



Cite this: RSC Adv., 2023, 13, 14249

2-Oxindole and related heterocycles: synthetic methodologies for their natural products and related derivatives

Shivangi Sharma, ^a Yukti Monga, ^b Ashu Gupta ^b and Shivendra Singh ^{*a}

Natural goods, medications, and pharmaceutically active substances all contain substituted oxindoles. Generally, the C-3 stereocenter of the substituents of oxindoles and their absolute arrangement have a substantial impact on the bioactivity of these substances. In this case, the desire for contemporary probe and drug-discovery programs for the synthesis of chiral compounds using desirable scaffolds with high structural diversity further drives research in this field. Also, the new synthetic techniques are generally simple to apply for the synthesis of other similar scaffolds. Herein, we review the distinct approaches for the synthesis of diverse useful oxindole scaffolds. Specifically, the research findings on the naturally existing 2-oxindole core and a variety of synthetic compounds having a 2-oxindole core are discussed. We present an overview of the construction of oxindole-based synthetic and natural products. In addition, the chemical reactivity of 2-oxindole and its related derivatives in the presence of chiral and achiral catalysts are thoroughly discussed. The data compiled herein provides broad information related to the bioactive product design, development, and applications of 2-oxindoles and the reported techniques will be helpful for the investigation of novel reactions in the future.

Received 4th April 2023
Accepted 25th April 2023
DOI: 10.1039/d3ra02217j
rsc.li/rsc-advances

1. Introduction

Oxindoles (1, Fig. 1) are a group of endogenous hetero-aromatic organic compounds (quinoline,^{1–3} indole,⁴ etc.) found in the tissues and bodily fluids of mammals and in the natural

products in different plants.⁵ The term “oxindole” and its derivatives are known as “1,3-dihydro-2H-indole-2-one(s)”⁶ because their structure consists of a six-membered benzene ring fused with a five-membered pyrrole ring and a carbonyl group at the C-2 position. It is well known that oxindoles have

^aDepartment of Applied Chemistry, Amity School of Engineering and Technology, Amity University Madhya Pradesh, Gwalior, Madhya Pradesh-474 005, India. E-mail: shivendrasngh0@gmail.com

^bShyamal College, Department of Chemistry, University of Delhi, Delhi-110032, India



Ms Shivangi Sharma has obtained her MSc in July 2021 from the School of Studies in Chemistry, Jiwaji University Gwalior (MP). Since then, she has been pursuing her PhD in Synthetic Organic Chemistry and Molecular Design at Amity University Madhya Pradesh. She is experienced in organic synthesis and molecular docking study using several software such as Autodock-Vina 4, Pymol,

& Discovery Studio Visualizer. She completed the “CSIR-SUMMER RESEARCH TRAINING PROGRAM” (CSIR-SRTP) JUNE–AUGUST 2020 and her research interests include synthesis, molecular design and biological activity of heterocyclic derivatives.



Dr Yukti Monga received her BSc (2007) from the University of Delhi. She obtained first rank in her Master's (2009) from Jiwaji University. She completed her PhD in Chemistry from the University of Delhi in 2016. She was awarded an Inspire fellowship from DST, INDIA. She has many publications in reputed international journals and book chapters with World Scientific and Wiley. Her research inter-

ests include the sustainable development of nanomaterials and their use in heterogeneous catalytic reactions.



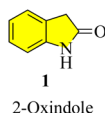


Fig. 1 Structure of oxindole.

two rapidly tautomerizing hydroxyl isomers.^{7–9} Several synthetic methods have been successfully applied to develop various derivatives and scaffolds with a range of biological activities, in addition to natural methods to obtain the oxindole nucleus.

New oxindole compounds with a distinct pharmacological profile and commendable efficacy are of great interest in many sectors of the pharmaceutical industry and academia. Nintedanib (2), a marketed highly potent drug, was recently approved in March 2020 in the United States for the treatment of interstitial lung diseases such as idiopathic pulmonary fibrosis (IPF) and chronic fibrosis with a progressive phenotype. It is one of the most potent indolinone compounds and has an effective antiproliferative property, which inhibits angiokine and limits the growth factor. Nintedanib is advertised commercially under the brand names “Ofev” and “Vargatef”.¹⁰ Sunitinib (3), a tiny molecule that primarily serves as a tyrosine kinase inhibitor and a well-known treatment for renal cell carcinoma and gastrointestinal stromal tumours, is another notable oxindole derivative. It was clinically approved by the Food and Drug Administration (FDA) in 2006 and is the first anticancer medicine to be approved for use on two distinct types of cancer cell lines simultaneously. As part of its mechanism of action, it indirectly targets numerous receptor tyrosine kinases in an effort to suppress cellular signaling.¹¹ The phase III trials for semaxanib (7), a tyrosine kinase inhibitor that targets angiogenesis and colon-rectal cancer *via* the vascular endothelial growth factor pathway, were unsuccessful.¹² Ropinirole (6), a commonly prescribed drug for the treatment of Parkinson's disease and restless legs syndrome (RLS), is a well-known

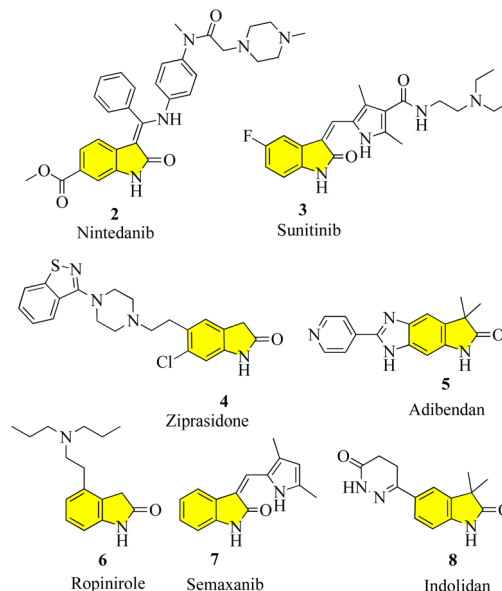


Fig. 2 Structure of commercially available drugs with an oxindole core.

medication, which contains the widely used active ingredient oxindole. It was developed in 1996 and functions as a complete agonist at D₂, D₃, and D₅ receptors, as well as a dopamine receptor (D₂) agonist. However, it has a relatively lower affinity for D₁ and D₅ receptors. The success of ropinirole as a medication is attributed to the characteristics of its structure, which are beneficial to its functionality, such as low molecular weight, accessibility, and a stereocenter-free chemical structure, attracting additional attention in ongoing research.¹³ Another oxindole derivative called ziprasidone (4) is a novel antipsychotic medication produced by the American pharmaceutical behemoth Pfizer and marketed and sold under the trade name “Geodon”. Presently, ziprasidone has received FDA approval for



Dr Ashu Gupta completed her PhD from the Department of Chemistry, University of Delhi, India in 2004 under the guidance of Prof. R. K. Sharma. Currently, she is an Associate Professor and Coordinator of the DBT Research Project in Shyam Lal College, University of Delhi. She is an active member of the Green Chemistry Network Centre. She has more than 20 year teaching and research

experience. Her research interests focus on sustainable material development for catalysis, sensors, and other energy applications.



Dr Shivendra Singh received his PhD in Organic Chemistry from the Department of Chemistry, Indian Institute of Technology (IIT), Indore, India in 2017. He worked as a Post-Doctoral Fellow at the Indian Institute of Science Education & Research (IISER), Mohali, India, and was a recipient of a National Post-Doctoral Fellowship (NPDF) from the Science and Engineering Research Board (SERB),

Department of Science and Technology (DST), India. Currently, he is working as an Assistant Professor-II at the Department of Applied Chemistry at Amity School of Applied Sciences, Amity University Madhya Pradesh, India. His research interests include molecular docking, synthesis, and application of organic molecules.



use in psychotherapy and the treatment of mental illnesses including schizophrenia and severe manic behavior. Ziprasidone (**4**) works similarly to ropinirole by blocking the dopamine (D2) receptor.¹⁴ Cardiovascular diseases and ischemic chest pains are actively treated with some other drugs that have potent vasodilatory and advantageous inotropic effects, notably adibendan (**5**) and indolidan (**8**) (Fig. 2).¹⁵

2. Natural products & biologically active compounds

To maximize the achievements in current probe- and drug-discovery studies, there is a great demand for synthetic libraries of chiral molecules that mimic the structural characteristics of privileged scaffolds frequently occurring in natural products and pharmaceuticals. One of these classes of scaffolds are 3,3-disubstituted oxindoles,¹⁶ which are the foundation of many bioactive natural products (Fig. 3). Many new drugs and lead compounds have been created by taking inspiration from these molecules. It is interesting to note that in all fully substituted stereocenters, whether spiro or not, all their carbons are either quaternary or tetrasubstituted with heteroatoms, including the C-3 carbon of oxindoles.^{17,18} The efficient construction of tetrasubstituted, and in particular, quaternary carbon stereocenters remains challenging, which has sparked intense interest in the catalytic enantioselective synthesis of 3,3-disubstituted oxindoles.^{19–22} In addition to providing libraries of structurally different oxindole derivatives for medicinal study, which aid in the discovery of more potent and selective analogues, enormous research effort has also been devoted to the development of new synthetic methodologies.²³

Notably, successful catalytic enantioselective reactions involving direct C–H bond²⁴ functionalization or highly stereoselective construction of adjacent all-carbon quaternary stereocenters are well documented.²⁵ A comprehensive overview on the development of the rapidly evolving field of catalytic enantioselective synthesis of 3,3-disubstituted oxindoles was reported in 2010.^{26,27} Meanwhile, it was noted that, although

elegant protocols are available, it is still highly desirable to create effective techniques for extracting oxindoles with a wide range of structural compositions from readily available starting materials. To achieve the catalytic²⁸ enantioselective synthesis of 3,3-disubstituted oxindoles, some programmes employ new chiral catalysts, new activation models, and tandem sequences.²⁹

Aspergilline A (**9**) was isolated in 2014 from the fungus *Aspergillus versicolor* by Hu and Gao.³⁰ Cladoquinazoline (**10**) is an active obtained from a mangrove-derived fungal strain known as *Cladosporium* sp. and was chosen for additional study as part of anti-influenza compounds. The chemical analysis of the EtOAc extracts of the mycelia of the fungus and its fermentation broth led to the discovery of new indole alkaloids, including cladoquinazoline and other indole alkaloids known to contain quinazoline.^{31,32} Alstonisine (**11**) was discovered in 1972 by Elderfield and Gilman, which is the first oxindole alkaloid related to macrolines, in the plant *Alstonia muelleriana* Domin.^{33,34}

Costinone B (**12**) was isolated from the Pakistani herb *Isatis costata* and found to inhibit lipoxygenases and butyl cholinesterases. The *N*-aryl substitution in these compounds is notable, and also the presence of the 3-oxygen substituent in costinone B. The presence of the 3'-oxygen substituents in isatinones is noteworthy.³⁵

Gelsemine (**13**, C₂₀H₂₂N₂O₂) is an indole alkaloid that acts as a paralytic and is extremely toxic, which was discovered in flowering plants of the genus *Gelsemium*, a plant native to the subtropical and tropical Americas and Southeast Asia. Exposure to gelsemine can be fatal. The family Loganiaceae, which includes the subtropical to tropical flowering plant genus *Gelsemium*, contains five species as of 2014, with the species *G. sempervirens* Ait. being more common in the Americas and *G. elegans* Benth. in China and East Asia.^{36,37}

The chemistry of marine bacteria has also been studied, but they represent a largely untapped source of unusual, bioactive chemical components. Prosurugatoxin (**14**) and a few brominated pyrroles are examples of products of marine bacterial metabolism that have been isolated thus far, demonstrating that the few genera under investigation appear to utilize a variety of secondary metabolic pathways.³⁸

Spirobrassinin (**15**) belongs to the class of organic compounds known as indolines including (*S*)-spirobrassinin. Indolines are substances that contain an indole moiety, which is created when a pyrrolidine ring is fused to a benzene ring to develop 2,3-dihydroindole. In different foods, including *Brassica oleracea* var. *botrytis*, *Brassica rapa*, and *Raphanus sativus*, spirobrassinin has been identified but not quantified. Consequently, (*S*)-spirobrassinin may one day serve as a biomarker for consuming these foods.³⁹

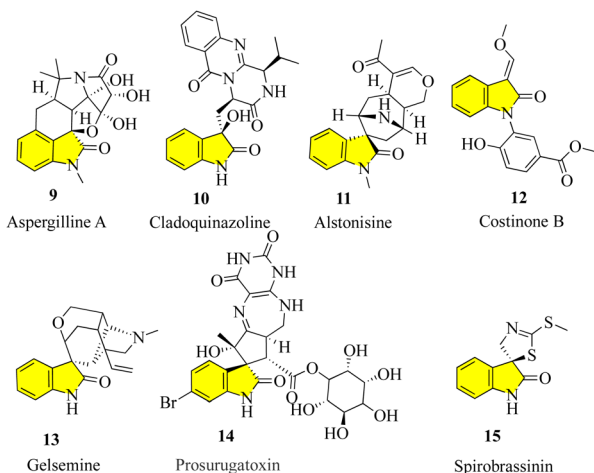


Fig. 3 Structure of natural-based bioactive natural products.

