryl ether locks into feasible (*as*)-*exo* atropisomers.

5. Catalytic atropselective synthesis of related amines

Despite being recognized as potential chemotypes in medicinal chemistry and many FDA-approved drugs,¹² the atropselective synthesis of biaryl amines has been completely overlooked.



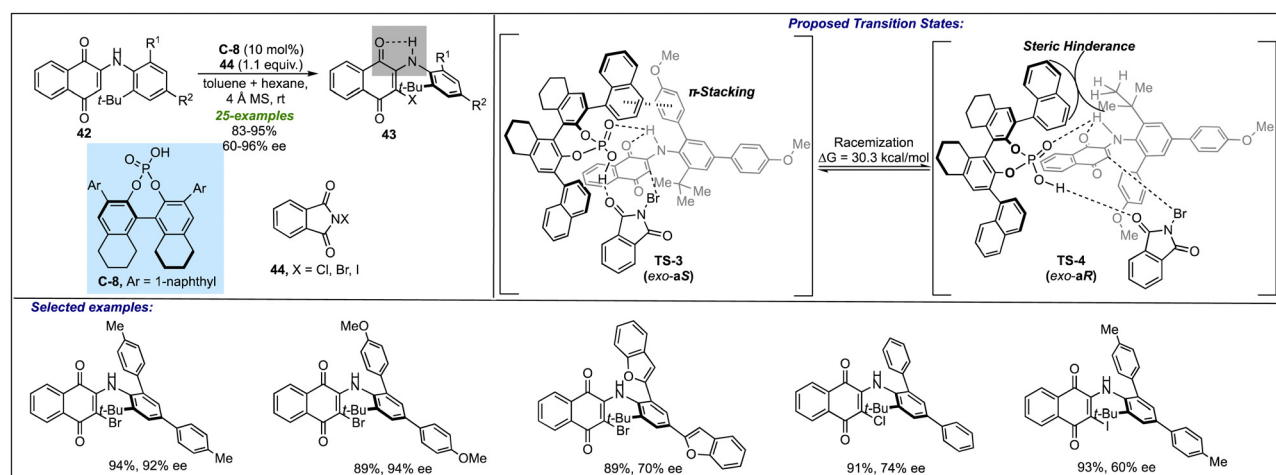
Scheme 11 Atropselective synthesis of naphthoquinone-based aryl ethers.

Besides the early efforts of Kawabata¹⁹ and Clayden²⁰ towards establishing the axial chirality among biaryl amines, Gustafson's group devised the atropselective synthesis of *N*-aryl quinoids-based biaryl amines **43** (Scheme 12).³¹ Their work disclosed a chiral phosphoric acid **C-8** catalyzed halogenation of *N*-aryl quinoids **42**, yielding the desired product **43** in excellent yields and enantioselectivities. Under ambient conditions, *N*-aryl quinoid-derived biaryl amines were found to be exhibiting strong N–H–O hydrogen bonding interactions, resulting in high stereochemical stability of the *exo*-product ($\Delta G^\ddagger = 30 \text{ kcal mol}^{-1}$) over *endo*-product in both protic and aprotic solvents. During the reaction, it was observed that replacing aryl substitution at *R*¹ and *R*² by methyl groups suddenly resulted in an immense loss of ee value (20% ee). This is mainly attributed to the fact that π – π -stacking between the catalyst and aryl rings of the substrates at the *R*¹ and *R*² positions play a crucial role in leading to high enantiomeric excess. It was also found that decreasing the bulk by replacing the *tert*-butyl group with *iso*-propyl and trifluoromethyl groups drastically decreased the enantioselectivity to 28% ee and 12% ee, respectively, probably due to a low barrier of racemization. It was postulated that at

onset, the **C-8** catalyst is involved in hydrogen bonding with the stable *exo*-conformation of the quinoid substrate (**TS-3**). The resulting hydrogen bonding increases the energy of HOMO, *i.e.* the enamine-like amino naphthoquinone. The rotation around the C–N axis allows the positioning of the *t*-Bu group to be far away from the catalyst, thus organizing into (*aS*)-conformation. This conformation allows the π – π -stacking of the *ortho*-aryl group with the naphthyl group of the catalyst. Simultaneously, the acidic proton of **C-8** activates the electrophilic halogenating reagent (NXS) *via* H-bonding, thus directing the halogenation towards the naphthoquinone site, and the axis will be further reinforced, leading to the selective formation of the product with (*aS*)-configuration, whereas the formation of (*aR*)-product is limited due to steric crowding and lack of π – π -stacking (**TS-4**).

