

Volume 10 | Number 14 | 21 July 2023

**10**  
YEARS  
ANNIVERSARY



# ORGANIC CHEMISTRY

## FRONTIERS



CHINESE  
CHEMICAL  
SOCIETY



ROYAL SOCIETY  
OF CHEMISTRY

[rsc.li/frontiers-organic](https://rsc.li/frontiers-organic)

## RESEARCH ARTICLE

View Article Online  
View Journal | View IssueCite this: *Org. Chem. Front.*, 2023, **10**, 3460Received 1st March 2023,  
Accepted 17th April 2023

DOI: 10.1039/d3qo00322a

rsc.li/frontiers-organic

## Synthesis of atropisomeric phosphino-triazoles and their corresponding gold(I) complexes†

Yiming Zhao,<sup>a</sup> Fernanda Meloni,<sup>a</sup> Louise Male,<sup>a</sup> Cécile S. Le Duff,<sup>a</sup> William D. G. Brittain,<sup>b</sup> Benjamin R. Buckley<sup>b</sup> and John S. Fossey<sup>b,†,‡</sup>

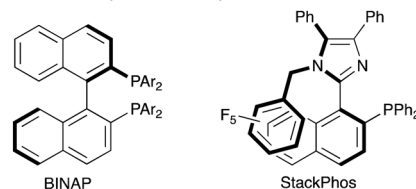
The synthesis of atropisomeric phosphino-triazoles is disclosed. It was found that the introduction of a phosphine functionality onto the 5-position of a 1,2,3-triazole ring could be highly restrictive towards the rotation around a triazole-aryl bond. VT NMR and chiral HPLC studies demonstrated that rotation was restricted even at high temperatures. Gold(I) chloride complexes of single-enantiomer phosphines were prepared and again demonstrated to be conformationally stable.

Atropisomerism is one of the fundamental families of chirality, where rotational restriction around a single bond in a molecule creates two non-superimposable mirror images.<sup>1–3</sup> Atropisomerism is present in many compounds with several natural products, ligands and organocatalysts incorporating restricted rotation.<sup>4–6</sup> Phosphine ligands for transition metals that rely on atropisomerism to engender chirality into products are well established, with compounds such as BINAP and the phosphoramidite family commonly being employed in a variety of asymmetric catalytic applications (Fig. 1a).<sup>7–9</sup> In general heterocyclic-containing phosphines have seen a wide variety of application and new scaffolds in this space have great value.<sup>10</sup>

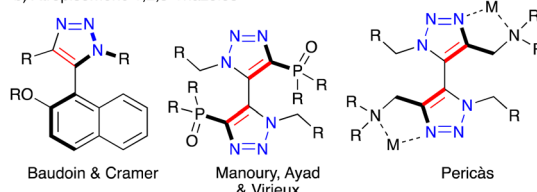
Whilst atropisomeric metal ligands are common, most are built around a 1,1'-bi-2-naphthyl scaffold.<sup>11–14</sup> Ligands built around the 1,2,3-triazole core are in comparison extremely sparse. Reports in this area are usually centred around a 5–5' bistriazole architecture<sup>15–17</sup> apart from a small number of notable exceptions (Fig. 1b).<sup>18,19</sup> In extension atropisomeric 1,2,3-triazole-phosphines to the best of our knowledge have not been previously reported and there appears to be only limited reports of atropisomeric 1,2,4-triazole-phosphines.<sup>20</sup> This is particularly intriguing as accessing 1,2,3-triazoles can be readily achieved through the ubiquitous copper-catalysed

azide-alkyne cycloaddition (CuAAC) which is part of the family of commonly employed “click” reactions.<sup>14,21–23</sup> Triazoles have been widely employed in biorthogonal conjugation, which in

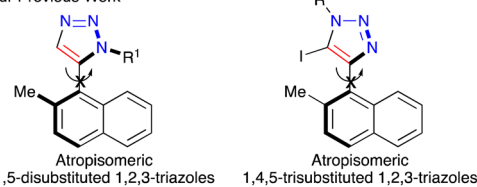
## a) Well-Established Atropisomeric Phosphines



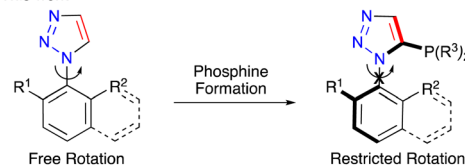
## b) Atropisomeric 1,2,3-Triazoles



## c) Our Previous Work



## d) This work



**Fig. 1** (a) Atropisomeric phosphines commonly employed in asymmetric catalysis (b) atropisomeric 1,2,3-triazoles (c) our previous work on the synthesis of atropisomeric 1,2,3-triazoles (d) this work on the synthesis of atropisomeric 1,2,3-triazole phosphines.