3. Pharmaceuticals containing the 2-oxindole moiety

Despite the large number of 3'-alkyl-oxindoles reported, a Beilstein database search, which was limited to structures from

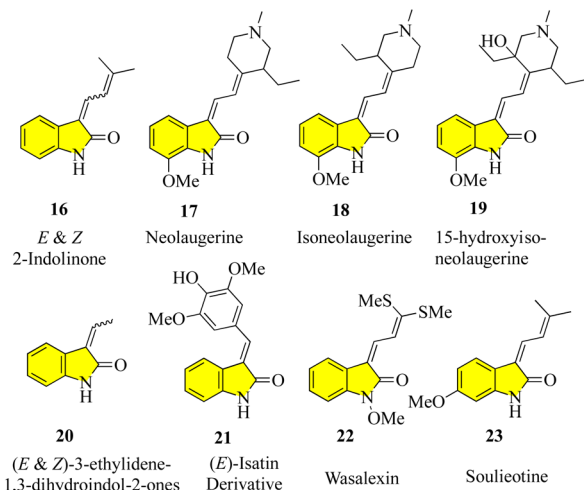


Fig. 4 Natural products containing 3 alkyl-oxindole cores and 3 alkyl-tri-substituents.

natural sources, yielded only 13 compounds with a tri-substituted unit (Fig. 4). The first naturally occurring 3'-alkyloxindoles were (*E*) and (*Z*)-3-(2-methyl-2-butene)-2-indolinones **16** of type 2. In 1978, the two yellow pigments were isolated from the stem of the plant *Cimicifuga dahurica*, which is used in traditional Chinese medicine (Bei Sheng Ma), and especially known for its antipyretic properties. However, it should be noted that the complete description of the structure of these compounds was only reported in 1981 by the same authors. In 1993, three new oxindole alkaloids, *i.e.*, neolaugerine (**17**), isoneolaugerine (**18**), and 15-hydroxyisonolaugerine (**19**) (no stereo morphisms were fully understood), were isolated from the root of the small evergreen tree *Neolaugeria resinosa*, which is distributed in the Bahamas and West Indies. In 1996, very simple (*E*)-3-ethylidene-1,3-dihydroindol-2-ones **20** were

isolated from the *Colletotrichum fragariae* fungus and described as self-growth inhibitors. Oxide alkaloid *E*-isatin derivative **21** was isolated in 1997 from the roots of *Isatis indigotica*, a component of the widely used traditional Chinese medicine Ban-Lan-Gen. In 1999, two plant alkaloids, wasalexins, were isolated from the foliar tissue of Wasabi (*Wasabia japonica*, syn. *Eutrema wasabi*) and wasalexin **22** exhibits antifungal activity against *Phoma lingam*. The functions of the C-3-nitrogen substitute and *N*-methoxy-oxindole are remarkable in these examples. In 2005, it was reported that soulieotine (**23**) was isolated from the roots of *Souliea vaginata*, a plant used as an anti-inflammatory analgesic in traditional Chinese medicine.^{40–47} The simplest examples (without additional rings) are the two new antifungal alkaloid isatinones A (**24**) and B (**25**) isolated from *Isatis costata* (non-stereochemistry defined in the alcohol side chain) in 2007, which expanded the structure search to include naturally occurring 3'-alkenyl-oxide compounds with a 3'-alkenyl unit substituted by tetrasubstituted 3'-alkenyl-oxide compounds, producing the compounds shown in Fig. 5. The presence of a substitute of three oxides in isatinones is noteworthy (see costinone B (**12**)). There are a sufficient number (six) of natural products of 3'-alkenyloxindole, and the 3'-alkenyl substitute is part of an additional ring. The purple pigment violacein (**26**) was first isolated from the Amazon bacteria *Chromobacterium violaceum* 1934, has a variety of biological activities, including *in vitro* antitumor effects. The main component of deoxyviolacein (**27**) was isolated from the same source in 1958, and pseudodeoxyviolacein (**28**) was isolated from *Chromobacterium violaceum* in 1994. Biosynthetic studies of violacein and deoxyviolacin have identified three alkyl-oxindoles related to violacin as possible intermediates. Indirubin (**29**) was first isolated in 1986 from plasma, urine, and hemofiltrate in human blood, and in 2001 from the leaves of *Isatis tinctoria*, its isotopes, isoindirubin (**30**) and isoindigo (**31**), were isolated together.^{48–59}

4. Synthesis of spiro oxindoles & related natural products

4.1 Construction of oxindole-based synthetic products

The atroposelective synthesis of axial chiral molecules has attracted considerable attention from chemists due to the importance of these molecules. However, due to the low rotational barriers and low configuration stability of these molecules, the catalytic asymmetric synthesis of axial chiral styrene or vinyl arene is underdeveloped and difficult. Therefore, the development of powerful strategies for the selective catalytic synthesis of axial cyclic or vinyl arenes is extremely important. In one study, the first selective access to the axially chiral styrene based on oxindole by a catalytic kinetic resolution strategy was developed. This strategy provides two types of oxindole-based axially chiral derivatives of styrene with good diastereoselectivity (94:6dr) and excellent enantioselectivity (98% ee) with high selectivity factors (*S* up 106). This strategy not only facilitates access to the axially chiral styrene based on oxindole, but also provides a robust method for the synthesis of

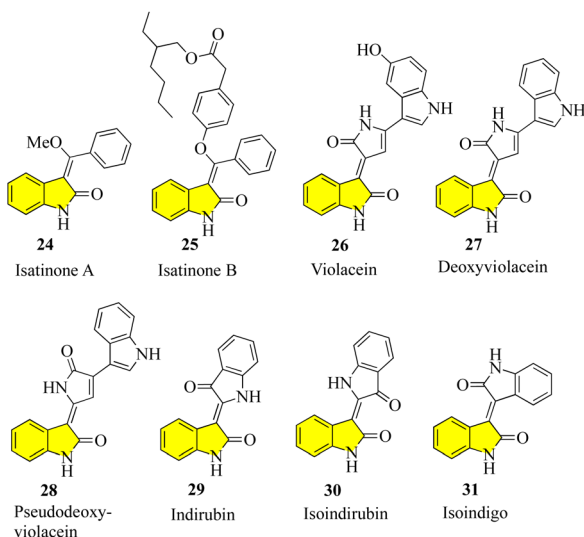
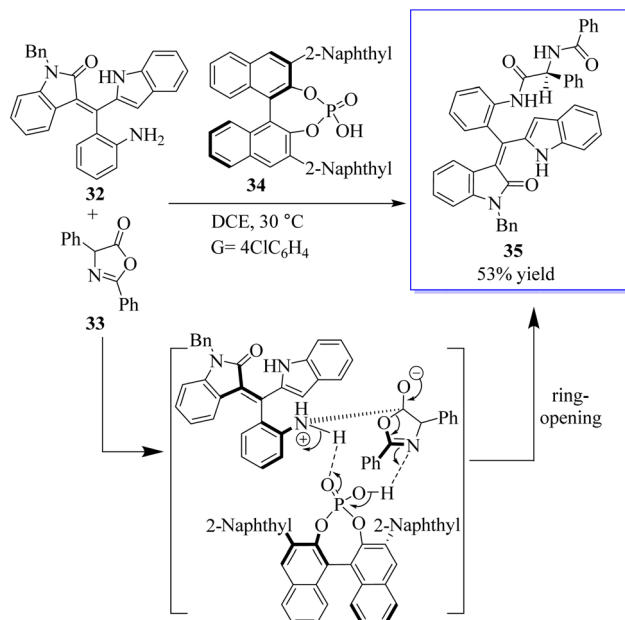


Fig. 5 Natural products with a 3-alkenyl-oxindole core and 3-alkene tetra-substituent.

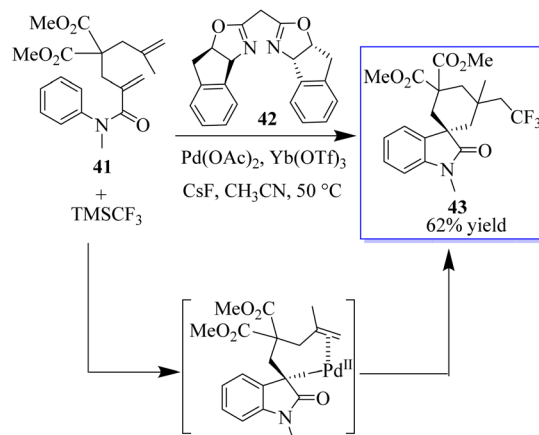




Scheme 1 Construction of the tetracyclic core of lycorine-type alkaloids and its application in the formal synthesis of α -lycorane.

bisamide derivatives **35** with both axial and central chirality. More importantly, this strategy added new members to the axially chiral styrene family (Scheme 1).⁶⁰

The 3,3'-pyrrolidinylspirooxindole scaffold is a privileged structural motif that can be found in a wide range of natural products and pharmaceuticals, which possesses various biological activities, such as antitumor, antidiabetic, anti-inflammatory, and antitubercular activities, among others. Because of these significant bioactivities, 3,3'-pyrrolidinylspirooxindole has emerged as an attractive target, and some elegant



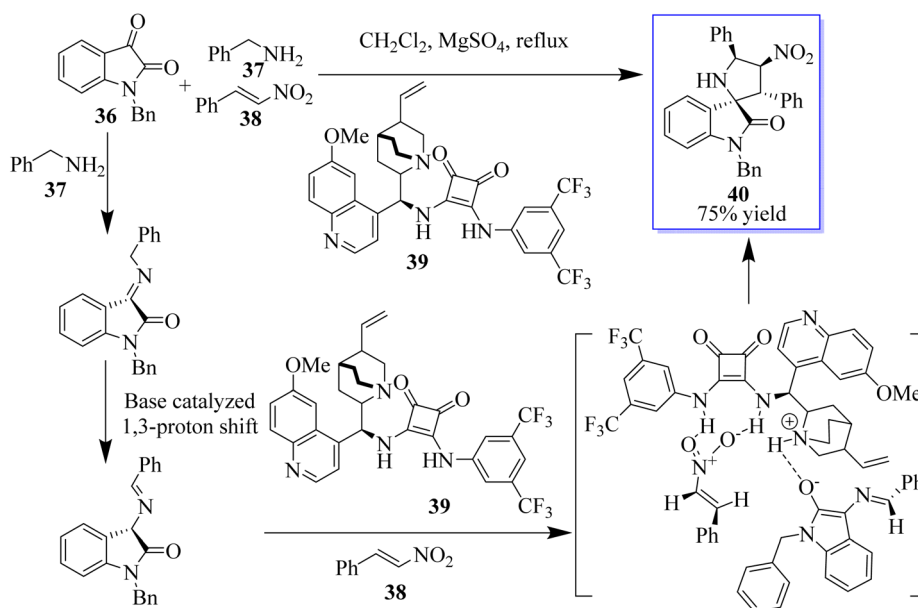
Scheme 3 Pd-catalyzed aryl trifluoromethylation of alkenes.

strategies for its construction have been established. An enantioselective approach to assemble heterocycle **40** involving a three-component reaction of isatins **36**, amines **37**, and nitroalkenes **38** catalyzed by chiral bifunctional squaramide **39** is well documented in this direction (Scheme 2).³⁶

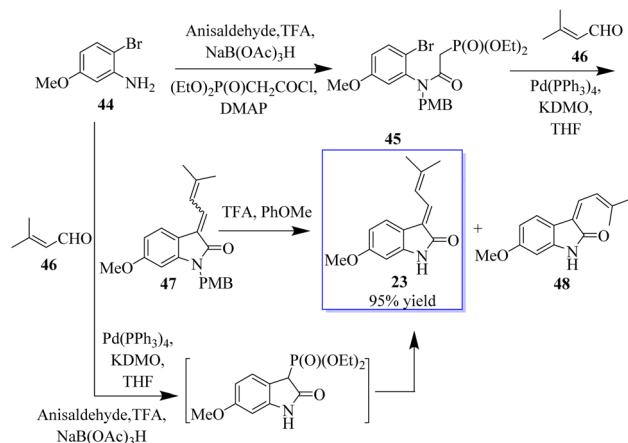
The oxidative aryl trifluoromethylation reaction of activated alkyls catalyzed by palladium was studied in intramolecular and molecular oxidation reactions. The reaction allows the efficient synthesis of various CF_3 -containing oxides **43** in moderate to good yield. The preliminary mechanical studies indicated that the reaction involves an intermediate of $Csp^3Pd^{IV}(CF_3)$, which is reduced to enable the formation of Csp^3CF_3 bonds (Scheme 3).⁶¹

4.2 Construction of the oxindole-based natural products

The enolate arylation/HWE sequence was also used for the first synthesis of the simple natural product soulieotine (**23**)

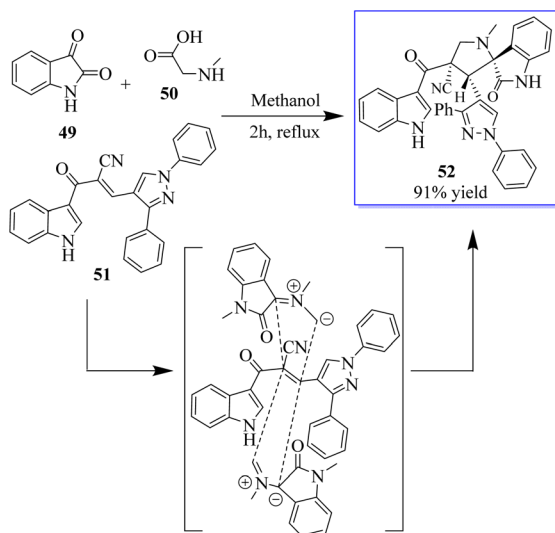


Scheme 2 Catalytic enantioselective 1,3-proton shift/[3 + 2] cycloaddition for the synthesis of spirooxindoles.