Conclusions & outlook

Widespread applications and interesting properties of atropisomers have led to the discovery of new synthetic methods for



Scheme 12 Atropselective synthesis of naphthoquinone-based aryl amines.

selectively synthesizing these classes of compounds. This highlight article has summed up the importance, early precedents and asymmetric methods leading to a unique class of biaryl atropisomers *viz* biaryl ethers and related amines. Whereas the evidence of atropisomers in such molecules has long been recognized, their asymmetric synthesis has gained recognition only recently. In this context, desymmetrization reactions under the action of chiral catalysts have emerged as an efficient strategy. In most cases, the desymmetrization step imparts the desired enantio-differentiation; however, the role of KR has also been observed in improving the enantioselectivity. Apart from the first example on the catalytic pathway leading to enantioselective synthesis of biaryl ethers by using enzymes, recently, the potential of organocatalysts such as chiral Brønsted acid and NHC have been explored for accessing such biaxial biaryls. In addition, a transition metal-catalyzed desymmetrization/KR process has been developed to achieve dual axially chiral biaryl ethers. Regardless of the major challenges due to flexible C–X {X = O, N, S} axes and the prolonged distance between the groups imparting restricted rotations, all the seminal contributions discussed above provide excellent enantioselectivities of the biaryl ethers with a sufficiently longer half-life of racemization. Impressively, diverse precursors with wide functional group tolerance, including biologically relevant compounds, have been well accommodated under mild reaction conditions. In a couple of cases, the applications of newly developed enantiopure diaryl ethers have been successfully explored as chiral catalysts.

Though the evidences of dual axial chirality due to the hindrance in rotation were apparent for related biaryl amines and sulfides, nothing much has been seen in terms of asymmetric methods for accessing these scaffolds. At this stage, we believe that these current developments will open a new area in asymmetric synthesis,³² which will not only be limited to the atropselective synthesis of biaryl ethers but may extend up to the enantioselective synthesis of dual axial biaryls bearing C–N–C, C–S–C, C–P–C linkage, *etc.* The enantioselective synthesis of these biaxial biaryls is also expected to extend beyond the desymmetrization process involving the site-specific modifications of a selected aryl ring or the substituent attached to the aryl ring. Beyond desymmetrization, at this stage, we can anticipate that the developing asymmetric methods relying on coupling reactions or direct C–H functionalization may also lead to the atropselective synthesis of biaryl ethers and related molecules.

Author contributions

N. Kotwal and P. Chauhan have contributed to the original draft preparation, editing, and proofreading of the manuscript of this highlight article.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgements

We are highly grateful to the researchers who have contributed to developing synthetic methods for accessing the enantiopure biaryl ethers and related scaffolds. This activity was supported by the Science & Engineering Research Board (SERB) India (File Number: CRG/2022/003212). N. K. thanks the Department of Science & Technology, India, for the INSPIRE fellowship.

