

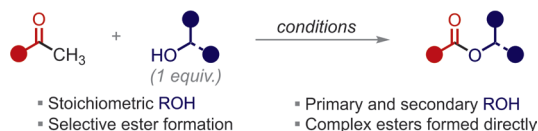
# Synform

People, Trends and Views in Chemical Synthesis

2024/09

## A Stoichiometric Haloform Coupling for Ester Synthesis with Secondary Alcohols

Highlighted article by A. C. Rowett, S. G. Sweeting, D. M. Heard, A. J. J. Lennox



### Contact

Your opinion about Synform is welcome,  
please correspond if you like:  
[marketing@thieme-chemistry.com](mailto:marketing@thieme-chemistry.com)

## Dear Readers,

In this September issue of Synform we welcome a new Editorial Board Member: Prof. Dr. Tanja Gulder (Saarland University, Germany) who joined the Editorial Board of SYNLETT with effect of July 2024. Tanja has been interviewed by Synform about her scientific interests, her feelings about becoming a Thieme Chemistry Editorial Board Member, as well as her extra-lab interests. A warm welcome to Tanja!!

The first Literature Coverage article focuses on a novel version of an evergreen classic of organic chemistry: the haloform reaction, the scope of which has been expanded by the group of A. J. J. Lennox (UK) to include a stoichiometric coupling-type reaction with secondary alcohols to give directly the corresponding esters. The next article is a Young Career Focus interview with one of the most recent Thieme Chemistry Journals Awardees, B. Schmidt (Germany), who talks with us about his scientific and extra-lab interests. The issue is wrapped up by another Literature Coverage article, highlighting a novel enantioselective synthesis of  $\alpha$ -tertiary ethynylamines and related tandem sequence to provide also azacycles, as reported in *Nat. Chem.* by the group of J. Zhou (P. R. of China).

Enjoy your reading!



### In this issue

— Editorial Board Focus

**Editorial Board Focus: Prof. Dr. Tanja Gulder (Saarland University, Germany)** ..... A136

— Literature Coverage

**A Stoichiometric Haloform Coupling for Ester Synthesis with Secondary Alcohols** ..... A138

— Young Career Focus

**Young Career Focus: Dr. Bernd M. Schmidt (Heinrich Heine Universität, Germany)** ..... A141

— Literature Coverage

**Enantioselective Propargylic Amination and Related Tandem Sequences to  $\alpha$ -Tertiary Ethynylamines and Azacycles** ..... A143

Coming soon ..... A147

### Contact

If you have any questions or wish to send feedback, please write to Matteo Zanda at: [synform@outlook.com](mailto:synform@outlook.com)

## Editorial Board Focus: Prof. Dr. Tanja Gulder (Saarland University, Germany)

**Background and Purpose.** From time to time, SYNFORM portraits Thieme Chemistry Editorial Board or Editorial Advisory Board members who answer several questions regarding their research interests and revealing their impressions and views on the developments in organic chemistry as a general research field. This Editorial Board Focus presents Prof. Dr. Tanja Gulder (Saarland University, Germany) who joined the Editorial Board of SYNLETT with effect of July 2024.

### Biographical Sketch



© Astrid Eckert, TUM

Prof. Dr. T. Gulder

**Tanja Gulder** is the Chair of Organic Chemistry at Saarland University, Germany and is heading the Synthesis of Natural-Product Derived Drugs group at the Helmholtz Center of Pharmaceutical Research Saarland (HIPS). Her laboratory is dedicated to biomimetic catalysis (enzyme mimicking), focusing on halogenations and deuterations and their application in synthesizing natural products, anti-

infectives, and diagnostics. Tanja Gulder studied chemistry at the University of Wuerzburg, Germany, where she received her diploma in 2004. After earning her Ph.D. with distinction under the supervision of Prof. G. Bringmann in 2008, she pursued postdoctoral studies with Prof. P. S. Baran at The Scripps Research Institute (La Jolla, CA, USA). After returning to Germany, she started her independent career in 2011, supported by a Liebig fellowship of the Fonds der Chemischen Industrie at RWTH Aachen. 2014 she moved to TU Muenchen, Germany, and was appointed Heisenberg-Professor of Biomimetic Catalysis in 2018. From 2020 to 2023, she was a Full Professor in Organic Chemistry at Leipzig University, Germany.

She received the Emmy-Noether grant, the prestigious support for exceptionally qualified early career scientists by the German Research Foundation. In 2023, she was elected as an ordinary member of the Saxonian Academy of Science. She was awarded the Publication Award Fluorine Chemistry Prize of the German Chemical Society for her significant contributions to developing transition-metal catalysis for transforming organic molecules to prepare value-added materials by site-selective functionalization, with impact on the agrochemical and pharmaceutical industries. In 2024, she joins SYNLETT as Associate Editor.

### INTERVIEW

**SYNFORM** *What fascinates you most about organic chemistry and synthesis?*

**Prof. Dr. T. Gulder** The creativity and complexity of building structurally complex molecules from simple substances is not just a scientific pursuit for me but a passion. The ability to design and construct molecules with specific, often complex structures with a particular function is like solving a complex puzzle, and this challenge is what drives my enthusiasm for organic chemistry and synthesis.

**SYNFORM** *What do you think about the modern role and prospects of organic chemistry?*

**Prof. Dr. T. Gulder** Organic chemistry, despite its long history, remains a field of endless discovery and innovation. It creates new tools that push the pharmaceutical, agrochemical, and material industries, impacting other scientific disciplines and everyone's lives. This significant impact comes with great responsibilities. One of the main aims of modern organic synthesis is to rethink chemical feedstocks, processes, and products concerning environmental safety and sustainability. This makes organic chemistry a perpetually exciting and rewarding science area, offering endless exploration opportunities and a promising future.

**SYNFORM** *Please comment on your role as a member of the Editorial Board of SYNLETT.*

**Prof. Dr. T. Gulder** As I am new on the SYNLETT Editorial Board, I am still figuring out my specific role. Besides the daily business of handling manuscripts, I started to develop on the first thematic issue for the journal, which may be devoted to enzymes in organic reactions, an emerging field in organic synthesis.