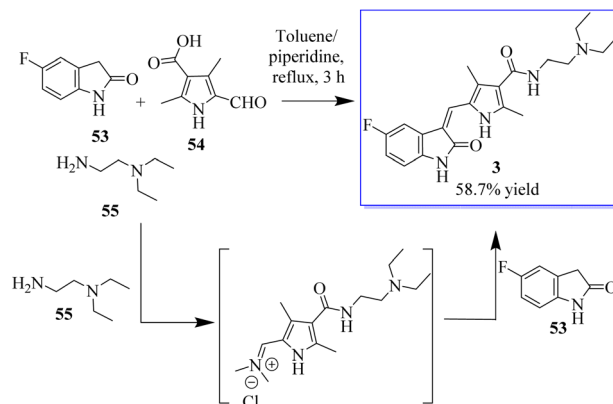
Scheme 4 Synthesis of *E* and *Z*-soulieotine.

(Scheme 4). Soulieotine was isolated from the rhizomes of *Souliea vaginata*, which is used as an anti-inflammatory and analgesic plant in traditional Chinese medicine. The PMB-protected cyclization precursor was easily obtained from aniline and using 3'-methylbutene as a trapping agent gave the expected conductor, with an unoptimized yield as an isomer mixture of alkene. It should be noted that enolate arylation using $\text{Pd}(\text{OAc})_2$ was not successful, probably because of the deprotection of PMB mediated by palladium(II) but proceeded with tetrakis(triphenylphosphine)palladium. After TFA protection, the mixture of *E* and *Z*-soulieotine (2 : 1) was separated by chromatography.⁶²

Under the optimum conditions, a series of novel highly functionalized spiropyrrolidine-oxindoles **52** was synthesized *via* the 1,3-dipolar-cycloaddition of azomethine ylides derived from isatine **49** and various amino acids such as sarcosine, phosphate, and tyroline, and dipolarophile (1,3-diphenyl-1*H*-pyrazol-4-yl)-2-(1*H*-indole-3-carbonyl)acrylonitriles. All synthetic



Scheme 5 Synthesis of spiropyrrolidine-oxindoles by [3 + 2] cycloaddition.



Scheme 6 Synthesis of sunitinib.

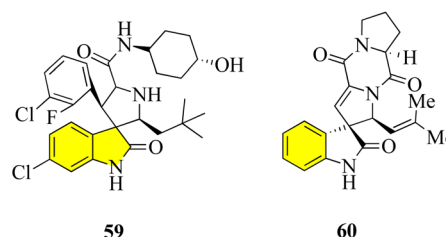
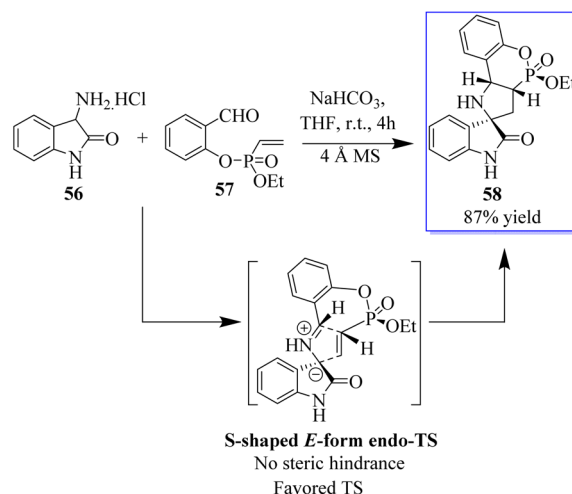


Fig. 6 Bioactive molecules containing spiro-[pyrrolidin-2,3'-oxindole].

compounds have been evaluated for antimicrobial activity and showed significant activity (Scheme 5).⁶³ The process involves converting 5-fluoroindolin-2-one, 5-formyl-2,4-dimethyl-1*H*-pyrrole-3-carboxylic acid and its derivatives, and *N*¹,*N*¹-diethylethane-1,2-diamine to give sunitinib **3**. Sunitinib produced by the above-mentioned process has only 93.87% purity, as shown in Scheme 6.⁶⁴

Pyrrolidin-2,3'-oxindoles **59–60** are representative spirooxindole compounds, which are notable heterocyclic frameworks

Scheme 7 Synthesis of oxindole spiro-*P,N*-polycyclic heterocycles.

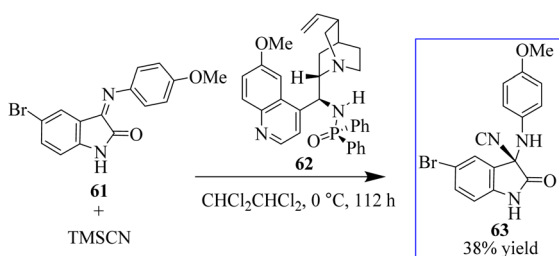
Review

because they are widespread in many natural products and synthetic compounds. They have a wide range of bioactivities such as anticancer, antibacterial, and MDM₂ inhibitory effects (Fig. 6).⁶⁵ A reliable stereoselective assembly strategy for the construction of pyrrolidin-2,3'-oxindole *cis*-fused phosphadihydrocoumarin **58** was established. This process involves the condensation of *O*-vinyl-phosphate salicylic aldehydes and 3'-aminooxindoles, followed by intermolecular cycloaddition with high diastereoselective and atom economy (Scheme 7).⁶⁶

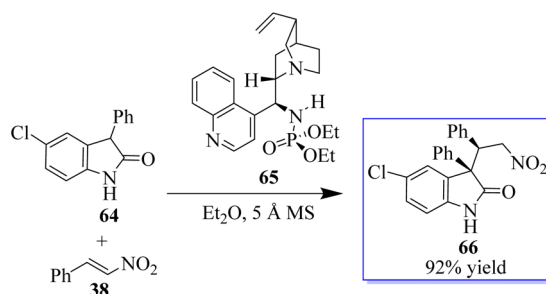
5. Chemical reactivity of 2-oxindole and related derivatives in the presence of chiral catalysts

Quinine-based catalysts **62** were first applied in the Strecker reaction of *N*-aryl isatin ketimines **61** with trimethylsilyl cyanide (TMSCN) (Scheme 8) because at that time, the bifunctional tertiary-amine catalyzed Strecker reaction was undeveloped,⁶⁷ although Deng had reported a highly enantioselective chiral tertiary-amine mediated ketone cyanosilylation to produce **63**.⁶⁸ Preliminary studies indicated that 10 mol% phosphinamide catalyst afforded oxindole based on an aminonitrile.^{69,70} Despite the unsatisfactory result, the phosphinamide catalyst afforded clearly better enantioselectivity than analogous catalysts with amide or thiourea as the H-bond donor.

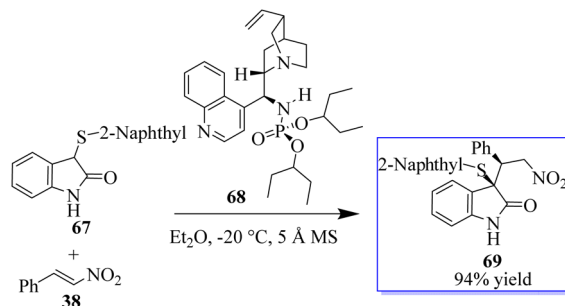
This result exhibited their potency in other reactions and triggered subsequent research on the catalytic enantioselective addition of nucleophiles to isatin ketimines for the synthesis of chiral 3-substituted aminooxindoles. The bifunctional phosphoramides were subsequently found to be potent catalysts for the Michael addition of 3-substituted oxindoles **64** to



Scheme 8 Isatin ketimine asymmetric Strecker reaction.



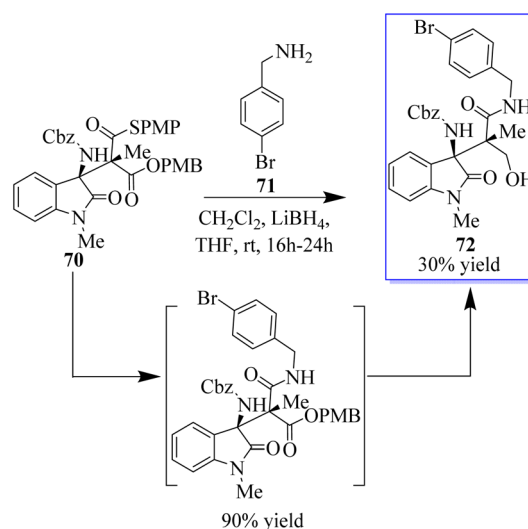
Scheme 9 Asymmetric Michael addition.



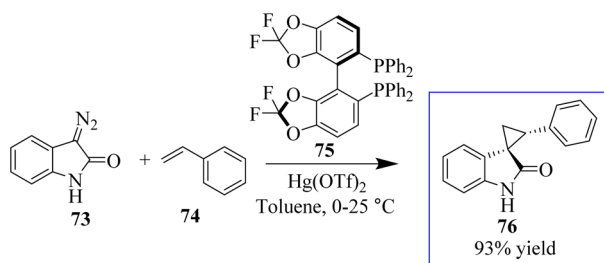
Scheme 10 3-Thioxindole addition by Michael reaction.

nitroolefins **38** (Scheme 9).^{71–73} The resulting adducts are valuable synthons to access quaternary oxindole **66** and indoline derivatives; however, previous studies relied on the use of highly active *N*-protected 3-substituted oxindoles and had limited substrate scope. Unprotected 3-substituted oxindoles were less reactive but more convenient and atom-efficient to prepare. The simple and easily available cinchonidine-derived phosphoramidate is used to achieve high to excellent diastereo- and enantioselectivity. Notably, both 3-aryl- and 3-alkyloxindoles as well as aryl- and alkyl-substituted nitroolefins are viable substrates, giving the desired quaternary oxindoles with excellent enantioselectivity. Later, with phosphoramidate having a bulky ester group, the highly enantioselective Michael addition of 3-alkylthio- and 3-arylthiooxindoles was developed, giving various 3-substituted 3-thiooxindoles **69** in high yield (Scheme 10). This reaction could be run on a gram scale with only 1.0 mol% catalyst **68**.^{74a}

The oxides with adjacent tetrasubstituted stereocenters were obtained in moderate yields and stereoselectivities by the monothiomalonate (MTM) monocatalyzed conjugate addition reaction of *N*-Cbz ketimines derived from isatin. This method requires 2 mol% catalyst load and operates under moderate reaction conditions. Both enantiomers can be used with



Scheme 11 Functionalization of the thioester and oxo-ester moieties.

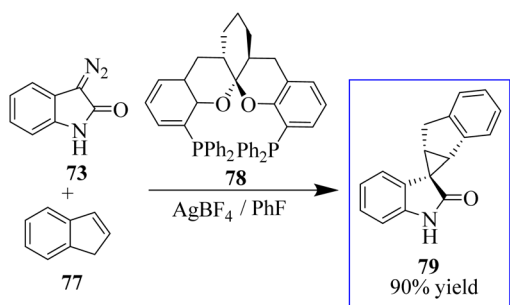


Scheme 12 Asymmetric olefin cyclopropanation reaction catalyzed by $\text{Hg}(\text{II})$.

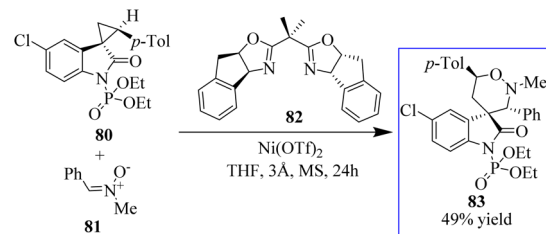
Takemoto's catalyst or an alkaloid derivative of cinchona, with good yields and excellent stereoselectivities. The synthetic methodology allowed a direct route to the derivatives of the antagonist of the agonist of the gene/cholecystokinin-B receptor AG-041R. The reactivity of the connected functional groups can be decreased by molecular crowding at nearby carbon atoms that are fully substituted. In this reaction, the authors treated ent-70 with 4-bromobenzylamine 71, which solely interacted with the thioester group to afford the amide (intermediate), to test if the differentially accessible oxo and thioester moieties could be further functionalized. Alcohol 72 was produced by further selective reduction of the oxoester moiety, which crystallised, allowing the unambiguous assignment of the absolute and relative stereochemistry of the addition products (Scheme 11).^{74b}

The first known instance of diazo compound-mediated $\text{Hg}(\text{II})$ -catalysed olefin cyclopropanation was revealed. Spirocyclopropyl oxindoles can act as donor-acceptor cyclopropanes for complexity-generating synthesis,⁷⁵ in addition to being useful pharmacophores. In this reaction, enantioselective cyclopropanation to obtain 76 was revealed using chiral difluorophos 75 in conjunction with $\text{Hg}(\text{OTf})_2$ (Scheme 12).⁷⁶

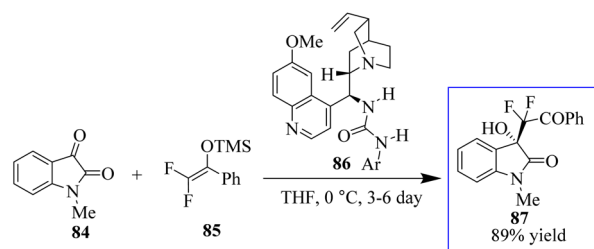
To produce cyclopropanes with high enantioselectivity, both unprotected and *N*-methyl diazooxindoles performed well. Ligand acceleration effects were observed because using 0.4 equivalent of chiral ligand in comparison to $\text{Hg}(\text{OTf})_2$ produced comparable results. Furthermore, changing the counter anion improved the catalytic properties, where difluorophos/ $\text{Hg}(\text{OTf})_2$ failed in these reactions, whereas difluorophos/ $\text{Hg}(\text{PF}_6)_2$ achieved high activity, albeit with moderate enantioselectivity, in



Scheme 13 Cyclopropanation of olefins catalyzed by $\text{Au}(\text{I})$.