References

- 1 J. E. Smyth, N. M. Butler and P. A. Keller, *Nat. Prod. Rep.*, 2015, **32**, 1562–1583.
- 2 (a) S. R. LaPlante, P. J. Edwards, L. D. Fader, A. Jakalian and O. Huckle, *ChemMedChem*, 2011, **6**, 505–513; (b) S. R. LaPlante, L. D. Fader, K. R. Fandrick, D. R. Fandrick, O. Huckle, R. Kemper, S. P. F. Miller and P. J. Edwards, *J. Med. Chem.*, 2011, **54**, 7005–7022.
- 3 (a) J. M. Lassaletta, *Atropisomerism and Axial Chirality*, World Scientific Publishing, Singapore, 2019; (b) D. Parmar, E. Sugiono, S. Raja and M. Rueping, *Chem. Rev.*, 2014, **114**, 9047–9153.
- 4 (a) Y. Chen, S. Yekta and A. K. Yudin, *Chem. Rev.*, 2003, **103**, 3155–3212; (b) Y. M. Li, F. Y. Kwong, W. Y. Yu and A. S. C. Chan, *Coord. Chem. Rev.*, 2007, **251**, 2119–2144.
- 5 For selected recent reviews on asymmetric synthesis of atropisomers: (a) J. W. Delord, A. Panossian, F. R. Leroux and F. Colobert, *Chem. Soc. Rev.*, 2015, **44**, 3418–3430; (b) J. K. Cheng, S. H. Xiang, S. Li, L. Ye and B. Tan, *Chem. Rev.*, 2021, **121**, 4805–4902; (c) N. Kotwal, Tamanna and P. Chauhan, *Chem. Commun.*, 2022, **58**, 11031–11044; (d) P. R. Salamanca, R. Fernandez, V. Hornillos and J. M. Lassaletta, *Chem. – Eur. J.*, 2022, **28**, e202104442; (e) X. F. Bai, Y. M. Cui, J. Cao and L. W. Xu, *Acc. Chem. Res.*, 2022, **55**, 2545–2561; (f) H. H. Zhang, T. Z. Li, S. J. Liu and F. Shi, *Angew. Chem., Int. Ed.*, 2024, **63**, e202311053.
- 6 For selected examples see: (a) E. A. Couladouros and V. I. Moutsos, *Tetrahedron Lett.*, 1999, **40**, 7027–7030; (b) L. Ciasullo, A. Casapullo, A. Cutignano, G. Bifulco, C. Debitus, J. Hooper, L. G. Paloma and R. Riccio, *J. Nat. Prod.*, 2002, **65**, 407–410.
- 7 (a) K. C. Nicolaou, C. N. C. Boddy, S. Bräse and N. Winssinger, *Angew. Chem.*, 1999, **111**, 2230–2287 (*Angew. Chem., Int. Ed.*, 1999, **38**, 2096–2152); (b) K. C. Nicolaou and C. N. C. Boddy, *J. Am. Chem. Soc.*, 2002, **124**, 10451–10455; (c) B. M. Crowley and D. L. Boger, *J. Am. Chem. Soc.*, 2006, **128**, 2885–2892.
- 8 R. D. Kavthe, J. R. A. Kincaid and B. H. Lipshutz, *ACS Sustainable Chem. Eng.*, 2022, **10**, 16896–16902.
- 9 M. Basilaia, M. H. Chen, J. Secka and J. L. Gustafson, *Acc. Chem. Res.*, 2022, **55**, 2904–2919.
- 10 (a) D. G. Brown and H. J. Wobst, *J. Med. Chem.*, 2021, **64**, 2313–2338; (b) R. K. Sadineni, R. K. Raplolu, V. V. N. K. V. P. Raju, S. Srinivasu, S. Malladi and N. Mulakayala, *Chem. Pap.*, 2021, **75**, 1475–1483; (c) S. Guo, X. Pang, L. Peng, M. Zhan, L. Fan, Y. Gong, F. Kang and Y. Chen, *Bioorg. Med. Chem. Lett.*, 2015, **25**, 2425–2428.
- 11 (a) F. Fang, M. M. Xue, M. Ding, J. Zhang, S. Li and X. Chen, *Chem. – Asian J.*, 2021, **16**, 2489–2494; (b) R. Wang, Y. Wu, J. Wang, H. Huang, Y. Wang, S. Xu and F. Zhao, *J. Mol. Struct.*, 2022, **1257**, 132642.
- 12 For selected examples, see: (a) N. M. Levinson and S. G. Boxer, *PLoS One*, 2012, **7**, e29828; (b) P. Koelblinger, J. Dornbierer and R. Dummer, *Future Oncol.*, 2017, **13**, 1755–1766; (c) Q. Liu, Y. Sabnis, Z. Zhao, T. Zhang, S. J. Buhrlage, L. H. Jones and N. S. Gray, *Chem. Biol.*, 2013, **20**, 146–159.
- 13 J. A. McRae, R. Y. Moir, J. J. Ursprung and H. H. Gibbs, *J. Org. Chem.*, 1954, **19**, 1500–1508.
- 14 M. Dahlgard and R. Q. Brewster, *J. Am. Chem. Soc.*, 1958, **80**, 5861–5863.
- 15 K. Fuji, T. Oka, T. Kawabata and T. Kinoshita, *Tetrahedron*, 1998, **39**, 1373–1376.
- 16 M. S. Betson, J. Clayden, C. P. Worrall and S. Peace, *Angew. Chem., Int. Ed.*, 2006, **45**, 5803–5807 (*Angew. Chem.*, 2006, **118**, 5935–5939).
- 17 J. Clayden, J. Senior and M. Helliwell, *Angew. Chem., Int. Ed.*, 2009, **48**, 6270–6273 (*Angew. Chem.*, 2009, **121**, 6388–6391).

