

Cross-Coupling-Reaction of Copper Carbene Intermediate with Terminal Alkyne

Subjects: [Chemistry](#), [Organic](#)

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Copper-catalyzed cross-coupling reaction of a copper carbene intermediate with terminal alkynes was one of the most powerful protocols for the construction of C–C bonds. However, in early works, a mixture of alkynoates and allenates was generated in combined moderate yields under harsh reaction conditions. Until 2004, Fu reported the first example of the copper-catalyzed coupling reaction of terminal alkynes with diazo esters or diazo amides to yield 3-alkynoate or 3-butyneamide products selectively with minimal amount of allene byproducts under no-basic conditions. Consequently, a variety of copper-catalyzed coupling reactions of terminal alkynes with various carbene precursors have been developed independently.

copper catalysis

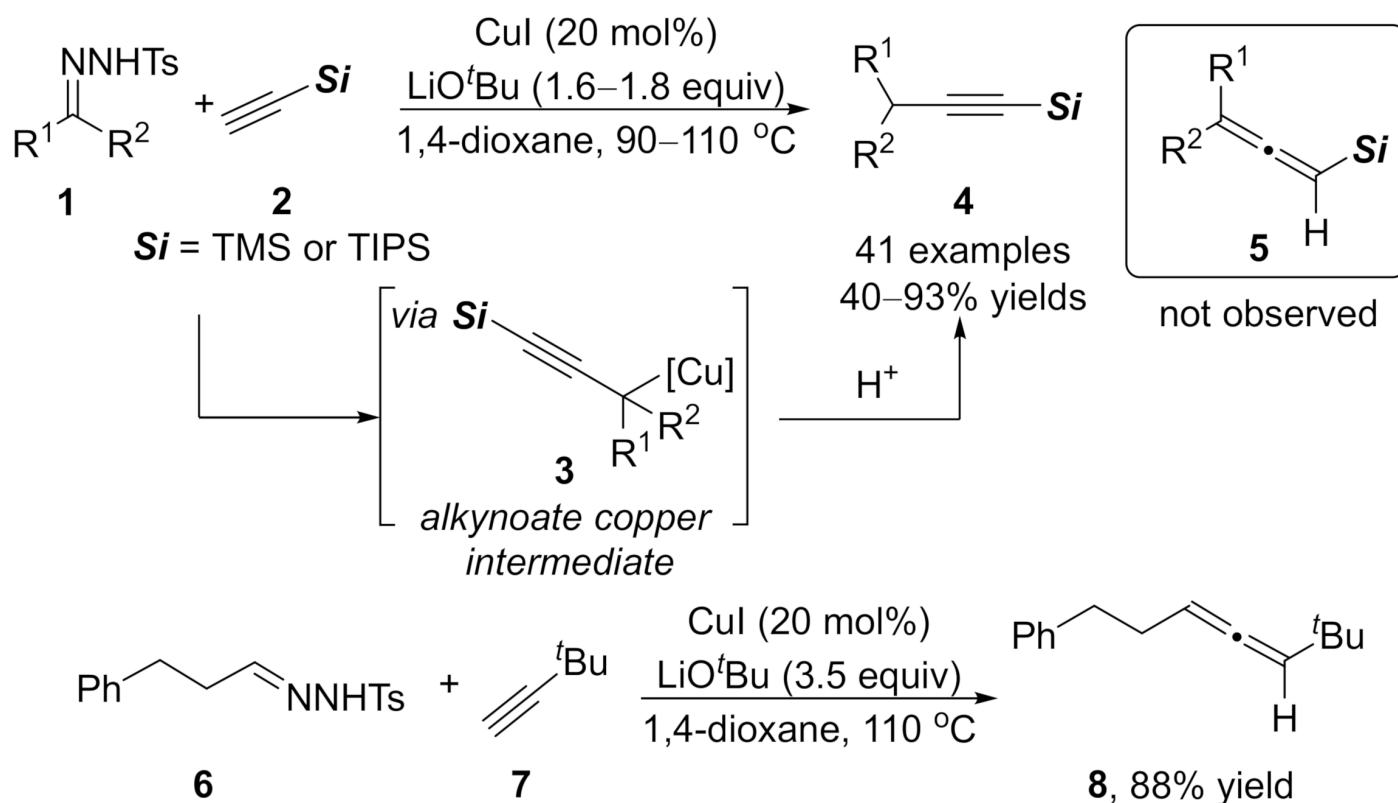