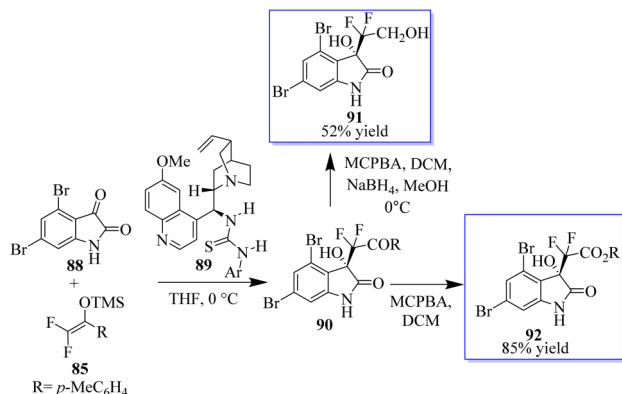
Scheme 14 [3 + 3] cycloaddition of nitrones in an enantioselective manner.



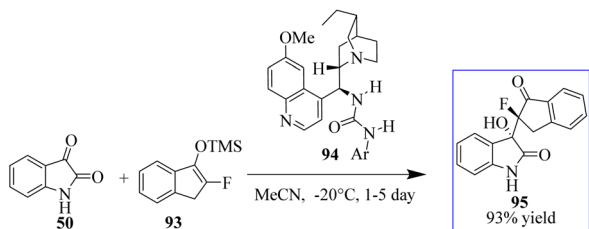
Scheme 15 Reaction of difluoroenoxyisilanes via Mukaiyama-aldol reaction.

the cyclopropanation of disubstituted olefins. These findings demonstrated that ligands could be used to modify the catalytic properties of mercury dioxide ($\text{Hg}(\text{II})$). The cyclopropanation of di- and trisubstituted olefins was then performed with high enantioselectivity using an $\text{Au}(\text{I})$ catalyst. In fact, it is challenging to develop a general catalyst for the full control of stereoselectivity in the cyclopropanation of *trans*- or *cis*-1,2-disubstituted and trisubstituted alkenes due to the high sensitivity of metallocarbenes to the steric hindrance and geometry of alkenes. Potent catalyst ((*S,S*)-SKP L1) 78 enabled highly stereoselective cyclopropanation with a variety of alkenes, including monosubstituted, *cis*- and *trans*-1,2-disubstituted, 1,1-disubstituted, and even trisubstituted alkenes, which is Ding's spiroketal bisphosphine-derived complex (Scheme 13).⁷⁷

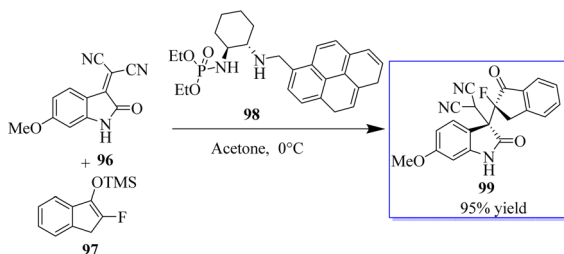
Adjusting the electron-withdrawing *N*-protecting group, the synthesis of spirocyclopropyl oxindoles was revealed. The activation by an appropriate Lewis acid produced *N*-diethoxyphosphoryl oxide 80, which can be converted into spirocyclic oxindoles 83 and 3,5-disubstituted pyrrolidinone by [3 + 3] cycloaddition with nitrone 81, cyclization with 1,4-dithiane-2,5-diol, and ring opening/cyclization with primary amine. Under $\text{Cu}(\text{OTf})_2$ catalysis, the *N*-benzoyl oxindole performed better in the [3 + 2] cycloaddition with aldehyde (Scheme 14). The absence of these transformations in the presence of unprotected or *N*-methyl spirocyclopropyl oxindoles proves the unmistakable activation effects of *N*-protecting groups. This activation strategy works well to produce oxindole-based spirocyclic tetrahydro-1,2-oxazine in catalytic enantioselective reactions using spirocyclopropyl oxindoles. Surprisingly, acetophenone-derived ketonitrones are also good substrates and can be used to make spirooxindoles with nearby quaternary and tetrasubstituted carbon stereocenters. Notably, this is the



Scheme 16 Synthesis of convolutamydine difluoro analogue.



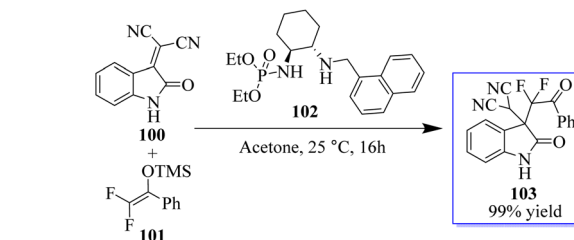
Scheme 17 Mukaiyama aldol reaction of isatins.



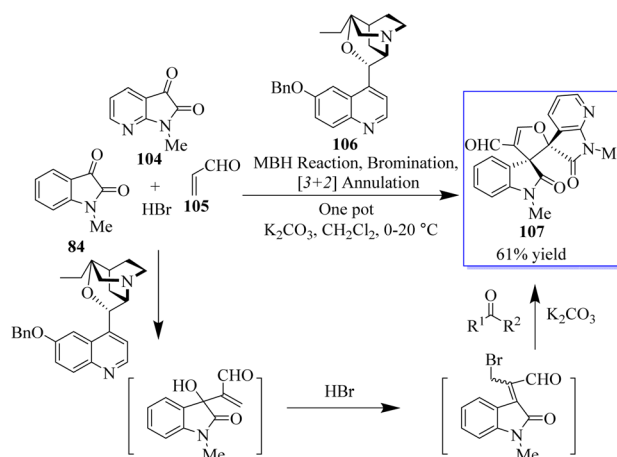
Scheme 18 Mukaiyama–Michael addition reaction.

first method to use the inactivated ketonitrone in enantioselective catalytic synthesis; however, this field has not received much attention.^{78–83}

It is common practice to modify the pharmaceutical properties of organic molecules by selectively adding a fluoroalkyl group. Consequently, oxindoles with a fluoroalkyl group at the C-3 position make intriguing targets for the creation of pharmaceuticals and biological probes. However, although enantioselective trifluoromethylation has been extensively investigated, enantioselective mono- or difluoroalkylation techniques have not.^{84,85} The creation of chiral carbons with an easily accessible mono- or difluoromethylated ketone moiety *via* selective fluoroalkylation⁸⁶ is challenging. It has been revealed that amines can activate fluorinated silyl enol ethers (FSEEs) for enantioselective synthesis with the bifunctional tertiary-amine-catalyzed Strecker reaction using TMSCN. Consequently, the highly enantioselective Mukaiyama-aldol reaction involving isatin **84** and difluoroenoxyisilane **85** was catalyzed by urea



Scheme 19 Implications of phosphoramidate.



Scheme 20 Triple asymmetric sequence.

derived from quinine **86**, producing 3-hydroxyoxindoles **87** (Scheme 15).⁸⁷

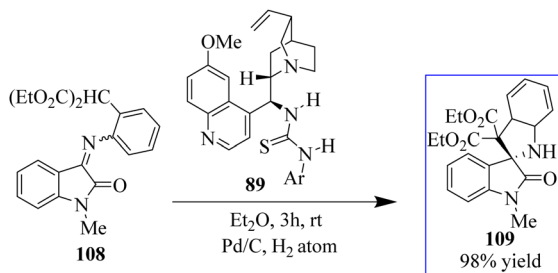
Given that the configuration of convolutamydines is *R*, they are generated by the reaction of isatin **88** with olefin **85** in the presence of quinidine-based catalyst **89**. The difluoro analogue of convolutamydine **E** (**91**) was easily produced in 52% yield after the Baeyer–Villiger oxidation of **90** generated **92** in 85% yield without the loss of ee (Scheme 16).^{19,88–90}

According to Liu, the reaction of isatins **50** and **93** catalyzed by **94** generates a quaternary chiral centre at the C-3 position of oxindole. Even after 3 days, the reaction in THF at 20 °C moved slowly, producing the desired product **95** in 95% yield and with modest stereoselectivities (Scheme 17).⁹¹

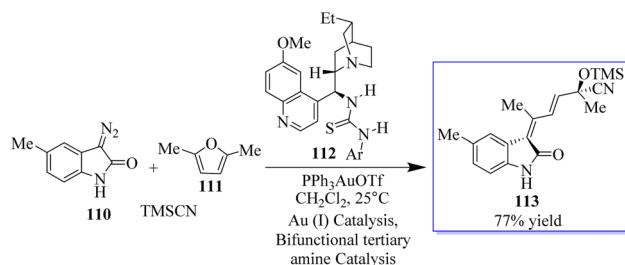
Monofluorinated enol silyl ethers **97** derived from either an α -fluoroindanone or benzofuranone were also effectively activated by the chiral secondary amine phosphoramidate **98** for reaction with isatylidene malononitrile **96** to produce adjacent and fully substituted carbon stereocenters. However, in this instance, **98**, which had a 1-pyrenylmethyl group on the secondary amine moiety, had the best outcome. Monofluorinated oxindole derivatives **99** were produced under these reaction conditions in excellent yield with high to excellent dr and ee values (Scheme 18).^{92–94} Surprisingly, catalyst **102** enabled the reaction between **100** and **101** to be completed in about 16 h at room temperature with acetone as the solvent, producing **103** in 99% yield (Scheme 19).⁹⁴

A novel MBH/bromination/annulation sequence made up of three intermolecular reactions to build on the highly





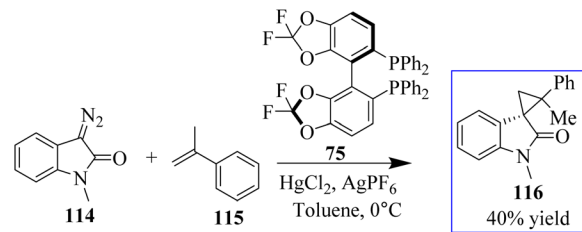
Scheme 21 Catalysis of asymmetric triple sequence.



Scheme 22 Tertiary amine nucleophilic catalysis with Au(I).

enantioselective Morita–Baylis–Hillman (MBH) reaction of isatins **84** and acrolein **105** has been developed. The oxindole-based tetrasubstituted alkenes were produced by the tertiary-amine-catalyzed MBH reaction, and then they underwent a highly stereoselective [3 + 2] annulation with a variety of activated ketones to produce bis(spiro)oxindoles **107** and spirocyclic oxindoles with adjacent quaternary/tetrasubstituted carbon (Scheme 20).^{95–97} Using 10 mol% of bifunctional thiourea **89** produced from quinine in Et₂O at room temperature, the range of the asymmetric 6 π electrocyclization was investigated. The required spirocyclic oxindole derivatives **109** were produced in good yield by a variety of malonate–ketimines **108** with various substituents on the isatin framework (Scheme 21). This process effectively combined the hydrogenation of nitrobenzene by Pd, keta-imine formation by Brønsted acid, and asymmetric electrocyclization by bifunctional tertiary amine in one step. Because the one-pot process avoided racemic cyclization of malonate–anilines during the purification by column chromatography and significant yield losses linked to the purification of malonate–ketimines **108**, the synthetic efficiency significantly increased. Also, the background electrocyclization was suppressed and the bifunctional tertiary-amine-mediated enantioselective reaction was not adversely affected by using only 4 mol% TsOH to promote ketimine formation.^{98,99}

Chiral tertiary amine **112** serves as a Brønsted base to deprotonate and activate the nucleophilic reaction partner of the Michael addition or amination reaction, while the Au-catalyzed reaction provides the 3-substituted oxindole as the nucleophile. As an alternative, the synthesis of chiral 3-alkenyloxindoles **113** from diazo-oxindoles **110**, disubstituted furans **111**, and TMSCN was highly enantioselective due to the integration of Au-catalyzed enone formation and tertiary-amine-mediated cyanosilylation of ketones (Scheme 22).¹⁰⁰

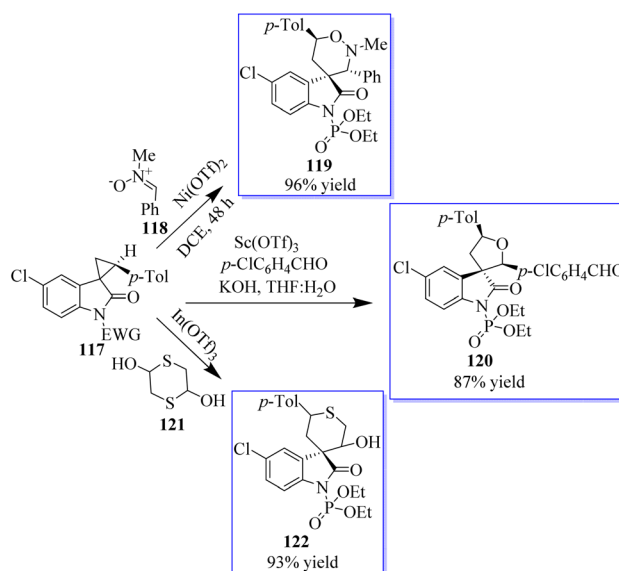


Scheme 23 Cyclopropanation of diazooxindole and styrene.