**SYNFORM** *Finally, on a personal note, what do you do in your free time?*

**Prof. Dr. T. Gulder** In my free time, I like to read books, do gardening, and cook together with my family. My most favorite thing is to spend time with my daughter and my husband outside.



Mattes Fank

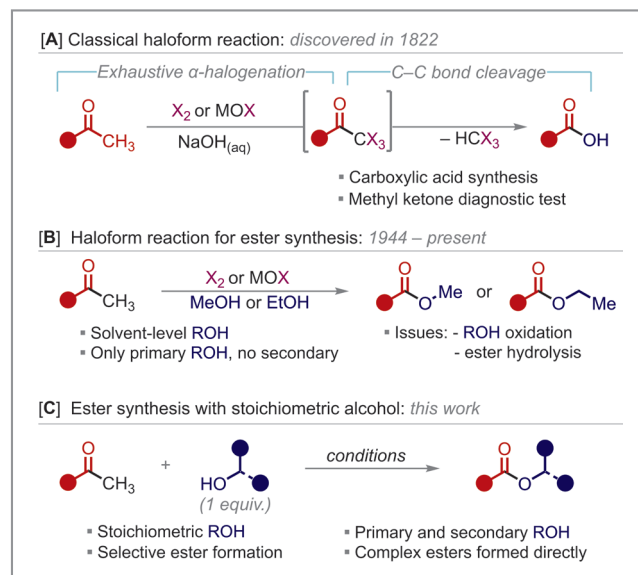
# A Stoichiometric Haloform Coupling for Ester Synthesis with Secondary Alcohols

*Angew. Chem. Int. Ed.* **2024**, *63*, e202400570

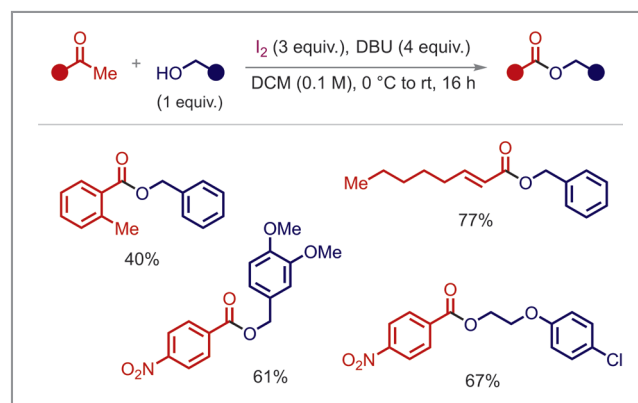
The haloform reaction, first reported in 1822 by Georges Simon Serullas, is a staple of undergraduate organic chemistry courses taught in universities. It takes methyl ketones to carboxylic acids through the triple  $\alpha$ -halogenation of the methyl group to form a good nucleofuge, which is replaced by hydroxide (Scheme 1A). This reaction has been widely applied in the pharmaceutical and fragrance industries. It has since evolved to form methyl or ethyl esters when the reaction is performed in methanol or ethanol, respectively (Scheme 1B). “Undergraduate textbooks tell you this is all that is possible,” said Dr. Alastair Lennox (University of Bristol, UK), who – together with his group – wondered why it was not possible to use more complicated and extravagant alcohols to make more interesting esters. “We conducted an in-depth literature search on whether it had been done before,” said Dr. Lennox, continuing: “When my PhD student, Bert [Albert Rowett, first author of the title paper], came back to me to tell me it had not been done, I told him to go back and look again as I could not believe there was nothing on it. As well as finding no examples of complicated alcohols, we also found that secondary alcohols had hardly been attempted; we found one example of the use of isopropanol. Hence, based on these surprisingly limited publications, we were inspired to explore whether it was possible to derive conditions with one equivalent of a complex primary or secondary alcohol (Scheme 1C).”

When the authors first started exploring this reaction, they realised that carboxylic acid would competitively form very readily and the alcohol nucleophile would oxidise, removing it from the reaction and providing the opportunity for other esters to form. Dr. Lennox explained: “We explored many different combinations of halogenating reagents and bases, and eventually found that  $I_2$  with DBU was especially proficient and facilitated 1 equivalent of our model primary alcohol to be used in the reaction (Scheme 2). This is because it was possible to dry these reagents well, which avoids the formation of carboxylic acid, and also slow the rate of alcohol oxidation down relative to the desired iodination of the methyl ketone.”

However, when applying these optimised conditions to secondary alcohols, the yield decreased significantly. “To understand why this occurred and how to adapt our conditions, we turned to kinetic studies and modelling,” remarked Dr.

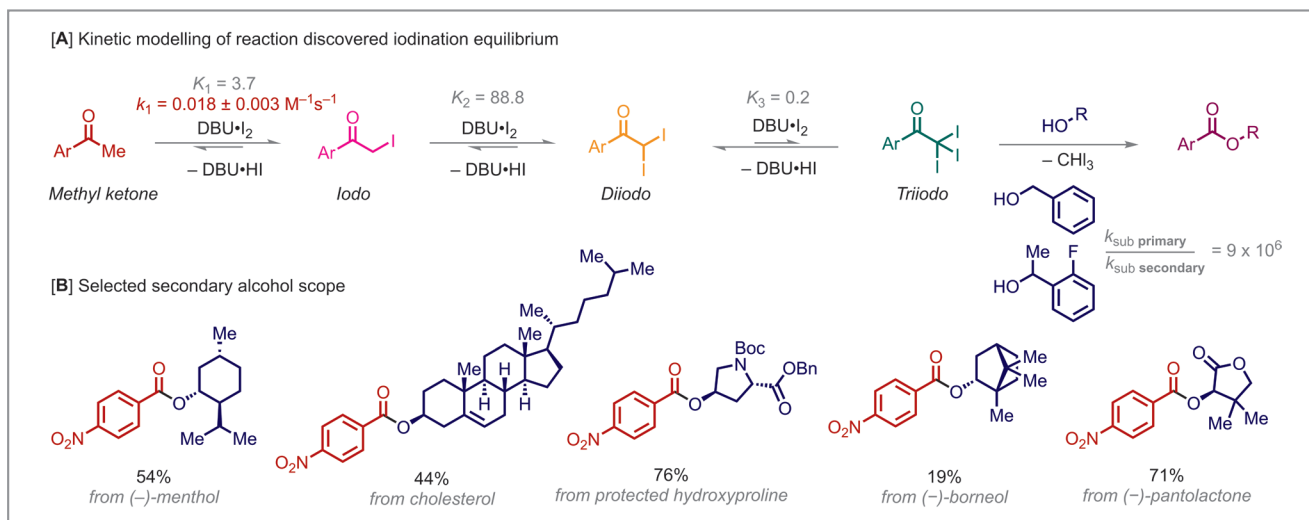


**Scheme 1** The haloform reaction forms carboxylic acids and esters with solvent-level primary alcohol from methyl ketones. This work presents the synthesis of new complex esters using stoichiometric primary and secondary alcohols.