- 18 (a) D. Casarini, L. Lunazzi and G. Placucci, *J. Org. Chem.*, 1987, **52**, 4721–4726; (b) D. P. Curran, H. Qi, S. J. Geib and N. C. DeMello, *J. Am. Chem. Soc.*, 1994, **116**, 3131–3132.
- 19 (a) T. Kawabata, C. Jiang, K. Hayashi, K. Tsubaki, T. Yoshimura, S. Majumdar, T. Sasamori and N. Tokitoh, *J. Am. Chem. Soc.*, 2009, **131**, 54–55; (b) K. Hayashi, N. Matubayasi, C. Jiang, T. Yoshimura, S. Majumdar, T. Sasamori, N. Tokitoh and T. Kawabata, *J. Org. Chem.*, 2010, **75**, 5031–5036.
- 20 R. Costil, H. J. A. Dale, N. Fey, G. Whitcombe, J. V. Matlock and J. Clayden, *Angew. Chem., Int. Ed.*, 2017, **56**, 12533–12537.
- 21 B. Yuan, A. Page, C. P. Worrall, F. Escalettes, S. C. Willies, J. J. W. McDouall, N. J. Turner and J. Clayden, *Angew. Chem., Int. Ed.*, 2010, **49**, 7010–7013 (*Angew. Chem.*, 2010, **122**, 7164–7167).
- 22 L. Dai, Y. Liu, Q. Xu, M. Wang, Q. Zhu, P. Yu, G. Zhong and X. Zeng, *Angew. Chem., Int. Ed.*, 2023, **62**, e202216534 (*Angew. Chem.*, 2023, **135**, e202216534).
- 23 H. Bao, Y. Chen and X. Yang, *Angew. Chem., Int. Ed.*, 2023, **62**, e202300481.
- 24 S. Shee, S. S. Ranganathappa, M. S. Gadhave, R. Gogoi and A. T. Biju, *Angew. Chem., Int. Ed.*, 2023, **62**, e202311709.
- 25 For reports on cooperative desymmetrization/kinetic resolution, see: (a) Y. Wu, M. Li, J. Sun, G. Zheng and Q. Zhang, *Angew. Chem., Int. Ed.*, 2022, **61**, e202117340; (b) M. Zhu, H. J. Jiang, I. Sharanov, E. Irran and M. Oestreich, *Angew. Chem., Int. Ed.*, 2023, **62**, e202304475.
- 26 B. A. Zhou, X. N. Li, C. L. Zhang, Z. X. Wang and S. Ye, *Angew. Chem., Int. Ed.*, 2024, **63**, e20231422.
- 27 Y. Liu, L. Yuan, L. Dai, Q. Zhu, G. Zhong and X. Zeng, *ChemRxiv*, 2023, preprint, DOI: [10.26434/chemrxiv-2023-78vb3](https://doi.org/10.26434/chemrxiv-2023-78vb3).
- 28 L. Li, W. Ti, T. Miao, J. Ma, A. Lin, Q. Chu and S. Gao, *J. Org. Chem.*, 2024, **89**, 4067–4073.
- 29 X. Han, L. Chan, Y. Yan, Y. Zhao, A. Lin, S. Gao and H. Yao, *ACS Catal.*, 2024, **14**, 3457–3481.
- 30 A. N. Dinh, R. R. Noorbehesht, S. T. Toenjes, A. C. Jackson, M. A. Saputra, S. M. Maddox and J. L. Gustafson, *Synlett*, 2018, 2155–2160.
- 31 S. D. Vaidya, S. T. Toenjes, N. Yamamoto, S. M. Maddox and J. L. Gustafson, *J. Am. Chem. Soc.*, 2020, **142**, 2198–2203.
- 32 During the preparation of this article, a few related works were published in the literature: (a) Y. Wu, X. Guan, H. Zhao, M. Li, T. Liang, J. Sun, G. Zheng and Q. Zhang, *Chem. Sci.*, 2024, **15**, 4564–4570; (b) L. Dai, X. Zhou, J. Guo, Q. Huang and Y. Lu, *Chem. Sci.*, 2024, **15**, 5993–6001; (c) Y. Wang, R. Mi, S. Yu and X. Li, *ACS Catal.*, 2024, **14**, 4638–4647; (d) J. Xu, W. Lin, H. Zheng and X. Li, *ACS Catal.*, 2024, **14**, 6667–6673; (e) Z. Ye, W. Xie, D. Wang, H. Liu and X. Yang, *ACS Catal.*, 2024, **14**, 4958–4967.