Diazooxindoles are multifunctional cyclic diazo reagents for the diversity-oriented synthesis (DOS) of substituted oxindoles through reagent-controlled catalytic diversification, including insertion, cyclopropanation, and cycloaddition reactions. Diazooxindoles **114** are easily produced from isatins on a large scale. When the counteranion was changed from OTf to PF₆, the less reactive disubstituted alkenes were also suitable substrates for this reaction. Consequently, when 5.0 mol% of *in situ*-created catalyst **75**/Hg(PF₆)₂ was utilised, *R*-methylstyrene **115** produced **116** in moderate yield, with two neighbouring quaternary stereogenic carbon centres (Scheme 23).⁷⁶

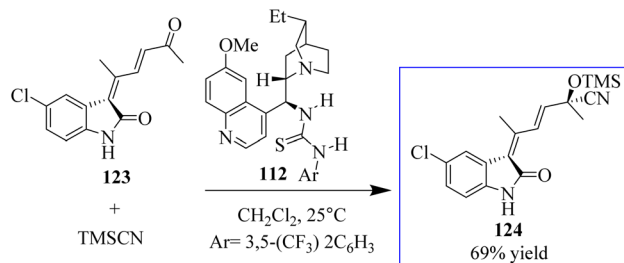
6. Miscellaneous reactivities of 2-oxindole and related derivatives

Although protocols were restricted to substituted cyclopropane-1,1-dicarboxylates or diketones, the cycloaddition of doubly activated cyclopropanes is a successful method for the enantioselective synthesis of cyclic compounds. A particular variety of mono-activated cyclopropane is spirocyclopropyl oxindoles. The low reactivity of these substances prevents the development of new reactions and the associated catalytic enantioselective studies, whereas Carreira invented and demonstrated the value



Scheme 24 Spirocyclopropyl oxindoles-based DOS (diversity-oriented synthesis).





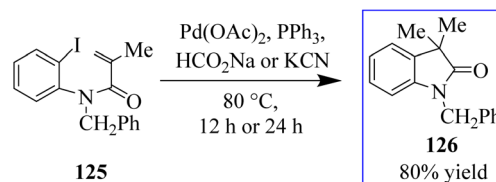
Scheme 25 Catalysis of tertiary amine by asymmetric Au(I).

of their cycloaddition with imines. A method for activating spirocyclopropyl oxindoles **117** using an *N*-protecting group that draws electrons was described. This modification allows for the bidentate coordination of oxindoles to chiral metal complexes for improved enantiofacial control and effectively stabilizes the negative charge developed at C-3 of an oxindole through charge separation upon the activation of a Lewis acid (Scheme 24).^{78,81,82} Due to the superiority of cationic Au(I) catalysis in the olefin cyclopropanation of diazooxindoles, the advantage is the previously unrecognized sequential Au(I)/chiral tertiary amine catalysis as an alluring method to develop diversity-oriented asymmetric tandem reactions, enabling the quick creation of scaffold diversity from diazooxindoles. The success of these tandem reactions, brought about by the high activity of cationic Au(I) catalysis, allowed the use of only 1.0 mol% gold complex to realize these transformations of diazooxindoles, although cationic Au(I) catalysis is known to be incompatible with tertiary amine catalysts. The performance of chiral tertiary amines, which are used at a 10 mol% concentration, is not significantly impacted by the remaining gold catalyst.

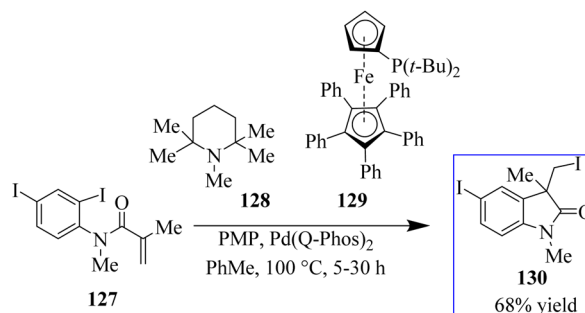
The successful synthesis of multifunctional enone **123** paved the door for the one-pot synthesis of chiral 3-alkenyloxindoles by combining gold-catalysed enone production with catalytic enantioselective addition of a nucleophile to the carbonyl group of **123**. Based on our findings in the Strecker reaction employing TMSCN catalysed by bifunctional tertiary amines, the authors initially attempted the asymmetric cyanosilylation of enone. Fortunately, bifunctional quinine-derived squaramide or (thio) urea **112** could mediate this reaction, and up to 69% of the desired product **124** was produced when **112** was utilised but the reaction progressed slowly even with a 20% catalyst at 25 °C (Scheme 25).¹⁰⁰

7. Miscellaneous synthetic approaches for 2-oxindole and related heterocycles

The palladium-catalyzed Mizoroki–Heck reaction has been a staple in the arsenal of organic chemists ever since it was discovered in the 1970s. A C–X electrophile is added oxidatively in this reaction, and followed by 1,2-migratory insertion. Base-mediated HX reductive elimination regenerates the Pd(0) catalyst, while a subsequent β-H elimination produces the unsaturated cross-coupled product. The suppression of β-H



Scheme 26 Pd-catalyzed carboiodination reaction.



Scheme 27 Polyiodinated compounds employed in the carboiodination reaction.

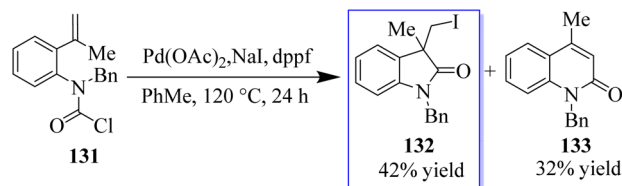
elimination has historically been accomplished in several ways, including electronic and steric bias of the catalyst or the production of a neopentyl organometallic species devoid of β-hydrogens. By using a tethered disubstituted olefin in an intramolecular 6-*endo*-trigtype cyclization of **125**, the Grigg group developed a method to disrupt the typical Mizoroki–Heck-type mechanism in the early 1990s (Scheme 26).^{101–104}

In a migratory insertion, a neopentyl species is formed, which is incapable of undergoing β-H elimination in the presence of Pd(Q-Phos)₂ **129** and 1,2,2,6,6-pentamethylpiperidine **128** in toluene at 100 °C. Under these conditions, **130** was isolated in 68% yield. With the optimized reaction conditions in hand, the authors examined a series of diiodinated compounds **127**. By extending these findings to a one-pot multistep reaction where both the carboiodination reaction and a conventional palladium-catalyzed Mizoroki–Heck reaction on the other halogen moiety were carried out, it showed that these scaffolds could be used as an “oxindole linchpin” molecule in organic synthesis (Scheme 27). Product **130** from both the intramolecular carboiodination reaction and the intermolecular Mizoroki–Heck reaction were identified by *in situ* NMR analysis of these reactions (precyclization).¹⁰⁰ According to these findings, the catalyst may be able to reversibly add to each carbon–iodine bond, enabling the two reactions to take place simultaneously with perfect selectivity.¹⁰⁵

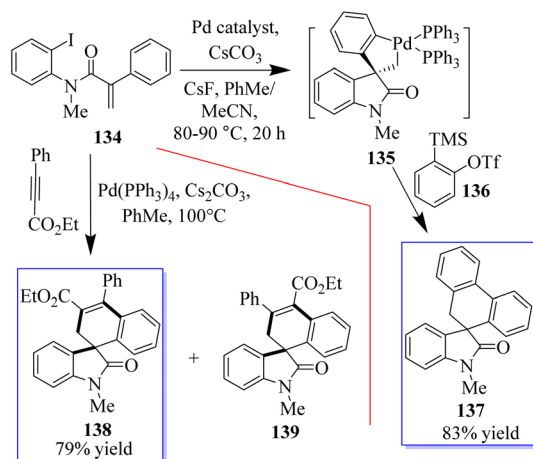
Fortunately, five members of benzo-fused lactam **132** were also easily isolated in 42% yield when compound **131** was used as a substrate in these reaction conditions. In addition to the carboiodination product **132**, Heck reaction product **133** was obtained with a yield of 32% probably by the 6-*endo* cyclization method (Scheme 28).¹⁰⁶

When substrate **134** reacts with the precursor of benzene **136** in MeCN 90 °C with CsF and a Pd(OAc)₂/PPh₃ (OAc) catalyst,





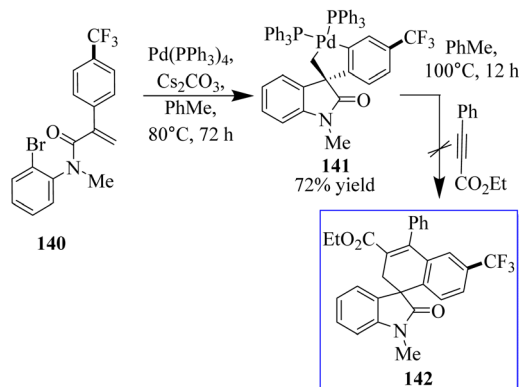
Scheme 28 Enantioselective carbamoyl iodination reaction catalyzed by nickel.



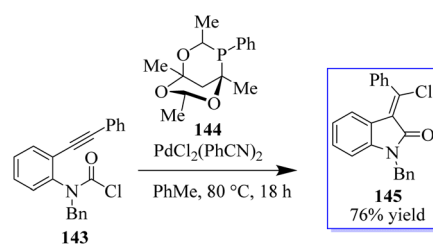
Scheme 29 Synthesis of Pd-catalyzed spirocyclic oxindole.

a small proportion of the expected product **137** was formed. Subjecting **134** to these reaction conditions gave trace amounts of **138** and **139**. The spirocyclic structure **138** was confirmed by spectroscopy and X-ray crystallography of single crystals. Taking advantage of this initial result, the reaction conditions were optimized to produce **138** in a yield of 79% and minimize the formation of side products (Scheme 29).^{107–109} With the help of an *o*-haloacrylamide substrate, it was possible to successfully isolate and characterize a spirocyclic oxindole palladacycle. In-depth research was done by the Garcia-Lopez group to clarify how these reactions work (Scheme 30). Palladacycle was synthesized from a halo acrylamide substrate and examined using single-crystal X-ray crystallography. The important five-membered palladacycle was formed after the σ -alkylpalladium complex was treated under C–H metalation conditions. Interestingly, spirocyclic oxindole **142** was produced after the alkyl-palladium complex was treated with *in situ*-produced benzyne. This provides evidence for both mechanisms. Inconclusive attempts were made to isolate any intermediates produced when benzyne was inserted into five-membered palladacycle **141**. Only the starting material and the final product were visible during NMR (nuclear magnetic resonance) reaction monitoring, indicating that the organometallic intermediates produced by benzyne insertion quickly broke down.^{108,109}

Using a PdCl_2L_2 catalyst, the chlorination of an alkyne can be achieved *via* chloropalladation reaction. The concurrent cyclization of carbamoyl chloride **143** enabled the highly *Z*-selective production of oxindoles **145** with the help of catalyst PA-Ph **144**



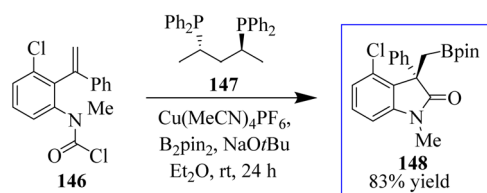
Scheme 30 Synthesis of Pd-catalyzed spirocyclic oxindole: potential mechanisms.



Scheme 31 Chlorocarbonylation reaction of Pd(II–IV).

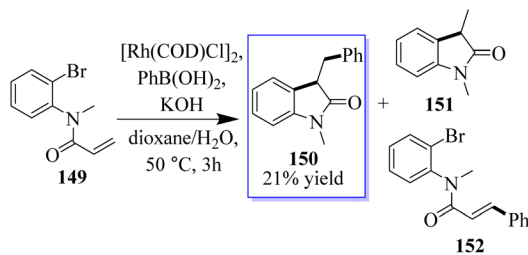
(Scheme 31).¹¹⁰ Subsequently, the oxindole products were produced by cyclization with carbamoyl chloride through a possible palladium(IV) intermediate. The teams of Hoveyda, Ito, and Yun made significant contributions to this field. Using styrenes containing tethered electrophiles, it is planned to apply this technique intramolecularly to quickly access heterocyclic scaffolds. In this procedure, the authors reported the use of carbamoyl chlorides in a copper-catalyzed cyclization method to synthesize enantioenriched substituted borylated oxindoles **148**, continuing our interest in carbamoyl chlorides **146** as electrophiles in metal-catalyzed reactions.

The active bis(diphenylphosphane) **147** catalyst is used in the enantioenriched oxindole synthesis *via* cyclization into the tethered carbamoyl chloride (Scheme 32).¹¹¹ The chemistry made possible by iodo-oxindoles is complemented by the divergent approach of this methodology to oxindole functionalization. Although conventional boronate oxidation produced the oxidized product, Suzuki coupling of the boronate handle produced the arylated oxindole with no erosion of

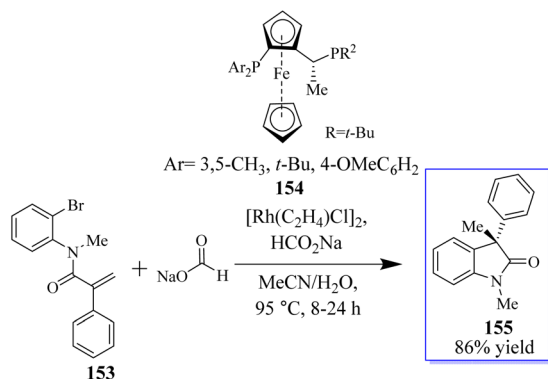


Scheme 32 Borylated oxindole production catalyzed by copper.