<sup>a</sup>School of Chemistry, University of Birmingham, Edgbaston, Birmingham, West Midlands, B15 2TT, UK. E-mail: J.S.Fossey@bham.ac.uk

<sup>b</sup>Department of Chemistry, Durham University, South Road, Durham DH1 3LE, UK. E-mail: william.d.brittain@durham.ac.uk

<sup>c</sup>Department of Chemistry, Loughborough University, Loughborough, Leicestershire, LE11 3TU, UK. E-mail: B.R.Buckley@lboro.ac.uk

† Electronic supplementary information (ESI) available. CCDC 2245022–2245029 and 2245473. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3qo00322a>

‡ In memory of John S. Fossey, a superb scientist, mentor and friend who will be sadly missed.

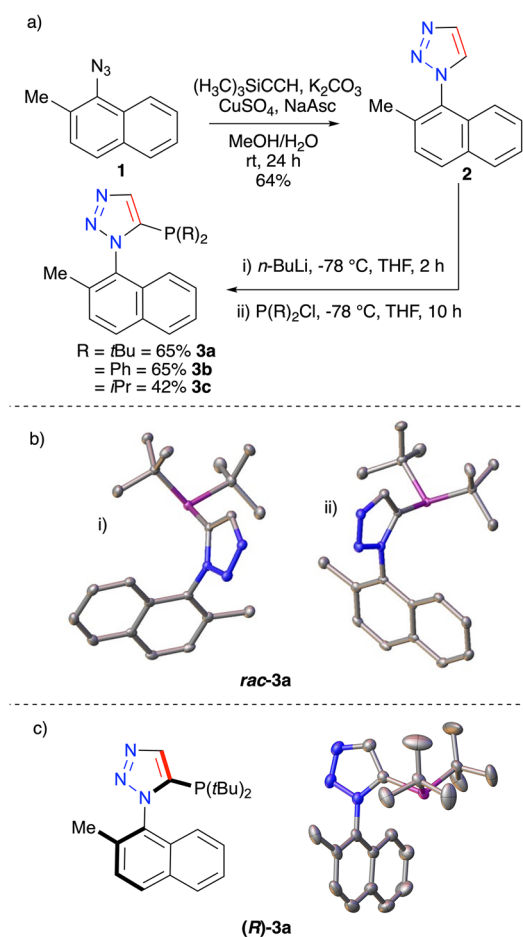


part led to the awarding of the 2022 Nobel Prize in Chemistry to Meldal, Sharpless and Bertozzi for their pioneering work in the field.<sup>24–26</sup> The CuAAC is well known as a reaction that can be carried out under mild conditions with exquisite regioselectivity and up to quantitative yields.<sup>27–29</sup>

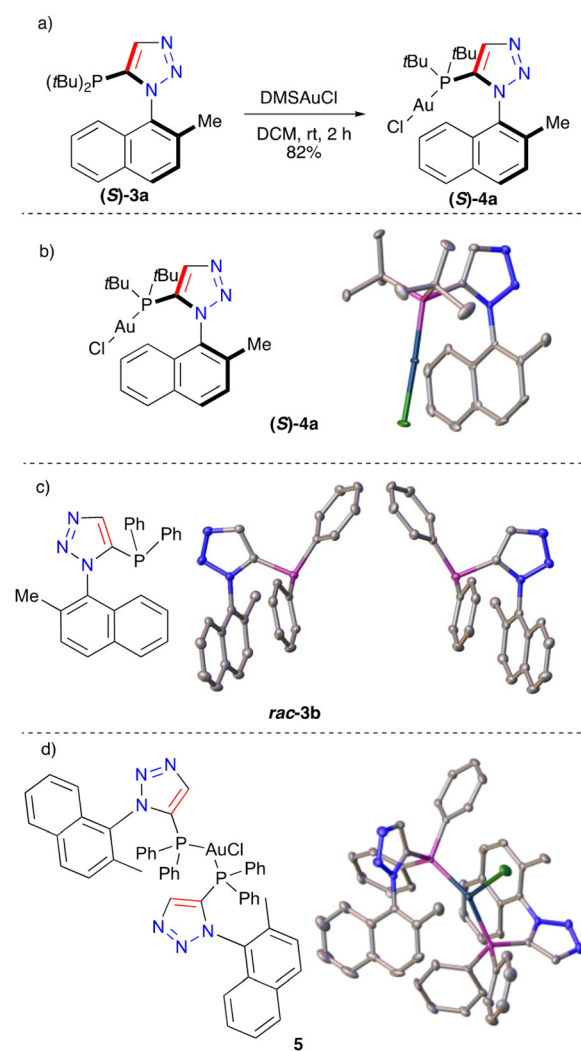
We have an ongoing interest into the utility of the CuAAC reaction and how it can be employed to generate chiral molecules. Previously we have reported on the generation of point chiral and axially chiral triazoles as well as utilising triazoles in a range of linkage applications.<sup>30–34</sup> Recently we have reported on the synthesis of a range of phosphine-appended triazoles<sup>35,36</sup> and on the establishment of how bi-aryl 1,2,3-triazoles can be rendered atropisomeric (Fig. 1c).<sup>37</sup> Therefore we wished to expand on these concepts and investigate if we could readily access atropisomeric and point chiral triazole-containing phosphines utilising the CuAAC reaction and in turn generate their corresponding gold complexes (Fig. 1d).