**Scheme 2** The haloform reaction conducted with 1 equivalent of a primary alcohol, selected examples

Lennox. He continued: “Through this, we discovered that the iodination steps were under equilibrium, which became apparent with secondary alcohols because the final substitution



**Scheme 3** Mechanistic studies led to the development of conditions for secondary alcohols

step is slower: the deiodination of the triiodo species started to compete with the substitution step that forms the ester (Scheme 3). This insight enabled a very simple solution, which was to increase the concentration of  $\text{DBU}\cdot\text{I}_2$  to increase the concentration of reactive triiodo species through Le Chatelier's principle, and therefore improve the rate of substitution for secondary alcohols. This change facilitated a range of complex secondary alcohols to be used in the reaction." Dr. Lennox concluded: "We now wonder how we could further change the reaction to work with different nucleophiles, increasing the available chemical space of this discovery."

*Matthew Fenske*

## About the authors



Dr. A. C. Rowett

**Albert C. Rowett** studied chemistry at the University of Bristol, UK (MSci, 2018). He then joined the EPSRC Centre for Doctoral Training (CDT) in Catalysis (University of Bath, University of Bristol and Cardiff University, UK), returning to Bristol to complete his PhD research under the supervision of Dr. Alastair Lennox. His research focused on (electro)halogenation reactions and, in particular, the development of the novel haloform coupling methodology discussed here.



S. G. Sweeting

**Stephen G. Sweeting** received his MSci Chemistry degree from the University of Bristol (UK) in 2021. He is now pursuing his PhD under the supervision of Dr. Alastair Lennox, also at the University of Bristol. His main area of study focuses on understanding fluorination reactions involving hydrogen fluoride complexes, as well as developing new electrochemical methodologies to achieve selective fluorinations.



Dr. D. M. Heard

**David M. Heard** studied at the University of Sheffield, UK (MChem, 2014) and completed a PhD in chemical synthesis at the University of Bristol, UK (2019), researching the structural elucidation and total synthesis of maleidride natural products with Prof. Chris Willis. David completed postdoctoral studies with Dr. Alastair Lennox (University of Bristol) on electrosynthetic methods and reactor technology, and with Prof. Dennis Hall (University of Alberta, Canada) investigating photoredox catalysis in organoboron chemistry. David now works in process development in the pharmaceutical industry.



Dr. A. J. J. Lennox

**Alastair J. J. Lennox** attained his PhD from the University of Bristol, UK (Prof. Guy Lloyd-Jones) and did postdoctoral studies in Rostock, Germany (Prof. Matthias Beller) as an Alexander von Humboldt Fellow and at the University of Wisconsin, Madison, USA (Prof. Shannon Stahl). In 2018, Alastair returned to the University of Bristol as a Royal Society University Research Fellow to start his independent research programme.

He was promoted to Associate Professor of Chemistry in 2022. His group are interested in the development of novel synthetic organic methods with sustainability and mechanism as themes that strongly underpin their approach to this. Specific interests include the exploration of electrochemistry as a tool for performing selective redox transformations, and also in the development of fluorination reactions and fluorinated building blocks.



## Young Career Focus: Dr. Bernd M. Schmidt (Heinrich Heine Universität, Germany)

**Background and Purpose.** SYNFORM regularly meets young up-and-coming researchers who are performing exceptionally well in the arena of organic chemistry and related fields of research, in order to introduce them to the readership. This Young Career Focus presents Dr. Bernd M. Schmidt (Heinrich Heine Universität, Germany).

### Biographical Sketch

Photo © NRW AWKI | Engel-Albustin



Dr. B. M. Schmidt

**Bernd M. Schmidt** completed his Ph.D. studies with Dieter Lentz at the Freie Universität Berlin, Germany, and with Hidehiro Sakurai at the Institute for Molecular Science in Okazaki, Japan, where he worked on aromatic buckybowls. Bernd received a Humboldt research fellowship to work with Makoto Fujita at the University of Tokyo, Japan, and Stefan Hecht at the Humboldt University of Berlin, Germany. Since February 2018, he has been an independent research group leader at Heinrich Heine Universität (HHU), Düsseldorf, Germany. He became a member of the Young Academy of the North Rhine-Westphalian Academy of Sciences, Humanities, and the Arts in 2020, where he served as speaker from 2021–2023. In 2023, he was accepted into the prestigious Heisenberg Programme from the German Research Foundation (DFG) to continue his group's work investigating functional and responsive supramolecular systems. Bernd is Germany's Young Investigator 2020 of the EuChemS Division of Organic Chemistry (YIW2020) and is a recipient of the Dr. Otto Röhm Memorial Foundation Award (2022), a JSPS BRIDGE Fellowship (2022), and a Thieme Chemistry Journals Award (2024).