Scheme 33 Enolate arylation and domino addition catalyzed by Rh.

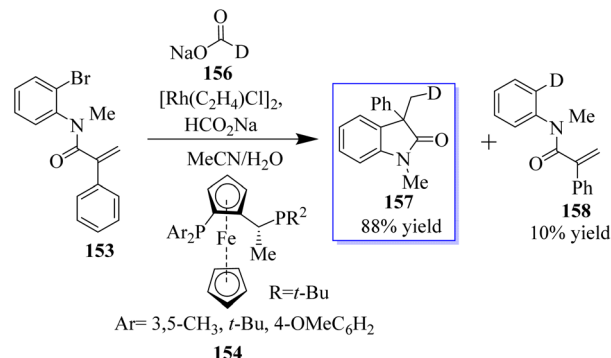


Scheme 34 Reductive arylation induced by enantioselective Rh catalysis.

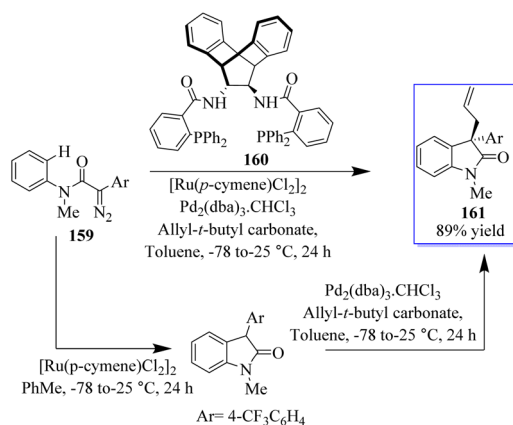
enantioselectivity. It should be noted that using a strong base caused the hydroxyoxindole product to completely lose its stereochemistry. This is probably because a retro-aldol-oxidation sequence took place. The desired oxidation was achieved with no loss of enantioselectivity by using a milder base.

Researchers conducted a study on different α -arylation techniques, and in the 1990s, Hayashi and Miyaura made the initial discovery of rhodium-catalyzed addition to unsaturated systems, which yielded metal enolates in a safe and consistent manner. Amine 149 was treated with $[\text{Rh}(\text{COD})\text{Cl}]$, $[\text{PhB}(\text{OH})_2]$, and KOH at 50°C in dioxane/water solution (10 : 1). After 3 h, 150 was isolated with a yield of 21% of a mixture of 151 and 152, which was eliminated through $\text{Rh}(\text{i})$ -catalyzed Heck-type processes. Attempts to suppress these by-products using substrates such as 152 resulted in complex mixtures (Scheme 33). Although a crucial component of this strategy is the oxidative addition of an $\text{Rh}(\text{i})$ species into a $\text{C}(\text{aryl})\text{X}$ bond, there are fewer examples of this than for palladium and nickel.^{112–119}

The objective to make enantioselective chiral quaternary oxindole using ferrocene chiral ligands 154 was revealed. The investigation into other substitutes and nucleophiles was sparked by this result, specifically the addition of a hydride to acrylamide 153. The 3,3-disubstituted oxindoles 155 were delivered efficiently in up to 86% yield (Scheme 34).¹²⁰ Conversely, 153 was subjected to the sodium deuterate formate and H_2O , making monodeuterated 157 the main product, which confirmed that the hydroxide actually came from sodium formate. In addition, small quantities of multi-deuterium and non-deuterium products were also observed in mass



Scheme 35 Enantioselective Rh-catalyzed reductive arylation.

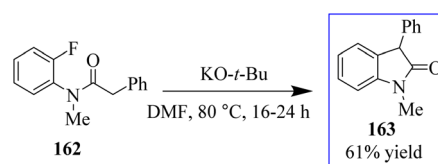


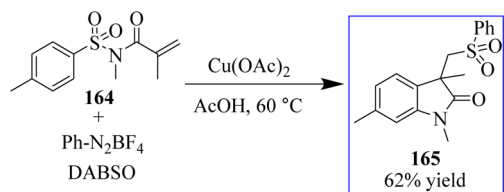
Scheme 36 Enantioenriched 3,3-disubstituted oxindoles through a dual-metal approach.

spectroscopy, possibly suggesting that deuterium incorporation can be a reversible process through the elimination of β -hydrides (Scheme 35).¹²⁰

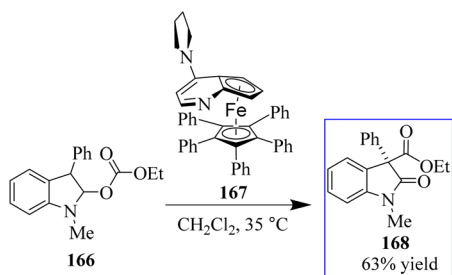
Numerous groups have extensively researched the use of allylic electrophiles such as π -allylpalladium species, with the pioneering work by the Tsuji and Trost group. The authors optimized the transition metal-catalyzed C–H functionalization of α -diazoamide 159 substituting 4(trifluoromethyl)-phenylsuccinate to produce a high yield of enantioenriched oxindoles 161 based on Pd-catalyzed asymmetric allylic alkylation (AAA) using amine-based catalyst 160. In 2016, allyloxindoles were synthesized under the influence of Ru- and Pd-based catalysis (Scheme 36).^{121–124}

An interesting procedure to synthesize substituted oxindoles by intramolecular α -arylation of fluoro- and chloro-substituted

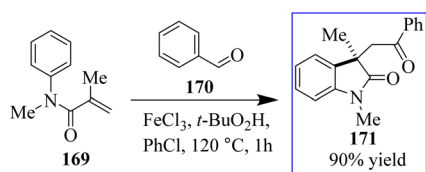
Scheme 37 Synthesis of oxindole derivative by intramolecular α -arylation.



Scheme 38 One-pot mechanism through radical addition reaction.



Scheme 39 Effect of the acyl group on the O-to-C rearrangements.

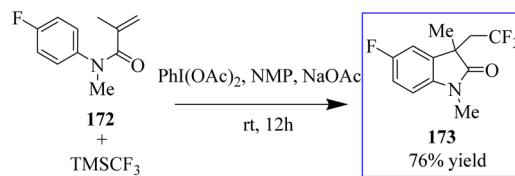
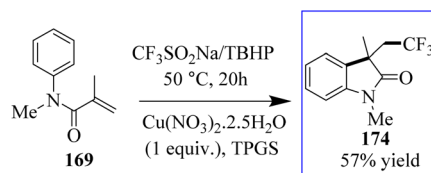


Scheme 40 Oxidative carbonylation of alkenes with aldehydes.

anilides **162** in dimethylformamide at 80 °C, mediated by potassium *tert*-butoxide was revealed. Consequently, product **163** was produced in 61% yield (Scheme 37).¹²⁵

A three-component reaction involving *N*-(arylsulfonyl)acrylamides **164**, DABSO (DABCO-bis(sulfur dioxide)), and phenyldiazonium tetrafluoroborates and Cu(OAc)₂ was used to access sulfonated oxindoles **165**. When aryldiazonium tetrafluoroborates and DABSO react, arylsulfonyl radicals are produced *in situ*, which initiates this transformation. Then, the formation of four new bonds occurs sequentially in one pot through radical addition, radical cyclization, and desulfonylative 1,4-aryl migration to produce the final product. This method exhibits high product yield and strong functional group tolerance (Scheme 38).¹²⁶

The common natural product **168** derived from indole has a quaternary stereocenter at the C-3 position of heterocycle. Catalyst **167** promotes the asymmetric synthesis of these compounds by rearranging *O*-acylated oxindole **166** (Scheme 39).¹²⁷ An effective method for the chiral synthesis of oxindole was established by carbonylation with aldehydes **170** and *N*-arylacrylamide **169**. Three functionalized oxides **171** were smoothly synthesized in high yield with FeCl₃ as the catalyst and tertiary butyl hydro-peroxide as the oxidizer. The obtained oxindoles can be used for further transformation to give various indole alkaloid structure motifs (Scheme 40).¹²⁸

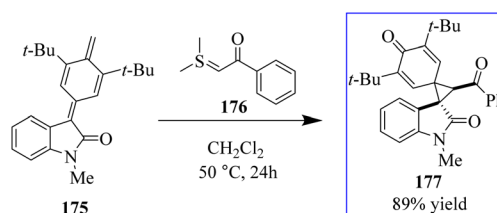
Scheme 41 Main approaches to CF₃-containing oxindoles.

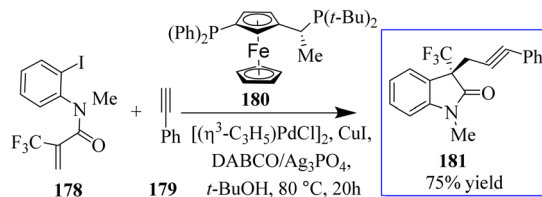
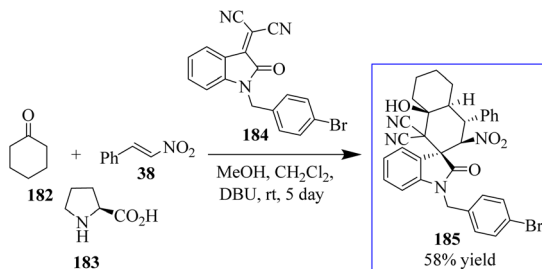
Scheme 42 Transition-metal-catalyzed aryltrifluoromethylation of alkene.

Under metal-free conditions, a light and efficient trifluoromethylation of oxindole **173** was developed using *N*-arylacrylamide **172**. This method is catalyzed by PhI(OAc)₂ and mediated through TMSCF₃. This method provides practical access to a variety of useful CF₃-containing oxides with moderate to good yields (Scheme 41).¹²⁹

The intramolecular trifluoromethylation of *N*-arylacrylamide **169** catalyzed by copper leads to oxide derivatives from the stable and cheap Langlois (CF₃SO₂Na) reagent to give corresponding oxindoles **174**. These reactions occur through a radical process in water at ambient temperature. This method is advantageous in terms of being a green approach (Scheme 42).¹³⁰

A catalyst-free and controllable reaction was developed between *para*-quinone methides derived from isatin **175** and sulfur ylides **176**. This protocol allows the synthesis of different valuable oxindole derivatives **177** in a wide spectrum with high stereoselectivity (Scheme 43).¹³¹ An asymmetric palladium and copper sequential Heck/Sonogashira reaction between *o*-iodoacryl anilides and final alkyls was developed to synthesize chiral oxindoles **181** by the reaction of methyl **178** and *para*-methoxyphenyl (PMB) **179** in the presence of catalyst **180**. A wide range of CF₃-substituted *O*-iodoacryl anilides react with terminal alkynes and provide the corresponding chiral oxindoles with quadruple stereogenic trifluoromethylated centres in high isolated yields and excellent enantioselectivity. This

Scheme 43 Applications of isatin-derived *p*-QMs for the synthesis of oxindole derivatives.

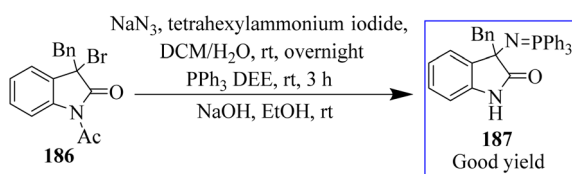
Scheme 44 Enantioselective synthesis of CF₃-substituted oxindoles.

Scheme 45 Synthesis of bioactive spirocyclic oxindoles.

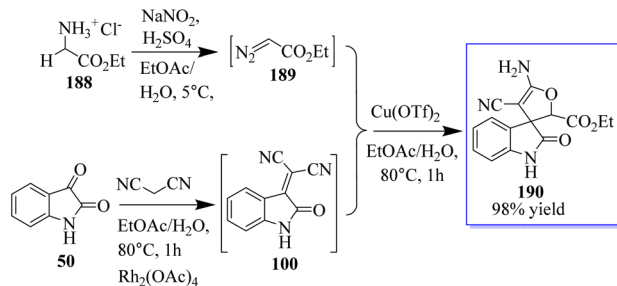
asymmetric Heck/Sonogashira reaction provides a general approach for the insertion of oxindole derivatives containing quaternary stereogenic centers, including those that replace CF₃ (Scheme 44).¹³²

The activation of cyclohexanone by enamine formation for cascade reactions is crucial for synthetic chemists. This concept was revealed using cyclohexanone **182**, nitrostyrene **38**, and L-proline **183**. Subsequently, the first test reaction using substrate **184**, in the presence of the basic organic catalyst DBU and MeOH to promote the domino Michael-aldol reaction was reported. The chiral oxindole having quaternary centre **185** was isolated in mild yield (Scheme 45).¹³³

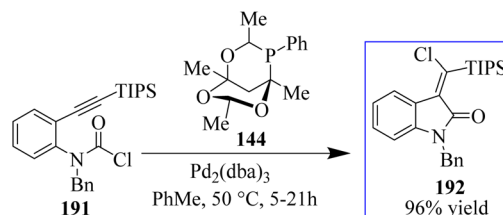
Using 3-alkyloxindole **186**, a derivative of oxindole was synthesized, beginning with the C-3 nitrogen atom. These derivatives were tested in five human tumor cell lines and in healthy donor primary cells (PBMCs), providing compounds with high anticancer effects in low micromolar areas of all cancer cells. The authors reported the green synthesis of iminophosphorane-based oxindole derivatives **187**, which are potentially useful in medicinal chemistry (Scheme 46).¹³⁴ The Rh₂(OAc)₄-catalyzed multi-component reaction of *in situ*-generated ethyl diazoacetate (EDA) **189** and isatylidene malononitrile **100** in the two-phase solvent of water and ethyl acetate using copper triflate was described for the synthesis of chiral oxindole. The Michael-type adducts of hydroxyl oxonium ylide



Scheme 46 Synthesis of 3,3-disubstituted oxindole derivatives.