To begin, we took inspiration from our previous studies on the 2-methyl-1-naphthyl moiety which can be incorporated

into atropisomeric triazoles which are stable up to high temperatures.<sup>37</sup> The triazole **2** was successfully synthesised in two steps from the commercially available 2-methyl-1-naphthylamine. Phosphine incorporation was then conducted using a lithiation followed by trapping with a range of chlorophosphines. This protocol led to the successful isolation of compounds **3a–c** incorporating <sup>t</sup>Bu, Ph and <sup>i</sup>Pr groups around the phosphorus in 65%, 65% and 42% yields, respectively (Fig. 2a). Immediately it was noticed in the <sup>1</sup>H NMR spectrum of **3a** that the resonances corresponding to the <sup>t</sup>Bu protons were diastereotopic. This was encouraging as it suggested the bi-aryl bond was no longer freely rotating. To confirm the



**Fig. 2** (a) Synthesis of 1,2,3-triazole phosphines (b) molecules from the crystal structure of *rac*-**3a** with ellipsoids drawn at the 50% probability level. Symmetry codes used to generate the equivalent atoms: (i)  $x, y, z$ , (ii)  $1 - x, 1 - y, 1 - z$ . Hydrogen atoms omitted for clarity (c) single-crystal X-ray diffraction structure of (*R*)-**3a**. Ellipsoids are drawn at the 50% probability level. Hydrogen atoms omitted for clarity.



**Fig. 3** (a) Synthesis of a single enantiomer gold(i) chloride complex (b) single-crystal X-ray diffraction structure of (*S*)-**4a**. Ellipsoids are drawn at the 50% probability level. A molecule of DCM has been omitted for clarity. Hydrogen atoms omitted for clarity. (c) Molecules from the crystal structure of *rac*-**3b** with ellipsoids drawn at the 50% probability level. Symmetry codes used to generate the equivalent atoms: (i)  $x, y, z$ , (ii)  $1 - x, 1 - y, 1 - z$ . Hydrogen atoms omitted for clarity. (d) Single-crystal X-ray diffraction structure of homo-chiral complex **5**. Ellipsoids are drawn at the 50% probability level. A molecule of DCM has been omitted for clarity. Hydrogen atoms have been omitted for clarity.



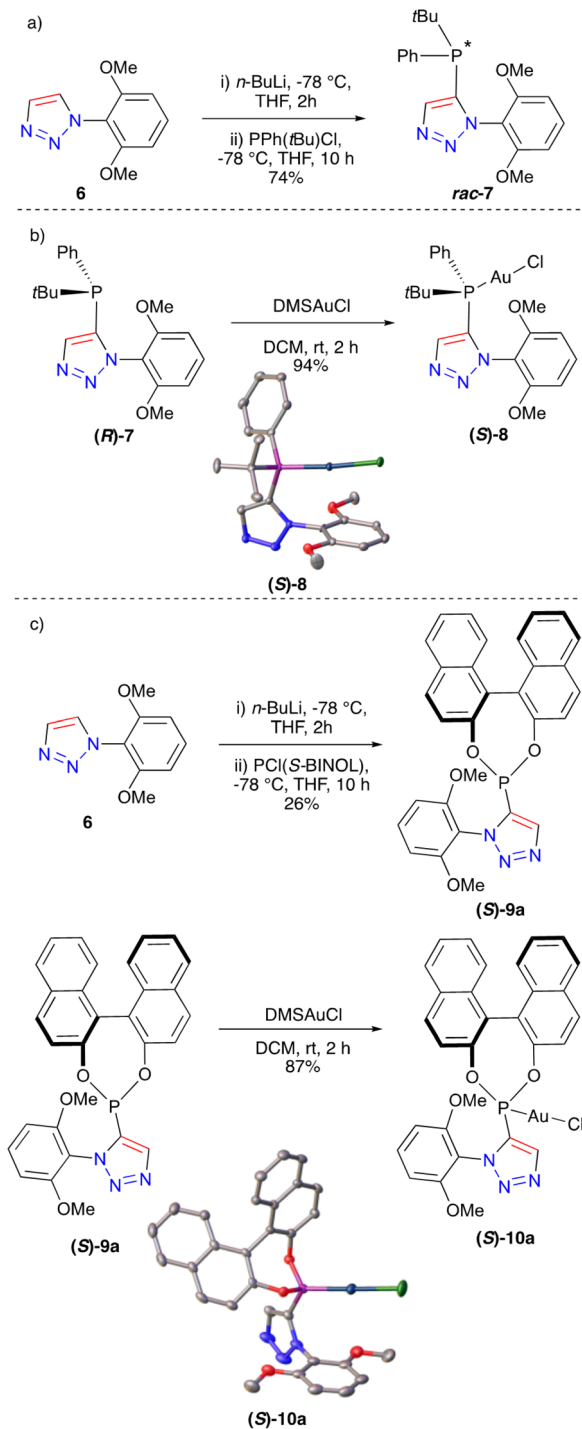
stability of the observed atropisomers, a VT NMR study was undertaken (see ESI†). In toluene-d<sub>8</sub> at 105 °C no coalescence of resonances was observed, pointing to a highly rotationally

restricted compound. In addition, a single-crystal X-ray structure was determined with both enantiomers of **3a** being clearly visible in the unit cell (Fig. 2b).