### INTERVIEW

**SYNFORM** Which field of organic chemistry are you interested in the most and why?

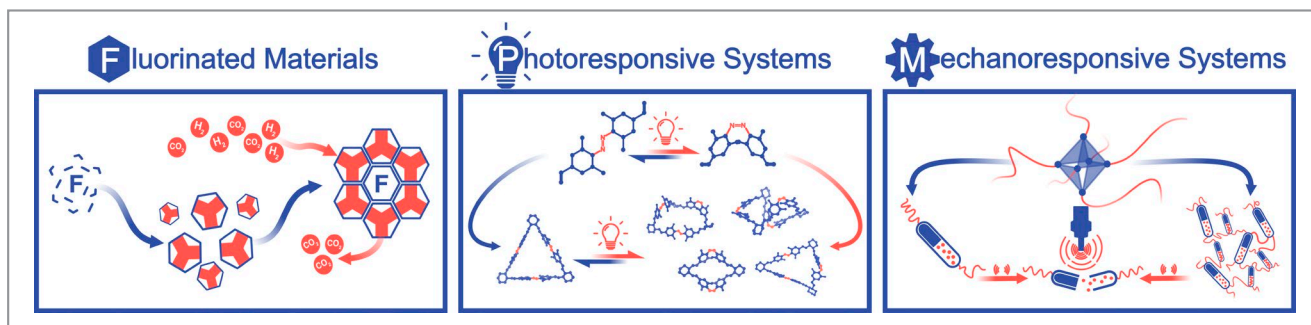
**Dr. B. M. Schmidt** I am particularly interested in supramolecular chemistry within the field of organic chemistry. In our research group, we leverage self-assembly to develop novel compounds and materials, spanning from individual molecules to intricate assemblies. Our aim is to explore the interdisciplinary applications of supramolecular structures in materials chemistry, photochemistry, macromolecular chemistry, and biochemistry.

**SYNFORM** Following that, what is the focus of your current research activity?

**Dr. B. M. Schmidt** My research group deals with various aspects of supramolecular chemistry (Figure 1), such as porous imine-based organic compounds containing fluorinated motifs, which we were able to establish with a series of publications. This is complemented by our pioneering work in the field of mechanoresponsive macromolecular supramolecular systems for releasing guests from aqueous solutions, as well as supramolecular building blocks that react to visible light. We combine the possibilities of organic synthesis chemistry with the self-organisation of complex molecular structures and molecular recognition in solution and in the solid state.

In the beginning of 2024, we published our work on multifold post-modification of macrocycles and cages by isocyanate-induced azadefluorination cyclisation. For this project, we designed suitable building blocks using organic chemistry knowledge, self-assembled them using dynamic-covalent imine chemistry, and transformed the supramolecular structures into (non-dynamic) covalent compounds by reduction with sodium borohydride. Reactive isocyanates can be used to introduce functional groups into the cages





**Figure 1** Current areas of research in the Schmidt group: fluorinated materials, photoswitchable supramolecular assemblies, and ultrasound-induced mechanochemistry in complex systems.

and macrocycles, leading to rigid and chemically stable molecules that can undergo further reactions. It is a dream to use concepts like this, combining the strength of supramolecular chemistry, the one-step self-assembly of complex structures, with the power of organic chemistry to make further precise modifications thereof.

**SYNFORM** *What do you think about the modern role and prospects of organic chemistry?*

**Dr. B. M. Schmidt** Organic chemistry provides the foundation for understanding the structure, reactivity, and synthesis of organic molecules, which are essential building blocks in supramolecular systems. Supramolecular chemists can then design and construct a wide range of molecular architectures with tailored properties and functions. In terms of prospects, organic chemistry continues to play a vital role in driving innovation and advancing scientific knowledge, extending over various fields, particularly within the dynamic and rapidly evolving field of supramolecular chemistry.

**SYNFORM** *Which difficulties are there for young upcoming chemists in your field? Do you have any tips?*

**Dr. B. M. Schmidt** The status of junior researchers in Germany, particularly in the phase before obtaining a lifetime professorship, is currently varied and often not aligned with their responsibilities in this important qualification phase. In contrast to counterparts abroad, who often have clearly defined roles (e.g., assistant professor) with specified rights and duties, Germany lacks nationwide uniform standards. Even when standards are proposed by funding organisations, such as for Emmy Noether research groups, implementation might vary. As a result, individuals frequently need to advocate for their rights in order to carry out their work effectively.

I co-authored a recent discussion paper on this topic in German ([link](#)).

**SYNFORM** *What is your most important scientific achievement to date and why?*

**Dr. B. M. Schmidt** My most significant achievement lies in the dedicated effort I've invested in nurturing and guiding the next generation of scientists. I take pride in training and preparing them for the challenges they may encounter in their future roles, regardless of the specific positions they choose to pursue. I am very grateful to see how they surpass themselves and find success.

The most important scientific achievement is always the next big project or paper for me, I do not like looking back.

**SYNFORM** *Could you tell us something about yourself outside the lab, such as your hobbies or extra-work interests?*

**Dr. B. M. Schmidt** I am an absolutely crazy foodie, and I love cooking (and eating), exploring things, restaurants, and everything. I also got into wine during the pandemic, so there's a whole new field to explore for me, too. I even interned in a restaurant before studying chemistry because I was wondering if I should become a chef.