carbene intermediate

alkyne functionalization

1. Alkynylation

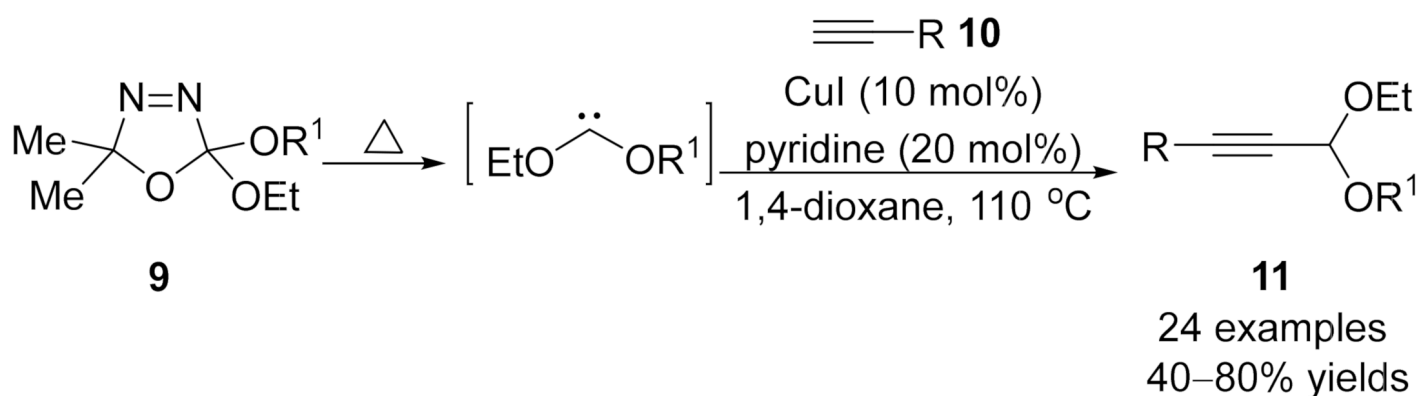
1.1. Alkynylation Terminated by Protonation

In 2012, Wang reported a copper-catalyzed cross-coupling of *N*-tosylhydrazones **1** with trialkylsilylalkynes **2**, leading to the alkynylated products **4** via the formation of C(sp)–C(sp³) bonds ([Scheme 1](#)). Mechanism study shows that migratory insertion of copper carbene species gives the alkynoate copper intermediate **3**, and sequential protonation affords the target products **4**. This coupling reaction proceeded efficiently with *N*-tosylhydrazones derived from aromatic and aliphatic aldehydes or ketones in moderate to excellent yields without detecting the formation of allene byproducts **5**. However, when a *tert*-butyl substituted alkyne **7** was employed, the corresponding allene product **8** was formed selectively [\[1\]](#).