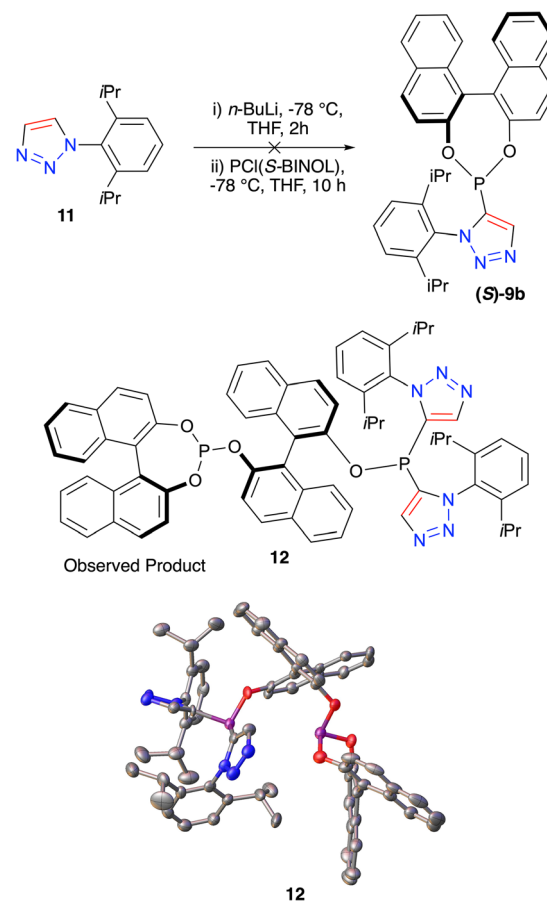
Compound **3b**, in which two phenyl groups are attached to the phosphorus was also successfully crystallised and again two enantiomers were visible within the unit cell (Fig. 3c). The isopropyl-substituted compound **3c** was observed to display characteristics of atropisomerism with diastereotopic protons being clearly visible in the <sup>1</sup>H NMR spectrum. In all, this series of compounds was highly encouraging as all substitutions around the phosphorous appeared to significantly restrict the rotational freedom of the molecules.

The next step was to obtain single-enantiomer compounds. Initially a Staudinger reaction was trialed to resolve the two enantiomers. Unfortunately, this was unsuccessful and thus led us to pursue chiral-prep HPLC to separate and isolate the single-enantiomer triazoles. Separations for both compounds (**R**)-**3a** and (**S**)-**3a** were successful (see ESI†) and the single enantiomers of **3a** were obtained in sufficient quantities for characterisation and metal complexation.

X-ray crystallographic studies of (**R**)-**3a** were conducted and the conformation around the restricted bi-aryl bond was deter-



**Fig. 4** (a) Synthesis of 1,2,3-triazolephosphine **rac-7** (b) synthesis and single-crystal X-ray diffraction structure of gold(i) chloride complex (**S**)-**8**. Ellipsoids are drawn at the 50% probability level (c) Synthesis of 1,2,3-triazole phosphonite (**S**)-**9a** and subsequent complexation. Single-crystal X-ray diffraction structure of (**S**)-**10a**. Ellipsoids are drawn at the 50% probability level.



**Fig. 5** Attempted synthesis of phosphonite (**S**)-**9b** and subsequent acquisition of the single-crystal X-ray diffraction structure **12**. Ellipsoids are drawn at the 50% probability level. Hydrogen atoms are omitted for clarity.