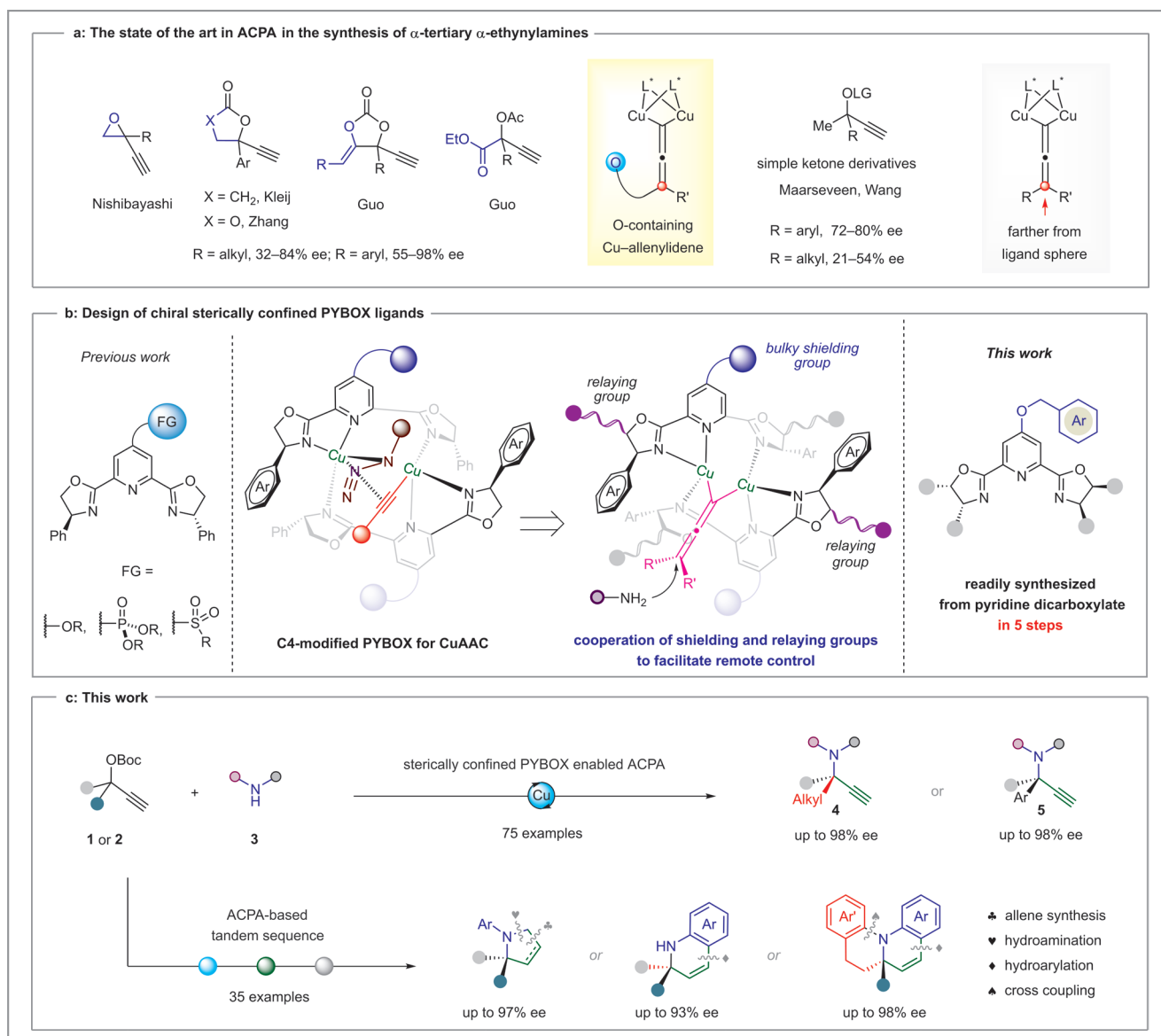
*Mattias Fenske*

# Enantioselective Propargylic Amination and Related Tandem Sequences to $\alpha$ -Tertiary Ethynylamines and Azacycles

*Nat. Chem.* **2024**, *16*, 521–532

Optically active amines and azacycles are widely present in drugs, bioactive compounds and agrochemicals. The synthesis of chiral  $\alpha$ -tertiary amines and related azacycles in sufficient structural diversity is important for drug development, but

constitutes a long-term challenge confronting organic chemists. “Enantioenriched  $\alpha$ -tertiary propargylamines featuring an  $\alpha$ -ethynyl substituent are valuable platform molecules, because their sp C–H bonds and carbon–carbon triple bonds