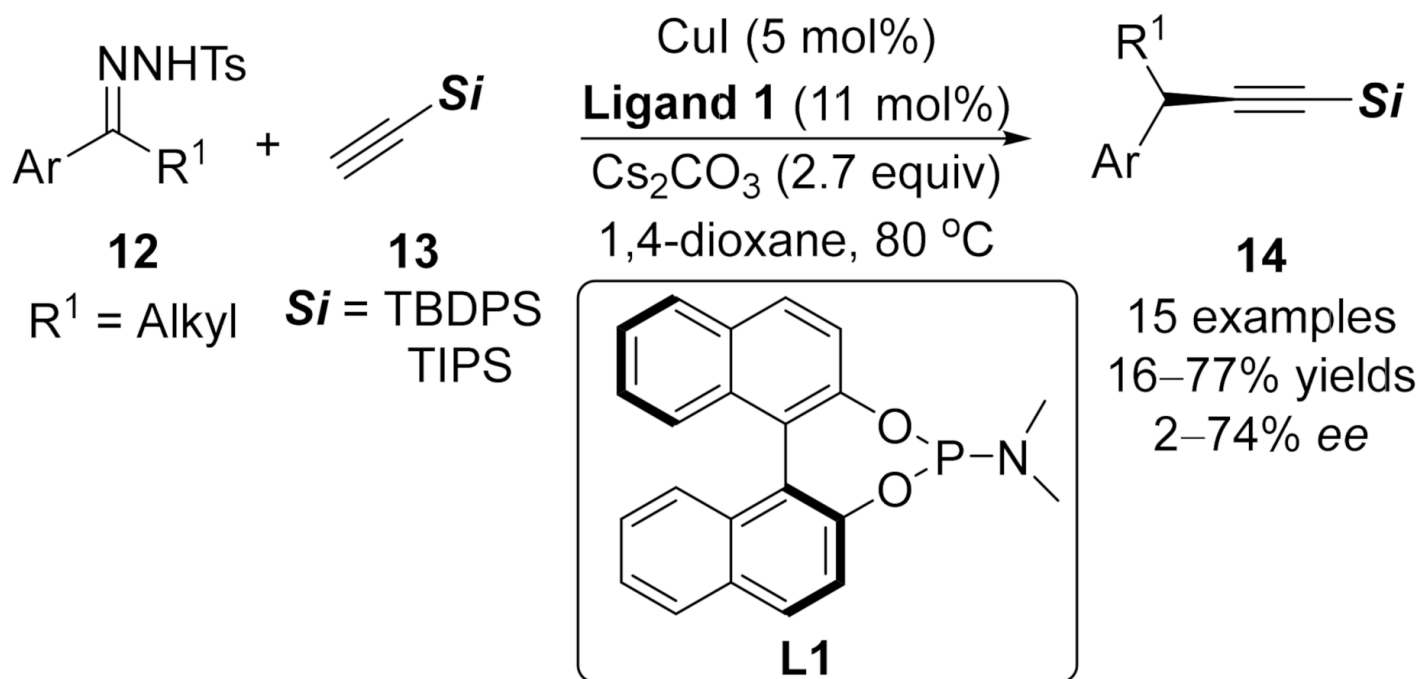
Scheme 1. Cross-coupling reaction of *N*-tosylhydrazones with terminal alkynes.

Later in 2014, Zhou reported a copper-catalyzed coupling reaction using dialkoxycarbenes **9** as carbene precursor [\[2\]](#), which provides unsymmetrical propargylic acetals **11** in moderate to good yields ([Scheme 2](#)).



Scheme 2. Cross-coupling reaction of dialkoxycarbenes with terminal alkynes.

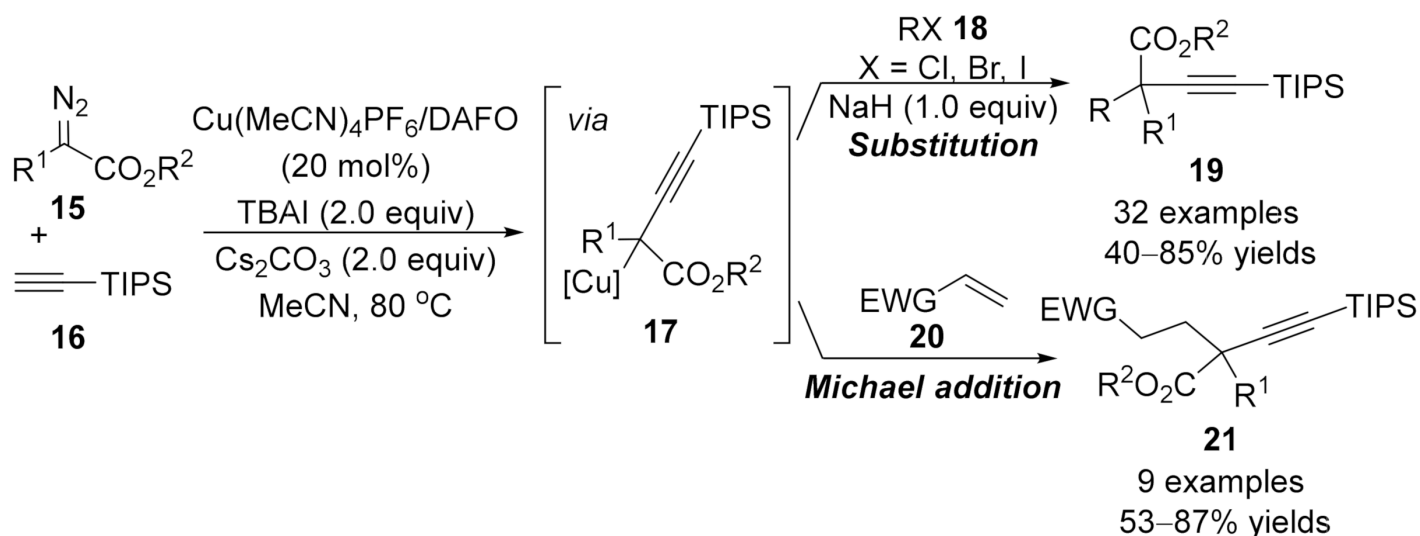
In 2018, an asymmetric coupling reaction of *N*-tosylhydrazones **12** with terminal alkynes **13** was achieved by Uozumi and co-workers using chiral copper(I)/phosphoramidite complex as the chiral catalyst ([Scheme 3](#)), and optically active alkynylated product **14** was generated in moderate to good yields and enantioselectivities [\[3\]](#).