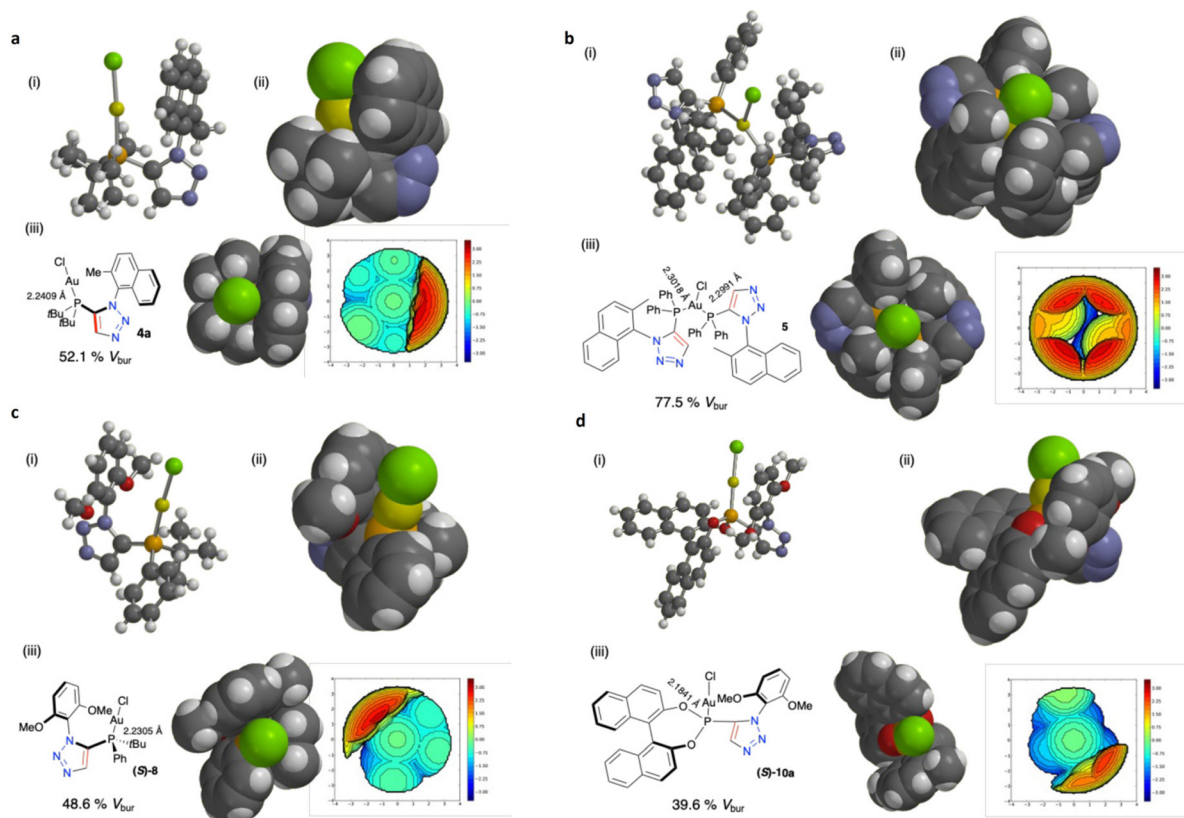
mined (Fig. 2c). This pure material (**S**)-**3a** was complexed with DMSAuCl, leading to the formation of (**S**)-**4a** in 82% and a corresponding crystal structure obtained (Fig. 3a and b). This structure demonstrated no racemisation had occurred during complexation and that a single-enantiomer complex had been successfully prepared. The phenyl substituted compound **3b** in its racemic form was trialled in a gold complexation and the homo-chiral complex **5** was obtained (Fig. 3c and d). We also attempted to use chiral-prep HPLC to obtain a single-enantiomer sample of **3c**, however after many attempts baseline resolution was not achieved. This series of compounds clearly demonstrated that the CuAAC reaction is a powerful tool to access atropisomerically stable phosphines and gold complexes.

Next, we wished to explore the ability to generate *P*-chirogenic triazole phosphines. This class of ligand was most famously used by Knowles in his seminal synthesis of L-DOPA using asymmetric hydrogenation.<sup>38,39</sup> However, *P*-chirogenic phosphines have been underutilised mainly due to their synthetic routes relying on phosphine oxide intermediates. We believed that through trapping a lithiated triazole with a chlorophosphine with two different substituents, an atropisomeric *P*-chirogenic ligand could be prepared. We have previously disclosed the preparation of triazole **6** which was utilised to prepare a range of bulky triazole phosphines. Taking **6** and treating it with *n*-BuLi followed by PPh(<sup>t</sup>Bu)Cl gave the triazole

*rac*-**7** in 74% yield (Fig. 4a). The <sup>1</sup>H NMR spectrum of this compound clearly showed diastereotopic protons with the OMe groups giving two clearly defined singlets of integration 3. Preparative chiral HPLC was then employed to resolve the enantiomers of **7**. Both enantiomers were successfully isolated (see ESI†) and the corresponding gold chloride complex (**S**)-**8** was prepared in 94% yield. Single-crystal X-ray crystallography was then used to unambiguously determine the chirality of the gold complex (**S**)-**8** (Fig. 4b). In addition a racemic crystal structure of **8** was also determined (see ESI†).

Finally, we wished to expand our study to investigate the synthesis of a new class of ligand incorporating the phosphonite functional group in combination with a triazole. It was decided that quenching the lithiated triazole **6** with PCl((*S*)-BINOL) would give a highly unusual species which incorporated both BINOL and 1,2,3-triazole units. Using this approach phosphonite triazole (**S**)-**9a** was successfully prepared in 26% yield (Fig. 4c). Taking this triazole and treating it with DMSAuCl furnished the gold complex (**S**)-**10a** with the stereochemical configuration confirmed through single-crystal X-ray analysis (Fig. 4c).

Interestingly whilst conducting the same reaction sequence with the *i*Pr substituted triazole **11**, there was no observed formation of the desired phosphonite (**S**)-**9b**; instead, a solid by-product was formed. Single-crystal X-ray analysis led to the



**Fig. 6** Single-crystal X-ray diffraction structures of (a) **4a**, (b) **5**, (c) **8** and (d) **10**: (i) Ortep representation, ellipsoid probability 50% (rendered in Spartan). (ii) Space-filling representation. (iii) Percentage buried volume determined from the crystal structures steric map of ligand depicted (right).



identification of the multi-aryl species **12** (Fig. 5). Whilst this was not the desired outcome this structure was highly unusual both with multiple phosphorus atoms and multiple stereodefined bi-aryls. This compound could therefore be of use in applications where highly sterically hindered phosphorus ligands are required.