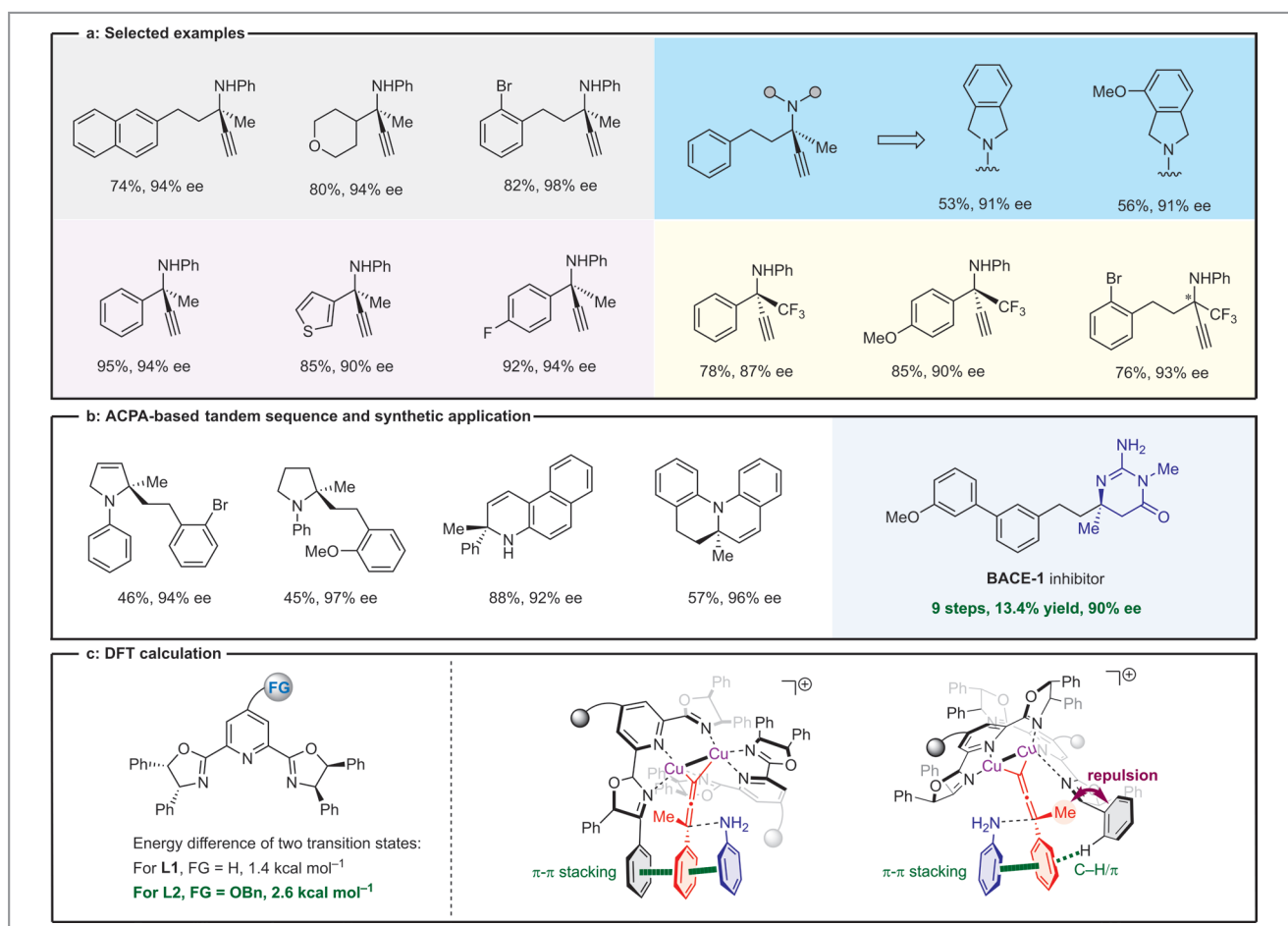
**Scheme 1** Working hypothesis

permit versatile diversifying reactions,” said Professor Jian Zhou [East China Normal University (ECNU), P. R. of China], adding: “Nevertheless, their facile access is undeveloped in terms of enantioselectivity, substrate scope and synthetic efficiency, as compared with internal propargylamines.”

According to Professor Zhou, asymmetric Cu(I)-catalysed propargylic amination (ACPA) is a promising strategy to access enantioenriched ethynylamines, following the seminal independent work of Marseveen,<sup>1a</sup> Nishibayashi<sup>1b</sup> and colleagues. However, despite previous enlightening studies, there remains much to explore (Scheme 1a).<sup>2</sup> This is because only substrates leading to Cu–allenylidene intermediates with both an aryl and an oxygen-containing moiety could give satisfactory results, while aliphatic compounds and, especially, simple ketone-derived propargylic alcohol derivatives still represent problematic substrates.<sup>3</sup> Moreover, the modular combination of ACPA into tandem sequences for diverse synthesis of azacycles remains an untrodden path.

“As the prochiral carbon in the Cu–allenylidene intermediate is further away from the ligand sphere, how to achieve excellent remote enantiofacial control is the key to success,” said Professor Zhou. He continued: “With our experience in asymmetric CuAAC reactions realized by PYBOX ligands featuring a bulky C4 shielding group, along with the fact that both reactions are proposed to involve di-copper catalysis, we wondered whether these C4-modified PYBOX ligands were effective for ACPA of challenging substrates.” Professor Zhou further stated: “Initial trials using PYBOX ligands with a single C4 shielding group all failed, thus prompting us to exploit sterically more confined PYBOX ligands by installing relaying groups on the 5-position of both oxazoline rings, to form a sterically constrained microenvironment to relay stereochemical information from the ligand chirality through steric interactions (Scheme 1b).”

Indeed, sterically confined PYBOX ligands with both C4 shielding group and relaying groups can achieve excellent



**Figure 1** Substrate expansion in ACPA reaction and ACPA-based tandem sequence

remote enantiofacial discrimination of the Cu–allenylidenes, thus allowing the highly enantioselective ACPA of aliphatic and alkyl aryl ketone-derived propargylic carbonates with a variety of primary and secondary amines (Scheme 1c). “The ACPA-based tandem sequences were also established as a new strategy to synthesize various chiral azacycles, such as dihydropyrroles, tetrahydropyrroles, dihydroquinines, or tetrahydroquinines with an aza-quaternary stereocenter (Figure 1a),” said Professor Zhou, adding: “The value of the thus-obtained chiral  $\alpha$ -tertiary  $\alpha$ -ethynylamines was demonstrated by various transformations and the enantioselective total synthesis of a selective multi-target  $\beta$ -secretase inhibitor (Figure 1b).” A collaboration with Prof. Xin Wang on theoretical calculations revealed that both the C4 bulky benzyloxy group and the phenyl relaying group on the oxazoline ring are responsible for the high enantioselectivity (Figure 1c). Professor Zhou explained: “With the combination of C4 benzyloxy and the phenyl relaying group, a more sterically confined chiral pocket should be formed, which enhances the favorable  $\pi$ – $\pi$  stacking interactions and also the unfavorable repulsion interaction, thus resulting in a higher free energy gap of 2.6 kcal/mol.”

Professor Zhou emphasized that the working model of sterically confined PYBOX ligands could also be extended to the asymmetric propargylic substitution of  $\alpha$ -CF<sub>3</sub> propargylic carbonate with O- and C-centered nucleophiles. “This work provides a flexible solution to develop asymmetric propargylic substitutions involving metal–allenylidene intermediates,” said Professor Zhou. He concluded: “Since the pioneering work of Nishibayashi, Hidai, Uemura and co-workers,<sup>5</sup> this field has grown exponentially, but the reaction of propargylic tertiary alcohol derivatives to  $\alpha$ -ethynyl quaternary stereocenters is still in its infancy. Considering the good performance of sterically confined PYBOX, there are enormous opportunities for the asymmetric propargylic substitution reactions via the development of new PYBOX ligands.”