Scheme 3. Chiral Cu(I)/phosphoramidite complex catalyzed asymmetric coupling reaction.

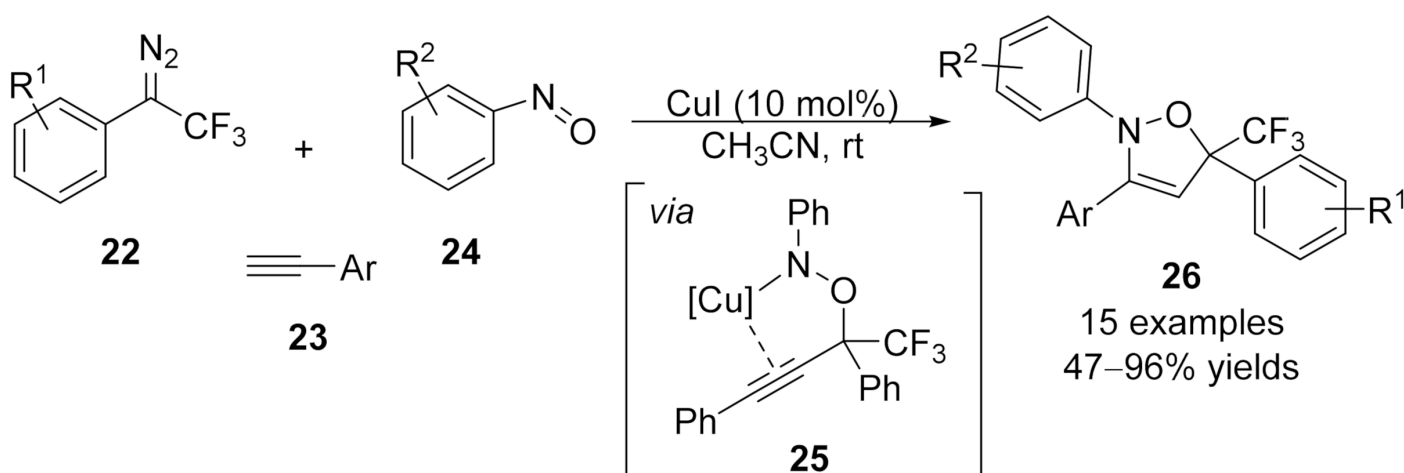
1.2. Alkynylation Terminated by Electrophilic Addition

Beyond the copper-catalyzed alkynylation terminated by protonation, the alkynoate copper intermediates, formed in situ from copper carbene species and terminal alkynes, could be intercepted through a nucleophilic substitution or electrophilic addition process. In 2015, Wang and co-workers contributed a three-component cross-coupling reaction of terminal alkyne with α -diazo ester and alkyl halide or Michael acceptor ^[4]. In this transformation, α -diazoesters **15** react firstly with the (triisopropylsilyl)acetylene **16** through a migratory insertion process to form the alkynoate copper intermediate **17**, followed by a nucleophilic substitution with alkyl halides **18** or Michael addition with electron-deficient alkenes **20** to produce the three-component products **19** and **21**, respectively ([Scheme 4](#)). This transformation represents a highly efficient method for the construction of alkynylation products with an all-carbon quaternary center in moderate to high yields. Notably, the copper catalyst works as the only catalyst to install two new C–C bonds on the carbenic carbon in this reaction.