We and others have previously discussed the importance of various parameters including steric effects of bulky phosphines relating to suitability and efficacy in catalysis (primarily as ligands for metals in metal-mediated catalysis).<sup>35,36</sup> The use of the Tolman cone angle has been an effective descriptor of ligand bulkiness for many years,<sup>40</sup> more recently Nolan's percentage buried volume parameter ( $\%V_{\text{bur}}$ ) has been widely applied to various ligand types (eg NHC's) to describe and correlate ligand/catalyst activity to the environment surrounding the metal catalyst. The use of  $\%V_{\text{bur}}$  has undoubtedly been accelerated through the use of the simple free web tool SambVca developed by Cavallo and co-workers.<sup>41</sup> So we set about determining some steric parameters of our novel Au phosphines using this tool.<sup>42-44</sup> Utilising the X-ray crystal structure CIF file data obtained for complexes **4a**, **5**, **8** and **10**, we were able to gain the  $\%V_{\text{bur}}$  data utilising the Au-P bond lengths observed (Fig. 6). Interestingly, the dimer complex **5** has a very significant buried volume surrounding the Au atom (77.5%  $V_{\text{bur}}$ ).

## Conclusions

Overall, we have been able to demonstrate that the formation of triazole-phosphines is a viable option for the generation of stable atropisomeric species. This is the first time to the best of our knowledge that a post CuAAC modification to a triazole ring has been utilised to engender atropisomerism. This approach offers added flexibility in being able to readily change the other groups around the phosphorous atom. We were also able to successfully access new classes of *P*-chirogenic phosphines and phosphonites. Preparative chiral HPLC was used to access enantiomerically pure samples of all of these species. These ligand architectures in general were readily amenable to the formation of gold(i) chloride complexes. We believe that the strategies outlined in chiral ligand design here may have an impact in catalysis and in particular gold- and palladium-mediated reactions.

## Author contributions

All authors contributed in varying degrees to planning the experiments, evaluating results and writing of the manuscript; specific contributions in addition to this are listed below: J. S. F. led and co-conceived the overarching project, providing critical assessment of data, directed most aspects, and supervised the experimental work; B. R. B. helped direct aspects of the research and gave input and critical assessment throughout the progress of the project; Y. Z., F. M., L. M. and W. D. G.

B co-conceived aspects of the project, conducted the synthesis and drafted a proportion of the ESI,<sup>†</sup> offered critical suggestions and conducted the XRD data collection and analysis herein, along with L. M. C. S. L. D. conducted and advised on the VT NMR experiments. W. D. G. B. and B. R. B. wrote the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors would like to thank the University of Birmingham, Durham University and Loughborough University for support. The CASE consortium is thanked for networking opportunities.<sup>45,46</sup>

## References

- 1 J. K. Cheng, S.-H. Xiang, S. Li, L. Ye and B. Tan, Recent Advances in Catalytic Asymmetric Construction of Atropisomers, *Chem. Rev.*, 2021, **121**, 4805–4902.
- 2 M. Basilaia, M. H. Chen, J. Secka and J. L. Gustafson, Atropisomerism in the Pharmaceutically Relevant Realm, *Acc. Chem. Res.*, 2022, **55**, 2904–2919.
- 3 G.-J. Mei, W. L. Koay, C.-Y. Guan and Y. Lu, Atropisomers beyond the C–C axial chirality: Advances in catalytic asymmetric synthesis, *Chem*, 2022, **8**, 1855–1893.
- 4 J. E. Smyth, N. M. Butler and P. A. Keller, A twist of nature – the significance of atropisomers in biological systems, *Nat. Prod. Rep.*, 2015, **32**, 1562–1583.
- 5 Z. Xu and L.-W. Xu, Development of Ar-BINMOL-Derived Atropisomeric Ligands with Matched Axial and sp<sup>3</sup> Central Chirality for Catalytic Asymmetric Transformations, *Chem. Rec.*, 2015, **15**, 925–948.
- 6 S. Brandes, B. Niess, M. Bella, A. Prieto, J. Overgaard and K. A. Jørgensen, Non-Biaryl Atropisomers in Organocatalysis, *Chem. – Eur. J.*, 2006, **12**, 6039–6052.
- 7 M. Berthod, G. Mignani, G. Woodward and M. Lemaire, Modified BINAP: The How and the Why, *Chem. Rev.*, 2005, **105**, 1801–1836.
- 8 T. H. Wöste and M. Oestreich, BINAP versus BINAP(O) in Asymmetric Intermolecular Mizoroki–Heck Reactions: Substantial Effects on Selectivities, *Chem. – Eur. J.*, 2011, **17**, 11914–11918.
- 9 J. F. Teichert and B. L. Feringa, Phosphoramidites: Privileged Ligands in Asymmetric Catalysis, *Angew. Chem., Int. Ed.*, 2010, **49**, 2486–2528.
- 10 M. H. Tse, P. Y. Choy and F. Y. Kwong, Facile Assembly of Modular-Type Phosphines for Tackling Modern Arylation Processes, *Acc. Chem. Res.*, 2022, **55**, 3688–3705.