Scheme 47 Multi-component cascade reactions.

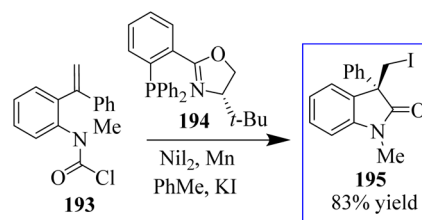


Scheme 48 Carbamoyl chlorination reaction catalyzed by Pd(0).

and malononitrile isatylidene underwent soft inner-molecular ring closure to obtain **190** as the final product with the excellent yield of 98% yield at 50 : 50 diastereoselectivity (Scheme 47).¹³⁵

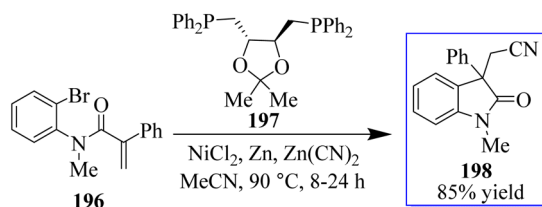
The desired products **192** were only produced in very small amounts because this strategy was incompatible with the synthesis of methylene oxindoles. Using carbamoyl chloride **191** tethered to an alkyne moiety, a different approach that may provide access to the highly desired halogenated methylene oxindole scaffold was discovered. The reaction, which made use of Pd₂(dba)₃ and the large phosphadadamantane ligand **144**, proved to be extremely selective by directing the reductive elimination *trans* to the initial carbopalladation site (Scheme 48).¹³⁶

Due to inability of nickel to form a C(sp³)-Cl bond, an external iodide source had to be added to carry out in-place halogen exchange and enable the reductive elimination of the C(sp³)-I bond. The nickel-catalyzed cross-coupling cyclization reactions were carried out with carbamoyl chlorides **193** as the electrophiles. The asymmetric transfer of a carbamoyl chloride surrogate across a tethered disubstituted styrene using Ni-*t*-BuPHOX catalyst in the presence of Mn(0) and a nucleophilic



Scheme 49 Carbamoyl iodination process catalyzed by enantioselective nickel.





Scheme 50 Reaction of arylcyanation catalyzed by nickel.

source of iodide (KI) produced oxindole derivatives **195** having a chlorine moiety (Scheme 49).¹³⁷

The authors reported that $\text{Zn}(\text{CN})_2$ was used as the source of cyanide in the $\text{NiCl}_2(\text{glyme})\text{DIOP}$ catalytic system, which produced cyanated oxindole products (Scheme 50). By adding activated zinc dust to the reaction mixture, the air-sensitive $\text{Ni}(\text{COD})_2$ was avoided, enabling the use of the manageable (S,S)-DIOP **197** as the catalyst. The “chain walking” reaction, which was popularized by the work of Martin and Marek and others, produced a remotely cyanated oxindole **198** scaffold in one substrate.^{138,139}

8. Conclusions

In this account, we discuss the outcomes of the design, development, and use of synthetic methodologies for the catalytic enantioselective synthesis of 3,3-disubstituted oxindoles. We reported several effective protocols based on different techniques that are categorized by the oxindole synthons used, making it simple to access oxindole-based natural products and synthetic derivatives with a wide range of structural diversity. Also, we reported the synthetic methodologies with a variety of potential bioactive applications. The development of quinine, thioquinine, and phosphoramidate-based chiral bifunctional catalysts or ligands is important given that the success of bifunctional catalysts in the Michael addition, which have some advantages over other H-bond donors. Additionally, the chemical reactivity of 2-oxindole-based derivatives was also described to understand the chemical reactivity pattern of this class of molecules. The use of Mizoroki–Heck-inspired domino cyclization reactions, 1,2-addition–cyclization domino sequences, and MCR strategy involving C–H functionalization can produce and derivatize these valuable structures. Although all these methods build the basic oxindole scaffold, each one has advantages and depends on a wide variety of fundamental reactivity. We anticipate that the techniques and approaches created will serve as models for developing approaches for other biologically significant scaffolds.

Author contributions

Shivangi Sharma: writing original draft, software. Yukti Monga: formal analysis. Ashu Gupta: formal analysis. Shivendra Singh: writing, review and editing.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

The authors are thankful to Prof. (Dr) Kuldeep Singh, HOD Applied Chemistry, Amity University, Gwalior, for his excellent support. We are also thankful to the Department of Applied Chemistry, Amity School of Engineering & Technology (ASET), Gwalior, for all the necessary facilities.

References

- 1 S. Sharma, K. Singh and S. Singh, *Curr. Org. Synth.*, 2022, **26**(15), 1453–1469.
- 2 S. Sharma and S. Singh, *Curr. Org. Chem.*, 2022, **26**, 1453–1469.
- 3 S. Sharma and S. Singh, *Interdiscip. Perspect. Infect. Dis.*, 2023, **2023**, 1–10.
- 4 S. Singh, A. Srivastava and S. Samanta, *Tetrahedron Lett.*, 2012, **53**(45), 6087–6090.
- 5 C. V. Galliford and K. A. Scheidt, *Angew. Chem., Int. Ed.*, 2007, **46**, 8748–8758.
- 6 H. Lin, S. J. Danishefsky, S. J. Danishefsky and H. Lin, *Angew. Chem., Int. Ed.*, 2003, **42**(1), 36–51.
- 7 C. Marti and E. M. Carreira, *Eur. J. Org. Chem.*, 2003, 2209–2219.
- 8 B. M. Trost and M. K. Brennan, *Synthesis*, 2009, **2009**(18), 3003–3025.
- 9 S. Ravi Suman Rudrangi, V. Kumar Bontha, V. Reddy Manda and S. Bethi, *Asian J. Res. Chem.*, 2011, **4**(3), 335–338.
- 10 R. Xu, M. Zhan, L. Peng, X. Pang, J. Yang, T. Zhang, H. Jiang, L. Zhao and Y. Chen, *J. Labelled Compd. Radiopharm.*, 2015, **58**, 308–312.
- 11 R. Roskoski, *Biochem. Biophys. Res. Commun.*, 2007, **356**, 323–328.
- 12 T. H. Yang, C. I. Lee, W. H. Huang and A. R. Lee, *Molecules*, 2017, **22**(6), 913.
- 13 M. Jörg, A. A. Kaczor, F. S. Mak, K. C. K. Lee, A. Poso, N. D. Miller, P. J. Scammells and B. Capuano, *Med. Chem. Commun.*, 2014, **5**, 891–898.
- 14 F. J. Urban, R. Breitenbach and D. Gonyaw, *Synth. Commun.*, 1996, **26**, 1629–1638.
- 15 E. M. de Marigorta, J. M. de Los Santos, A. M. O. de Retana, J. Vicario and F. Palacios, *Beilstein J. Org. Chem.*, 2019, **15**, 1065–1085.
- 16 S. Peddibhotla, *Curr. Bioact. Compd.*, 2009, **5**(1), 20–38.
- 17 X. P. Zeng, Z. Y. Cao, Y. H. Wang, F. Zhou and J. Zhou, *Chem. Rev.*, 2016, **116**, 7330–7396.
- 18 K. Shen, X. Liu, L. Lin and X. Feng, *Chem. Sci.*, 2012, **3**, 327–334.
- 19 Y. Liu, S. J. Han, W. B. Liu and B. M. Stoltz, *Acc. Chem. Res.*, 2015, **48**, 740–751.
- 20 J. S. Yu, H. M. Huang, P. G. Ding, X. S. Hu, F. Zhou and J. Zhou, *ACS Catal.*, 2016, **6**, 5319–5344.
- 21 K. W. Quasdorf and L. E. Overman, *Nature*, 2014, **516**, 181–191.
- 22 J. Zheng, L. Lin, L. Dai, Q. Tang, X. Liu and X. Feng, *Angew. Chem., Int. Ed.*, 2017, **56**, 13107–13111.
- 23 R. Dalpozzo, *Adv. Synth. Catal.*, 2017, **359**, 1772–1810.



- 24 J. R. Frost, S. M. Huber, S. Breitenlechner, C. Bannwarth and T. Bach, *Angew. Chem., Int. Ed.*, 2015, **54**, 691–695.
- 25 K. Ohmatsu, Y. Ando and T. Ooi, *J. Am. Chem. Soc.*, 2013, **135**, 18706–18709.
- 26 R. Dalpozzo, *Org. Chem. Front.*, 2017, **4**, 2063–2078.
- 27 M. Shibasaki and M. Kanai, *Chem. Rev.*, 2008, **108**, 2853–2873.
- 28 J. T. Mohr, M. R. Krout and B. M. Stoltz, *Nature*, 2008, **455**, 323–332.
- 29 D. I. S. P. Resende, P. Boonpothong, E. Sousa, A. Kijjoa and M. M. M. Pinto, *Nat. Prod. Rep.*, 2019, **36**, 7–34.
- 30 M. Zhou, M. M. Miao, G. Du, X. N. Li, S. Z. Shang, W. Zhao, Z. H. Liu, G. Y. Yang, C. T. Che, Q. F. Hu and X. M. Gao, Aspergillines A-E, highly oxygenated hexacyclic indole-tetrahydrofuran-tetramic acid derivatives from *Aspergillus versicolor*, *Org. Lett.*, 2014, **16**(9), 5016–5019.
- 31 J. Peng, T. Lin, W. Wang, Z. Xin, T. Zhu, Q. Gu and D. Li, *J. Nat. Prod.*, 2013, **76**, 1133–1140.
- 32 N. Netz and T. Opatz, *Mar. Drugs*, 2015, **13**, 4814–4914.
- 33 X. Z. Wearing and J. M. Cook, *Org. Lett.*, 2002, **4**, 4237–4240.
- 34 R. C. Elderfield and R. E. Gilman, Alkaloids of *Alstonia Muellieriana*, *Phytochemistry*, 1972, **11**(1), 339–343.
- 35 I. Fatima, I. Ahmad, S. A. Nawaz, A. Malik, N. Afza, G. Lutfullah and M. I. Choudhary, *Het*, 2006, **68**, 1421.
- 36 L. Tian, X. Q. Hu, Y. H. Li and P. F. Xu, *Chem. Commun.*, 2013, **49**, 7213–7215.
- 37 X. Chen, S. Duan, C. Tao, H. Zhai and F. G. Qiu, *Nat. Commun.*, 2016, **7**(1), 10304.
- 38 M. E. Hinze, J. L. Daughtry and C. A. Lewis, *J. Org. Chem.*, 2015, **80**, 11258–11265.
- 39 K. Ryu, S. Nakamura, S. Nakashima and H. Matsuda, *J. Nat. Med.*, 2021, **75**, 308–318.
- 40 K. Hata, K. Baba and M. Kozawa, *Chem. Pharm. Bull.*, 1978, **26**, 2279.
- 41 K. Baba, M. Kozawa, K. Hata, T. Ishida and M. Inoue, *Chem. Pharm. Bull.*, 1981, **29**, 2182.
- 42 B. Weniger, Y. Jiang, R. Anton, J. Bastida, T. Varea and J.-C. Quirion, *Phytochem*, 1993, **32**, 1587.
- 43 M. Inoue, N. Mori, H. Yamanaka, T. Tsurushima, H. Miyagawa and T. Ueno, *J. Chem. Ecol.*, 1996, **22**, 2111.
- 44 X. Wu, Y. Liu, W. Sheng, J. Sun and G. Qin, *Planta Med.*, 1997, **63**, 55.
- 45 M. S. C. Pedras, J. L. Sorensen, F. J. Okanga and I. L. Zaharia, *Bioorg. Med. Chem. Lett.*, 1999, **9**, 3015.
- 46 M. S. C. Pedras, P. B. Chumala and M. Suchy, *Phytochemistry*, 2003, **64**, 949.
- 47 L. Zhou, J. S. Yang, X. Wu, J. H. Zou, X.-D. Xu and G. Z. Tu, *Heter*, 2005, **65**, 1409.
- 48 I. Fatima, I. Ahmad, I. Anis, A. Malik and N. Afza, *Mol*, 2007, **12**, 155.
- 49 R. Wrede, *Hoppe-Seyler's Z. Physiol. Chem.*, 1934, **223**, 113.
- 50 F. M. Strong, *Sci*, 1944, **100**, 287.
- 51 P. S. Melo, G. Z. Justo, M. B. M. de Azevedo, N. Durán and M. Haun, *Toxics*, 2003, **186**, 217.
- 52 J. A. Ballantine, R. J. S. Beer, D. J. Crutchley, G. M. Dodd and D. R. Palmer, *Proc. Chem. Soc., London*, 1958, 232.
- 53 T. Hoshino, T. Hayashi and T. Uchiyama, *Biosci., Biotechnol., Biochem.*, 1994, **58**, 279.
- 54 M. Konzen, D. De Marco, C. A. S. Cordova, T. O. Vieira, R. V. Antônio and T. B. Creczynski-Pasa, *Bioorg. Med. Chem.*, 2006, **14**, 8307.
- 55 S. F. Brady, C. J. Chao, J. Handelsman and J. Clardy, *Org. Lett.*, 2001, **3**, 1981.
- 56 T. Hoshino, T. Hayashi and T. Odajima, *J. Chem. Soc., Perkin Trans. 1*, 1995, 1565.
- 57 H. Laatsch and H. Ludwig-Köhn, *Liebigs Ann. Chem.*, 1986, 1847.
- 58 S. U. Kim, K. S. Song, D. S. Jung, Y. A. Chae and H. J. Lee, *Planta Med.*, 1996, **62**(1), 54–56.
- 59 T. Maugard, E. Enaud, P. Choisy and M. D. Legoy, *Phytochemistry*, 2001, **58**, 897.
- 60 C. Ma, F. T. Sheng, H. Q. Wang, S. Deng, Y. C. Zhang, Y. Jiao, W. Tan and F. J. Shi, *JACS*, 2020, **142**, 15686–15696.
- 61 X. Mu, T. Wu, H. Y. Wang, Y. L. Guo and G. Liu, *J. Am. Chem. Soc.*, 2012, **134**, 878–881.
- 62 A. Millemaggi, A. Perry, A. C. Whitwood and R. J. K. Taylor, *Eur. J. Org. Chem.*, 2009, 2947.
- 63 D. Kathirvelan, J. Haribabu, B. S. Reddy, C. Balachandran and V. Duraipandian, *Bioorg. Med. Chem. Lett.*, 2015, **25**, 389–399.
- 64 G. Meng, C. Liu, S. Qin, M. Dong, X. Wei, M. Zheng, L. Qin, H. Wang, X. He and Z. Zhang, *Res. Chem. Intermed.*, 2015, **41**, 8941–8954.
- 65 A. Gollner, D. Rudolph, H. Arnhof, M. Bauer, S. M. Blake, G. Boehmelt, X. L. Cockroft, G. Dahmann, P. Ettmayer, T. Gerstberger and J. Karolyi-Oezguer, *J. Med. Chem.*, 2016, **59**, 10147–10162.
- 66 T. Huang, L. Liu, Q. Wang, M. Wu and D. Kong, *Synthesis*, 2020, **52**, 1387–1397.
- 67 Y. L. Liu, F. Zhou, J. J. Cao, C. bin Ji, M. Ding and J. Zhou, *Org. Biomol. Chem.*, 2010, **8**, 3847–3850.
- 68 S. K. Tian and L. Deng, *J. Am. Chem. Soc.*, 2001, **123**, 6195–6196.
- 69 Y. L. Liu and J. Zhou, *Synthesis*, 2015, **47**, 1210–1226.
- 70 P. Chauhan and S. S. Chimni, *Tetrahedron: Asymmetry*, 2013, **24**, 343–356.
- 71 F. Zhou, Y. L. Liu and J. Zhou, *Adv. Synth. Catal.*, 2010, **352**, 1381–1407.
- 72 M. Ding, F. Zhou, Y. L. Liu, C. H. Wang, X. L. Zhao and J. Zhou, *Chem. Sci.*, 2011, **2**, 2035–2039.
- 73 M. Ding, F. Zhou, Z. Q. Qian and J. Zhou, *Org. Biomol. Chem.*, 2010, **8**, 2912–2914.
- 74 (a) W. M. Gao, J. S. Yu, Y. L. Zhao, Y. L. Liu, F. Zhou, H. H. Wu and J. Zhou, *Chem. Commun.*, 2014, **50**, 15179–15182; (b) O. D. Engl, S. P. Fritz and H. Wennemers, *Angew. Chem.*, 2015, **127**, 8311–8315.
- 75 Z. Y. Cao and J. Zhou, *Org. Chem. Front.*, 2015, **2**, 849–858.
- 76 Z. Y. Cao, F. Zhou, Y. H. Yu and J. Zhou, *Org. Lett.*, 2012, **15**, 42–45.
- 77 Z. Y. Cao, X. Wang, C. Tan, X. L. Zhao, J. Zhou and K. Ding, *J. Am. Chem. Soc.*, 2013, **135**, 8197–8200.
- 78 P. W. Xu, J. K. Liu, L. Shen, Z. Y. Cao, X. L. Zhao, J. Yan and J. Zhou, *Nat. Commun.*, 2017, **8**(1), 1619.