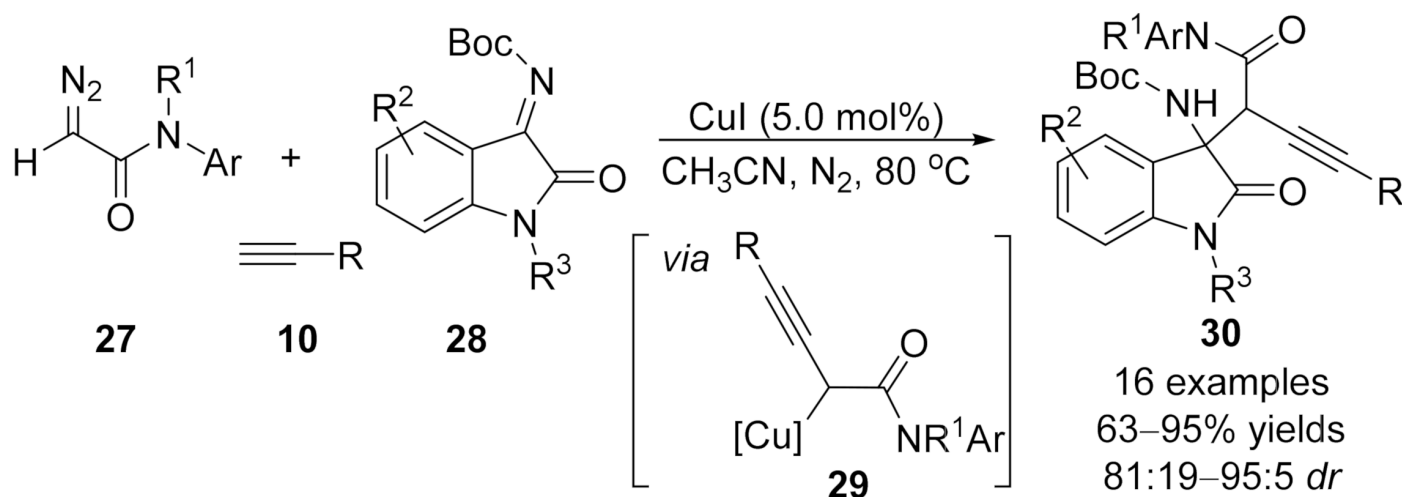
Scheme 4. Copper-catalyzed alkynylation terminated by nucleophilic substitution/Michael addition.

In 2018, Hu and co-workers reported a copper-catalyzed three-component [1+2+2]-cycloaddition of trifluoromethyl diazo compounds **22** with terminal alkynes **23** and nitrosoarenes **24** [5]. With this method, a series of trifluoromethyl-substituted dihydroisoxazoles **26** were obtained in high yields under mild conditions. Mechanistically, electrophilic trapping of the alkynoate copper intermediate by nitrosobenzenes was proposed as the key step in this cascade transformation, which forms a proposed intermediate **25**, followed by a copper catalyzed intramolecular annulation to deliver the target products **26** (Scheme 5).



Scheme 5. Three-component reaction of diazo compound with terminal alkyne and nitrosobenzene.