*Mattias Farnok*

## REFERENCES

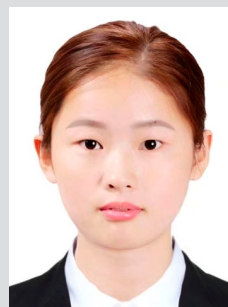
- (1) (a) R. J. Detz, M. M. E. Delville, H. Hiemstra, J. H. van Maarseveen *Angew. Chem. Int. Ed.* **2008**, *47*, 3777–3780. (b) G. Hattori, H. Matsuzawa, Y. Miyake, Y. Nishibayashi *Angew. Chem. Int. Ed.* **2008**, *47*, 3781–3783.
- (2) (a) G. Hattori, A. Yoshida, Y. Miyake, Y. Nishibayashi *J. Org. Chem.* **2009**, *74*, 7603–7607. (b) J. E. Gómez, W.-S. Guo, S. Gaspa, A. W. Kleij *Angew. Chem. Int. Ed.* **2017**, *56*, 15035–15038. (c) L. Tian, L. Gong, X. Zhang *Adv. Synth. Catal.* **2018**, *360*, 2055–2059. (d) W. Guo, L. Zuo, M. Cui, B. Yan, S. Ni *J. Am. Chem. Soc.* **2021**, *143*, 7629–7634. (e) T. Liu, S. Ni, W. Guo *Chem. Sci.* **2022**, *13*, 6806–6812.
- (3) (a) R. J. Detz, Z. Abiri, R. Griel, H. Hiemstra, J. H. van Maarseveen *Chem. Eur. J.* **2011**, *17*, 5921–5930. (b) Q. Cai, H. Rao, S.-J. Li, Y. Lan, K. Ding, X. Wang *Chem* **2024**, *10*, 265–282.
- (4) (a) R.-Y. Zhu, L. Chen, X.-S. Hu, F. Zhou, J. Zhou *Chem. Sci.* **2020**, *11*, 97–106. (b) K. Liao, Y. Gong, R.-Y. Zhu, C. Wang, F. Zhou, J. Zhou *Angew. Chem. Int. Ed.* **2021**, *60*, 8488–8493. (c) Y. Gong, C. Wang, F. Zhou, K. Liao, X.-Y. Wang, Y. Sun, Y.-X. Zhang, Z. Tu, X. Wang, J. Zhou *Angew. Chem. Int. Ed.* **2023**, *62*, e202301470.
- (5) Y. Nishibayashi, G. Onodera, Y. Inada, M. Hidai, S. Uemura *Organometallics* **2003**, *22*, 873–876.

## About the authors



Z. Zhang

**Zheng Zhang** was born in 1997 in Shandong province of China. After obtaining his bachelor's degree in 2019 from Qufu Normal University (P. R. of China), he joined Professor Jian Zhou's group in East China Normal University (P. R. of China). His research interests focus on the asymmetric propargyl substitution reaction and developing new PY-BOX ligands.



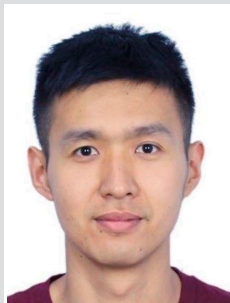
Y. Sun

**Ying Sun** was born in Shandong province, P. R. of China. She obtained her bachelor's degree in chemistry in 2021 from Qufu Normal University (P. R. of China). Since 2021, she has been studying for a master's degree at East China Normal University (P. R. of China) in the group of Prof. Jian Zhou.

>>



## About the authors



Dr. Y. Gong

**Yi Gong** received his Master's degree from the Guizhou University (P. R. of China) in 2018. Then he moved to East China Normal University (P. R. of China) to pursue his Ph.D. in the group of Jian Zhou. He received his Ph.D. in 2023. Currently, he is a postdoctoral fellow with Prof. Jian Zhou. His work revolves around the catalytic asymmetric synthesis of fully substituted carbon stereocenters.



D. Tang

**Daliang Tang** received his Master's degree from Yangzhou University (P. R. of China) in 2017. He then worked for Wuxi AppTec for four years and in 2021, he moved to East China Normal University (P. R. of China) to pursue his Ph.D. in the group of Jian Zhou. His work revolves around the catalytic asymmetric synthesis of the fully substituted carbon stereocenters and synthesis of novel chiral Pybox ligands.



H. Luo

**Hui Luo** was born in Jiangxi province, P. R. of China. He obtained his bachelor's degree in chemistry in 2021 from Yichun University (P. R. of China). Since 2021, he has been studying for a master's degree in East China Normal University (P. R. of China) under the supervision of Professor Feng Zhou in the group of Professor Jian Zhou. Now he works in Shanghai Yuanhua Pharmaceutical Technology Co., Ltd.



Z. Zhao

**Zhi-Peng Zhao** obtained his bachelor's degree from the China West Normal University (P. R. of China) in 2020. He then pursued his master's degree in the group of Prof. Jian Zhou at East China Normal University (P. R. of China), where he worked on organic synthesis methods. He received his master's degree in 2023.



Dr. F. Zhou

**Feng Zhou** was born in 1987 in Shandong province, P. R. of China. After obtaining his bachelor's degree in 2009 from Sichuan Normal University (P. R. of China), he joined Professor Jian Zhou's group at East China Normal University (P. R. of China) and obtained his PhD in 2014. Subsequently, he embarked on his academic career at East China Normal University and has been a professor there since 2021. His research interest centers on the catalytic asymmetric construction of tetrasubstituted stereogenic centers and the utilization of CO<sub>2</sub>.



Dr. X. Wang

**Xin Wang** obtained his PhD in 2002 from Sichuan University (P. R. of China). After spending two years working as a postdoctoral fellow with Professor Yuansheng Jiang and Professor Shuhua Li at Nanjing University (P. R. of China), he joined Sichuan University in 2004. He has been a professor at Sichuan University since 2014. His current research interests include molecular design and simulation of materials and theoretical study of organocatalytic reaction mechanism.



Prof. J. Zhou

**Jian Zhou** obtained his PhD in 2004 from the Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences (P. R. of China), under the guidance of Professor Yong Tang. After spending one year working as a postdoctoral fellow with Professor Shū Kobayashi at the University of Tokyo (Japan), and three years with Professor Benjamin List at Max-Planck-Institut für Kohlenforschung (Germany), he joined the Shanghai Key Laboratory of Green Chemistry and Chemical Processes at East China Normal University (P. R. of China) as a professor from the end of 2008. His group focuses on the development of new chiral catalysts and new catalytic asymmetric reactions allowing the catalytic, economical and enantioselective construction of tetrasubstituted stereogenic carbon centers.