- 79 W. M. Shi, X. P. Ma, G. F. Su and D. L. Mo, *Org. Chem. Front.*, 2016, **3**, 116–130.
- 80 R. R. Liu, S. C. Ye, C. J. Lu, G. L. Zhuang, J. R. Gao and Y. X. Jia, *Angew. Chem.*, 2015, **127**, 11357–11360.
- 81 H. K. Grover, M. R. Emmett and M. A. Kerr, *Org. Biomol. Chem.*, 2015, **13**, 655–671.
- 82 P. B. Alper, C. Meyers, A. Lerchner, D. R. Siegel and E. M. Carreira, *Angew. Chem., Int. Ed.*, 1999, **38**(21), 3186–3189.
- 83 T. F. Schneider, J. Kaschel and D. B. Werz, *Angew. Chem., Int. Ed.*, 2014, **53**, 5504–5523.
- 84 J. A. Ma and D. Cahard, *Chem. Rev.*, 2004, **104**, 6119–6146.
- 85 C. P. Zhang, Q. Y. Chen, Y. Guo, J. C. Xiao and Y. C. Gu, *Chem. Soc. Rev.*, 2012, **41**, 4536–4559.
- 86 C. Ni and J. Hu, *Chem. Soc. Rev.*, 2016, **45**, 5441–5454.
- 87 Y. L. Liu and J. Zhou, *Chem. Commun.*, 2012, **48**, 1919–1921.
- 88 X. Yang, T. Wu, R. J. Phipps and F. D. Toste, *Chem. Rev.*, 2015, **115**, 826–870.
- 89 Y. L. Liu, J. S. Yu and J. Zhou, *Asian J. Org. Chem.*, 2013, **2**, 194–206.
- 90 M. Decostanzi, J. M. Campagne and E. Leclerc, *Org. Biomol. Chem.*, 2015, **13**, 7351–7380.
- 91 Y. L. Liu, F. M. Liao, Y. F. Niu, X. L. Zhao and J. Zhou, *Org. Chem. Front.*, 2014, **1**, 742–747.
- 92 J. S. Yu and J. Zhou, *Org. Chem. Front.*, 2016, **3**, 298–303.
- 93 J. S. Yu and J. Zhou, *Org. Chem. Front.*, 2016, **3**, 298–303.
- 94 J. S. Yu, F. M. Liao, W. M. Gao, K. Liao, R. L. Zuo and J. Zhou, *Angew. Chem.*, 2015, **127**, 7489–7493.
- 95 J. Zhou, *Chem.-Asian J.*, 2010, **5**, 422–434.
- 96 Y. L. Liu, B. L. Wang, J. J. Cao, L. Chen, Y. X. Zhang, C. Wang and J. Zhou, *J. Am. Chem. Soc.*, 2010, **132**, 15176–15178.
- 97 Y. L. Liu, X. Wang, Y. L. Zhao, F. Zhu, X. P. Zeng, L. Chen, C. H. Wang, X. L. Zhao and J. Zhou, *Angew. Chem.*, 2013, **125**, 13980–13984.
- 98 X. P. Yin, X. P. Zeng, Y. L. Liu, F. M. Liao, J. S. Yu, F. Zhou and J. Zhou, *Angew. Chem.*, 2014, **126**, 13960–13965.
- 99 Y. L. Liu, X. P. Yin and J. Zhou, *Chin. J. Chem.*, 2018, **36**, 321–328.
- 100 Y. L. Zhao, Z. Y. Cao, X. P. Zeng, J. M. Shi, Y. H. Yu and J. Zhou, *Chem. Commun.*, 2016, **52**, 3943–3946.
- 101 R. F. Heck and J. P. Nolley, *J. Org. Chem.*, 1972, **37**(14), 2320–2322.
- 102 R. Grigg and V. Sridharan, *J. Organomet. Chem.*, 1999, **576**(1–2), 65–87.
- 103 R. Grigg, V. Santhakumar and V. Sridharan, *Tetrahedron Lett.*, 1993, **34**(19), 3163–3164.
- 104 B. Burns, R. Grigg, V. Santhakumar, V. Sridharan, P. Stevensoga and T. Worakun, *Tetrahedron*, 1992, **48**(35), 7297–7320.
- 105 D. A. Petrone, M. Lischka and M. Lautens, *Angew. Chem., Int. Ed.*, 2013, **52**, 10635–10638.
- 106 C. Chen, J. Hu, J. Su and X. Tong, *Tetrahedron Lett.*, 2014, **55**, 3229–3231.
- 107 M. Pérez-Gómez and J. A. García-López, *Angew. Chem., Int. Ed.*, 2016, **55**, 14389–14393.
- 108 H. Yoon, A. Lossouarn, F. Landau and M. Lautens, *Org. Lett.*, 2016, **18**, 6324–6327.
- 109 H. Yoon, M. Rölz, F. Landau and M. Lautens, *Angew. Chem., Int. Ed.*, 2017, **56**, 10920–10923.
- 110 C. M. Le, T. Sperger, R. Fu, X. Hou, Y. H. Lim, F. Schoenebeck and M. Lautens, *J. Am. Chem. Soc.*, 2016, **138**, 14441–14448.
- 111 A. Whyte, K. I. Burton, J. Zhang and M. Lautens, *Angew. Chem., Int. Ed.*, 2018, **57**, 13927–13930.
- 112 R. Shintani, T. Yamagami and T. Hayashi, *Org. Lett.*, 2006, **8**, 4799–4801.
- 113 Y. J. Jang, H. Yoon and M. Lautens, *Org. Lett.*, 2015, **17**, 3895–3897.
- 114 Y. Harada, J. Nakanishi, H. Fujihara, M. Tobisu, Y. Fukumoto and N. Chatani, *J. Am. Chem. Soc.*, 2007, **129**, 5766–5771.
- 115 L. Souillart and N. Cramer, *Chem. Sci.*, 2014, **5**, 837–840.
- 116 D. Katayev, Y. X. Jia, A. K. Sharma, D. Banerjee, C. Besnard, R. B. Sunoj and E. P. Kündig, *Chem.-Eur. J.*, 2013, **19**, 11916–11927.
- 117 A. M. Taylor, R. A. Altman and S. L. Buchwald, *J. Am. Chem. Soc.*, 2009, **131**, 9900–9901.
- 118 S. Würtz, C. Lohre, R. Fröhlich, K. Bergander and F. Glorius, *J. Am. Chem. Soc.*, 2009, **131**, 8344–8345.
- 119 E. P. Kündig, T. M. Seidel, Y. X. Jia and G. Bernardinelli, *Angew. Chem., Int. Ed.*, 2007, **46**, 8484–8487.
- 120 Y. J. Jang, E. M. Larin and M. Lautens, *Angew. Chem., Int. Ed.*, 2017, **56**, 11927–11930.
- 121 P. Koschker and B. Breit, *Acc. Chem. Res.*, 2016, **49**, 1524–1536.
- 122 J. Tsuji and T. Susuki, *Tetrahedron Lett.*, 1965, **6**(34), 3027–3031.
- 123 B. M. Trost and M. L. Crawley, *Chem. Rev.*, 2003, **103**, 2921–2943.
- 124 K. Yamamoto, Z. Qureshi, J. Tsoung, G. Pisella and M. Lautens, *Org. Lett.*, 2016, **18**, 4954–4957.
- 125 A. Beyer, J. Buendia and C. Bolm, *Org. Lett.*, 2012, **14**, 3948–3951.
- 126 H. Wang, S. Sun and J. Cheng, *Org. Lett.*, 2017, **19**, 5844–5847.
- 127 I. D. Hills and G. C. Fu, *Angew. Chem., Int. Ed.*, 2003, **42**, 3921–3924.
- 128 F. Jia, K. Liu, H. Xi, S. Lu and Z. Li, *Tetrahedron Lett.*, 2013, **54**, 6337–6340.
- 129 W. Fu, F. Xu, Y. Fu, C. Xu, S. Li and D. Zou, *Eur. J. Org. Chem.*, 2014, **2014**, 709–712.
- 130 F. Yang, P. Klumphu, Y. M. Liang and B. H. Lipshutz, *Chem. Commun.*, 2014, **50**, 936–938.
- 131 Y. You, B. X. Quan, Z. H. Wang, J. Q. Zhao and W. C. Yuan, *Org. Biomol. Chem.*, 2020, **18**, 4560–4565.
- 132 X. Bai, C. Wu, S. Ge and Y. Lu, *Angew. Chem., Int. Ed.*, 2020, **59**, 2764–2768.
- 133 L. Straminelli, F. Vicentini, A. Di Sabato, C. M. Montone, C. Cavaliere, K. Rissanen, F. Leonelli and F. Vetica, *J. Org. Chem.*, 2022, **87**, 10454–10461.
- 134 M. S. Christodoulou, F. Nicoletti, K. Mangano, M. A. Chiacchio, G. Facchetti, I. Rimoldi, E. M. Beccalli and S. Giofrè, *Bioorg. Med. Chem. Lett.*, 2020, **30**, 126845.



- 135 T. Shi, S. Teng, Y. Wei, X. Guo and W. Hu, *Green Chem.*, 2019, **21**, 4936–4940.
- 136 M. L. Christine, M. Le, X. Hou, T. Sperger, F. Schoenebeck and M. Lautens, *Angew. Chem.*, 2015, **54**, 15897–15900.
- 137 A. D. Marchese, M. Wollenburg, B. Mirabi, X. Abel-Snape, A. Whyte, F. Glorius and M. Lautens, *ACS Catal.*, 2020, **10**, 4780–4785.
- 138 A. Yen and M. Lautens, *Org. Lett.*, 2018, **20**, 4323–4327.
- 139 H. Sommer, F. Juliá-Hernández, R. Martin and I. Marek, *ACS Cent. Sci.*, 2018, **4**, 153–165.