- 11 M. P. Walsh, J. M. Phelps, M. E. Lennon, D. S. Yufit and M. O. Kitching, Enantioselective synthesis of ammonium cations, *Nature*, 2021, **597**, 70–76.
- 12 D. Parmar, E. Sugiono, S. Raja and M. Rueping, Complete Field Guide to Asymmetric BINOL-Phosphate Derived Brønsted Acid and Metal Catalysis: History and Classification by Mode of Activation; Brønsted Acidity, Hydrogen Bonding, Ion Pairing, and Metal Phosphates, *Chem. Rev.*, 2014, **114**, 9047–9153.
- 13 J. M. Brunel, BINOL: A Versatile Chiral Reagent, *Chem. Rev.*, 2005, **105**, 857–898.
- 14 F. Yu, Y. Chen, H. Jiang and X. Wang, Recent advances of BINOL-based sensors for enantioselective fluorescence recognition, *Analyst*, 2020, **145**, 6769–6812.
- 15 P. Etayo, E. C. Escudero-Adán and M. A. Pericàs, 5,5'-Bistriazoles as axially chiral, multidentate ligands: synthesis, configurational stability and catalytic application of their scandium(III) complexes, *Catal. Sci. Technol.*, 2017, **7**, 4830–4841.
- 16 N. Sevrain, J.-N. Volle, J.-L. Pirat, T. Ayad and D. Virieux, 1,1'-Dibenzyl-bis-(triazolyl)diphenylphosphine dioxide: a new efficient organocatalyst for silicon tetrachloride-mediated enantioselective Abramov-type phosphorylation of aldehydes with trialkyl phosphites, *RSC Adv.*, 2017, **7**, 52101–52104.
- 17 C. Laborde, M.-M. Wei, A. van der Lee, E. Deydier, J.-C. Daran, J.-N. Volle, R. Poli, J.-L. Pirat, E. Manoury and D. Virieux, Double [3+2]-dimerisation cascade synthesis of bis(triazolyl)bisphosphanes, a new scaffold for bidentate bisphosphanes, *Dalton Trans.*, 2015, **44**, 12539–12545.
- 18 Q.-H. Nguyen, S.-M. Guo, T. Royal, O. Baudoin and N. Cramer, Intermolecular Palladium(0)-Catalyzed Atropo-enantioselective C–H Arylation of Heteroarenes, *J. Am. Chem. Soc.*, 2020, **142**, 2161–2167.
- 19 R. Vroemans, S. R. Ribone, J. Thomas, L. Van Meervelt, T. Ollevier and W. Dehaen, Synthesis of homochiral sulfanyl- and sulfoxide-substituted naphthyltriazoles and study of the conformational stability, *Org. Biomol. Chem.*, 2021, **19**, 6521–6526.
- 20 A. A. Kirilchuk, A. A. Yurchenko, Y. G. Vlasenko, A. N. Kostyuk and A. B. Rozhenko, Synthesis and Structure of Phosphanylated Bis-Triazoles as Potential Ligands for Chiral Catalysts, *Chem. Heterocycl. Compd.*, 2015, **50**, 1559–1566.
- 21 N. K. Devaraj and M. G. Finn, Introduction: Click Chemistry, *Chem. Rev.*, 2021, **121**, 6697–6698.
- 22 H. C. Kolb, M. G. Finn and K. B. Sharpless, Click Chemistry: Diverse Chemical Function from a Few Good Reactions, *Angew. Chem., Int. Ed.*, 2001, **40**, 2004–2021.
- 23 S. L. Scinto, D. A. Bilodeau, R. Hincapie, W. Lee, S. S. Nguyen, M. Xu, C. W. am Ende, M. G. Finn, K. Lang, Q. Lin, J. P. Pezacki, J. A. Prescher, M. S. Robillard and J. M. Fox, Bioorthogonal chemistry, *Nat. Rev. Methods Primers*, 2021, **1**, 30.
- 24 V. V. Rostovtsev, L. G. Green, V. V. Fokin and K. B. Sharpless, A Stepwise Huisgen Cycloaddition Process: Copper(I)-Catalyzed Regioselective “Ligation” of Azides and Terminal Alkynes, *Angew. Chem., Int. Ed.*, 2002, **41**, 2596–2599.
- 25 C. W. Tornøe, C. Christensen and M. Meldal, Peptidotriazoles on Solid Phase: [1,2,3]-Triazoles by Regiospecific Copper(I)-Catalyzed 1,3-Dipolar Cycloadditions of Terminal Alkynes to Azides, *J. Org. Chem.*, 2002, **67**, 3057–3064.
- 26 E. M. Sletten and C. R. Bertozzi, Bioorthogonal Chemistry: Fishing for Selectivity in a Sea of Functionality, *Angew. Chem., Int. Ed.*, 2009, **48**, 6974–6998.
- 27 J. E. Hein and V. V. Fokin, Copper-catalyzed azide–alkyne cycloaddition (CuAAC) and beyond: new reactivity of copper(I) acetylides, *Chem. Soc. Rev.*, 2010, **39**, 1302–1315.
- 28 M. Meldal and C. W. Tornøe, Cu-Catalyzed Azide–Alkyne Cycloaddition, *Chem. Rev.*, 2008, **108**, 2952–3015.
- 29 V. Hong, S. I. Presolski, C. Ma and M. G. Finn, Analysis and Optimization of Copper-Catalyzed Azide–Alkyne Cycloaddition for Bioconjugation, *Angew. Chem., Int. Ed.*, 2009, **48**, 9879–9883.
- 30 W. D. G. Brittain, B. R. Buckley and J. S. Fossey, Kinetic resolution of alkyne-substituted quaternary oxindoles via copper catalysed azide–alkyne cycloadditions, *Chem. Commun.*, 2015, **51**, 17217–17220.
- 31 W. D. G. Brittain, B. M. Chapin, W. Zhai, V. M. Lynch, B. R. Buckley, E. V. Anslyn and J. S. Fossey, The Bull–James assembly as a chiral auxiliary and shift reagent in kinetic resolution of alkyne amines by the CuAAC reaction, *Org. Biomol. Chem.*, 2016, **14**, 10778–10782.
- 32 W. D. G. Brittain, A. G. Dalling, Z. Sun, C. S. L. Duff, L. Male, B. R. Buckley and J. S. Fossey, Coetaneous catalytic kinetic resolution of alkynes and azides through asymmetric triazole formation, *Sci. Rep.*, 2019, **9**, 15086.
- 33 W. Zhai, L. Male and J. S. Fossey, Glucose selective bis-boronic acid click-fluor, *Chem. Commun.*, 2017, **53**, 2218–2221.
- 34 W. Zhai, B. M. Chapin, A. Yoshizawa, H.-C. Wang, S. A. Hodge, T. D. James, E. V. Anslyn and J. S. Fossey, “Click-fluors”: triazole-linked saccharide sensors, *Org. Chem. Front.*, 2016, **3**, 918–928.
- 35 Y. Zhao, M. G. Wakeling, F. Meloni, T. J. Sum, H. van Nguyen, B. R. Buckley, P. W. Davies and J. S. Fossey, Balancing Bulkiness in Gold(I) Phosphino-triazole Catalysis, *Eur. J. Org. Chem.*, 2019, 5540–5548.
- 36 Y. Zhao, H. van Nguyen, L. Male, P. Craven, B. R. Buckley and J. S. Fossey, Phosphino-Triazole Ligands for Palladium-Catalyzed Cross-Coupling, *Organometallics*, 2018, **37**, 4224–4241.
- 37 F. Meloni, W. D. G. Brittain, L. Male, C. S. Le Duff, B. R. Buckley, A. G. Leach and J. S. Fossey, Enantiomer stability of atropisomeric 1,5-disubstituted 1,2,3-triazoles, *Tetrahedron Chem*, 2022, **1**, 100004.
- 38 W. S. Knowles, Application of organometallic catalysis to the commercial production of L-DOPA, *J. Chem. Educ.*, 1986, **63**, 222.