## Coming soon

— Literature Coverage

### Phase-Transfer Catalysts Shift the Pathway to Transmetalation in Biphasic Suzuki–Miyaura Cross-Couplings

— Literature Coverage

### Copper-Catalysed Perarylation of Cyclopentadiene: Synthesis of Hexaaryl cyclopentadienes

— Literature Coverage

### Radical-Mediated $\alpha$ -*tert*-Alkylation of Aldehydes by Consecutive 1,4- and 1,3-(Benzo)thiazolyl Migrations

## Further highlights

**Synthesis** Review: 1,2-*trans*-Diaminocyclohexane (DACH) in Asymmetric Catalysis: Nearing Fifty Years of Faithful Service and Counting

(by S. Hanessian)

**Synlett** Account: Acetal Substitution Reactions: Stereoelectronic Effects, Conformational Analysis, Reactivity vs Selectivity, and Neighboring-Group Participation

(by K. A. Woerpel and co-workers)

**Synfacts** Synfact of the Month in category “Innovative Drug Discovery and Development”: Discovery and Development of CNS-Penetrant IRAK4 Inhibitors

For current SYNFORM articles, please visit [www.thieme-chemistry.com](http://www.thieme-chemistry.com)  
SYNFORM issue 2024/10 is available from September 13, 2024  
at [www.thieme-connect.com/ejournals](http://www.thieme-connect.com/ejournals)

#### Editor

Matteo Zanda  
C.N.R. – Istituto di Scienze e Tecnologie Chimiche (SCITEC)  
Via Mancinelli, 7, 20131 Milano, Italy  
Editorial Assistant: Alison M. Sage, [synform@outlook.com](mailto:synform@outlook.com)

#### Editorial Office

— Senior Director SYNTHESIS, SYNLETT, SYNFACTS:  
Susanne Haak, [susanne.haak@thieme.de](mailto:susanne.haak@thieme.de)  
— Scientific Editor:  
Selena Boothroyd, [selena.boothroyd@thieme.de](mailto:selena.boothroyd@thieme.de)  
— Development Editor:  
Stefanie Baumann, [stefanie.baumann@thieme.de](mailto:stefanie.baumann@thieme.de)  
— Editorial Assistant:  
Sabine Heller, [sabine.heller@thieme.de](mailto:sabine.heller@thieme.de)  
— Typesetting:  
Ulrike Holzwarth, Büro für Gestaltung, Stuttgart  
— Postal Address: Thieme Chemistry Journals, Georg Thieme Verlag KG,  
Rüdigerstraße 14, 70469 Stuttgart, Germany.  
— Homepage: [www.thieme-chemistry.com](http://www.thieme-chemistry.com)

#### Publication Information

Synform will be published 12 times in 2024 by Georg Thieme Verlag KG, Rüdigerstraße 14, 70469 Stuttgart, Germany, and is an additional online service for Synthesis, Synlett and Synfacts.

#### Product Names

Product names which are in fact registered trademarks may not have been specifically designated as such in every case. Thus, in those cases where a product has been referred to by its registered trademark it cannot be concluded that the name used is public domain. The same applies to labels, names or other signs.

#### Ordering Information for Synthesis, Synlett and Synfacts

The Americas: Thieme New York, 333 Seventh Avenue, New York, NY 10001, USA.  
Via e-mail: [customerservice@thieme.com](mailto:customerservice@thieme.com)  
Via website: [www.thieme-chemistry.com](http://www.thieme-chemistry.com)  
Phone: +1 212 760 0888; Fax: +1 212 947 0108  
Order toll-free within the USA: +1 800 782 3488

Europe, Africa, Asia, and Australia: Georg Thieme Verlag KG, Rüdigerstraße 14, 70469 Stuttgart, Germany.  
Via e-mail: [customerservice@thieme.de](mailto:customerservice@thieme.de)  
Via website: [www.thieme-chemistry.com](http://www.thieme-chemistry.com)  
Phone: +49 711 8931 421; Fax: +49 711 8931 410

Current list prices are available through [www.thieme-chemistry.com](http://www.thieme-chemistry.com).

#### Online Licensing Information

The online versions of Synform as well Synthesis, Synlett, Synfacts and SynOpen are available through [www.thieme-connect.com/products/ejournals/journals](http://www.thieme-connect.com/products/ejournals/journals) where it is also possible to register for a free trial account. For information on multi-site licenses and pricing for corporate customers, as well as backfiles, please contact our regional offices:

The Americas: [esales@thieme.com](mailto:esales@thieme.com), phone: +1 212 584 4695  
Europe, Africa, Asia (ex. India), and Australia: [eproducts@thieme.de](mailto:eproducts@thieme.de),  
phone: +49 711 8931 407  
India: [eproducts@thieme.in](mailto:eproducts@thieme.in), phone +91 120 455 6600

#### Manuscript Submission to Synthesis, Synlett, and SynOpen

Manuscript submissions will be processed exclusively online via <http://mc.manuscriptcentral.com/synthesis>, <http://mc.manuscriptcentral.com/synlett> and <http://mc.manuscriptcentral.com/synopen>, respectively. Please consult the Instructions for Authors before compiling a new manuscript. The current version and the Word template for manuscript preparation are available for download at [www.thieme-chemistry.com](http://www.thieme-chemistry.com).

#### Ownership and Copyright

© 2024. Thieme. All rights reserved.  
This publication, including all individual contributions and illustrations published therein, is legally protected by copyright for the duration of the copyright period. Any use, exploitation or commercialization outside the narrow limits set by copyright legislation, without the publisher's consent, is illegal and liable to criminal prosecution. This applies in particular to photocopy reproduction, copyright, cyclostyling, mimeographing or duplication of any kind, translating, preparation of microfilms, and electronic data processing and storage (CD-ROM, DVD, USB memory stick, databases, cloud-based service, ebook and other forms of electronic publishing) as well as making it available to the public (e.g., internet, intranet or other wired or wireless data networks), in particular by displaying on stationary or mobile visual display units, monitors, smart phones, tablets or other devices by download (e.g., e-pub, PDF, App) or retrieval in any other form.

#### Copyright Permission for Users in the USA

Authorization to photocopy items for internal or personal use, or the internal or personal use of specific clients, is granted by Georg Thieme Verlag KG Stuttgart · New York for libraries and other users registered with the Copyright Clearance Center (CCC) Transactional Reporting Service; [www.copyright.com](http://www.copyright.com). For reprint information in the USA, please contact: [journals@thieme.com](mailto:journals@thieme.com)