- 39 W. S. Knowles and R. Noyori, Pioneering Perspectives on Asymmetric Hydrogenation, *Acc. Chem. Res.*, 2007, **40**, 1238–1239.
- 40 C. A. Tolman, Steric effects of phosphorus ligands in organometallic chemistry and homogeneous catalysis, *Chem. Rev.*, 1977, **77**, 313–348.
- 41 SambVca 2.1 web application, <https://www.molnac.unisa.it/OMtools/sambvca2.1/index.html> (accessed February 2023).
- 42 A. Poater, B. Cosenza, A. Correa, S. Giudice, F. Ragone, V. Scarano and L. Cavallo, SambVca: A Web Application for the Calculation of the Buried Volume of N-Heterocyclic Carbene Ligands, *Eur. J. Inorg. Chem.*, 2009, 1759–1766.
- 43 L. Falivene, R. Credendino, A. Poater, A. Petta, L. Serra, R. Oliva, V. Scarano and L. Cavallo, SambVca 2. A Web Tool for Analyzing Catalytic Pockets with Topographic Steric Maps, *Organometallics*, 2016, **35**, 2286–2293.
- 44 L. Falivene, Z. Cao, A. Petta, L. Serra, A. Poater, R. Oliva, V. Scarano and L. Cavallo, Towards the online computer-aided design of catalytic pockets, *Nat. Chem.*, 2019, **11**, 872–879.
- 45 D. T. Payne, J. S. Fossey and R. B. P. Elmes, Catalysis and Sensing for our Environment (CASE2015) and the Supramolecular Chemistry Ireland Meeting (SCI 2015): Dublin and Maynooth, Ireland. 8th–11th July, *Supramol. Chem.*, 2016, **28**, 921–931.
- 46 J. S. Fossey and W. D. G. Brittain, The CASE 2014 symposium: Catalysis and sensing for our environment, Xiamen 7th–9th November 2014, *Org. Chem. Front.*, 2015, **2**, 101–105.