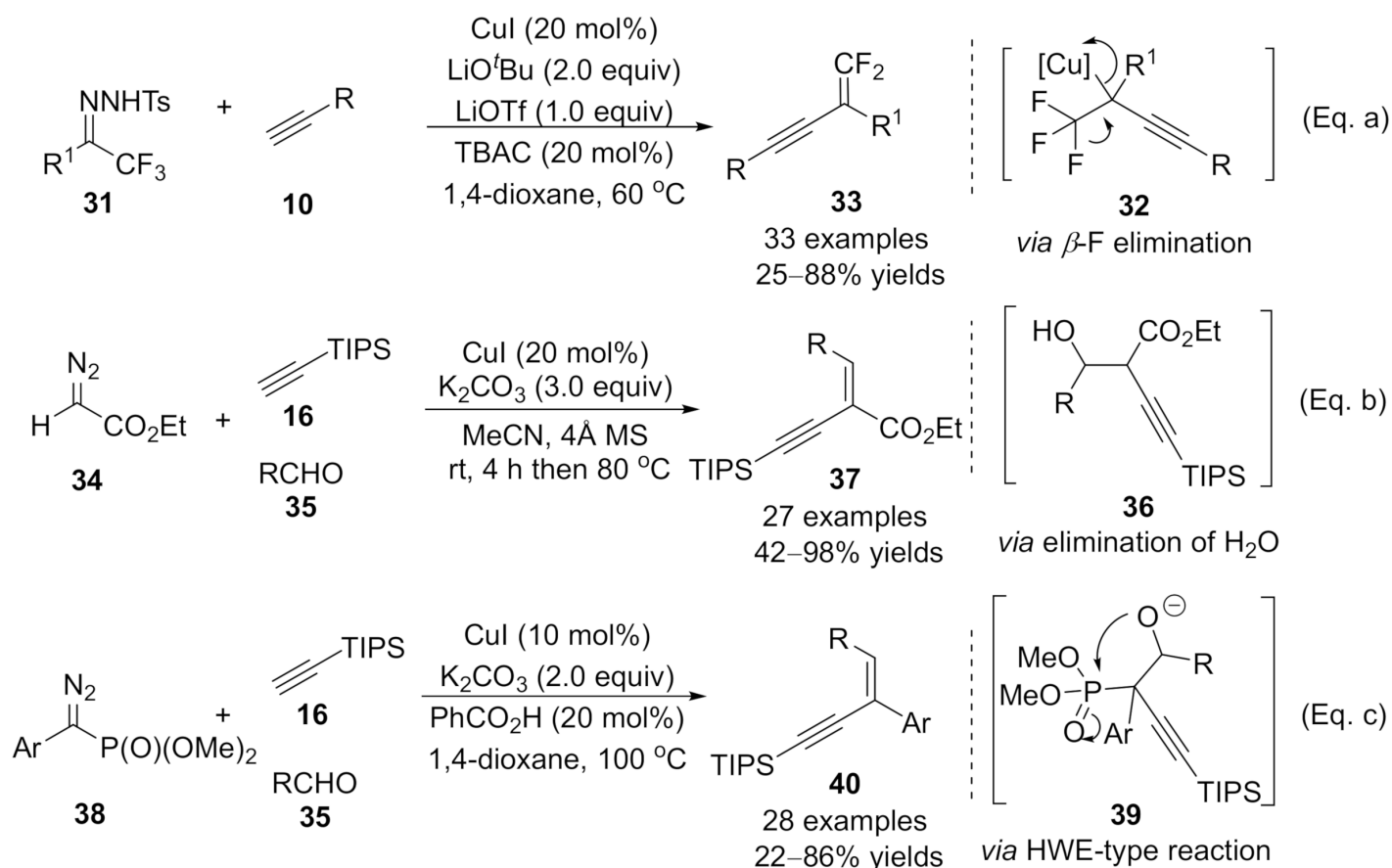
One year later, the same group reported a copper-catalyzed three-component reaction of terminal alkynes **10** with α -diazoamides **27** and isatin ketimines **28** in 2019 [6]. A series of alkynyl-containing 3,3-disubstituted oxindoles **30** were efficiently formed in high yields and diastereoselectivities through a Mannich type trapping of an in situ generated alkynoate copper intermediate **29** (Scheme 6).



Scheme 6. Copper-catalyzed three-component reaction of α -diazoamide with terminal alkyne and isatintetimine.

1.3. Alkynylation Terminated by β -Elimination

Besides the intermolecular trapping reactions of the alkynoate copper intermediate, β -H elimination could occur with this reactive species, providing a variety of conjugated enynes. Representative advances in this area have been reported by Wang's group [7][8][9]. In 2015, they reported a copper-catalyzed cross-coupling reaction of terminal alkynes **9** with trifluoromethyl ketone *N*-tosylhydrazones **31**, which provides an efficient synthesis of 1,1-difluoro-1,3-enyne derivatives **33** under mild reaction conditions [7]. Mechanistically, the alkynoate copper intermediate **32** was generated in situ through a migratory insertion of the copper carbene intermediate, followed by β -F elimination, leading to the *gem*-difluoroolefination products **33** in moderate to high yields [Scheme 7, Eq. (a)]. Later, researchers reported a copper-catalyzed three-component reaction of a (triisopropylsilyl)acetylene **16** with Ethyl diazoacetate **34** (EDA) and aldehydes **35** that provided an efficient method for the synthesis of α -alkynyl- α , β -unsaturated esters **37** [8]. In this cascade reaction, nucleophilic aldol addition of an alkynoate copper intermediate with aldehyde **35** formed product **36**, which delivered the desired products **37** as a single (*E*)-stereoisomers through an elimination process in good to excellent yields [Scheme 7, Eq. (b)]. In the same year, an analogous cross-coupling reaction with α -diazo phosphonates **38** instead of EDA through a sequential alkynylation/aldol addition/Horner–Wadsworth–Emmons (HWE) type reaction was disclosed by the same group [9]. This method provided straightforward access to conjugated enynes **40** with good stereoselectivity and excellent functional group compatibility [Scheme 7, Eq. (c)]. Moreover, one C–C bond and one C=C bond were formed successively in a one-pot manner, making those novel enynes synthesis methods practically useful.

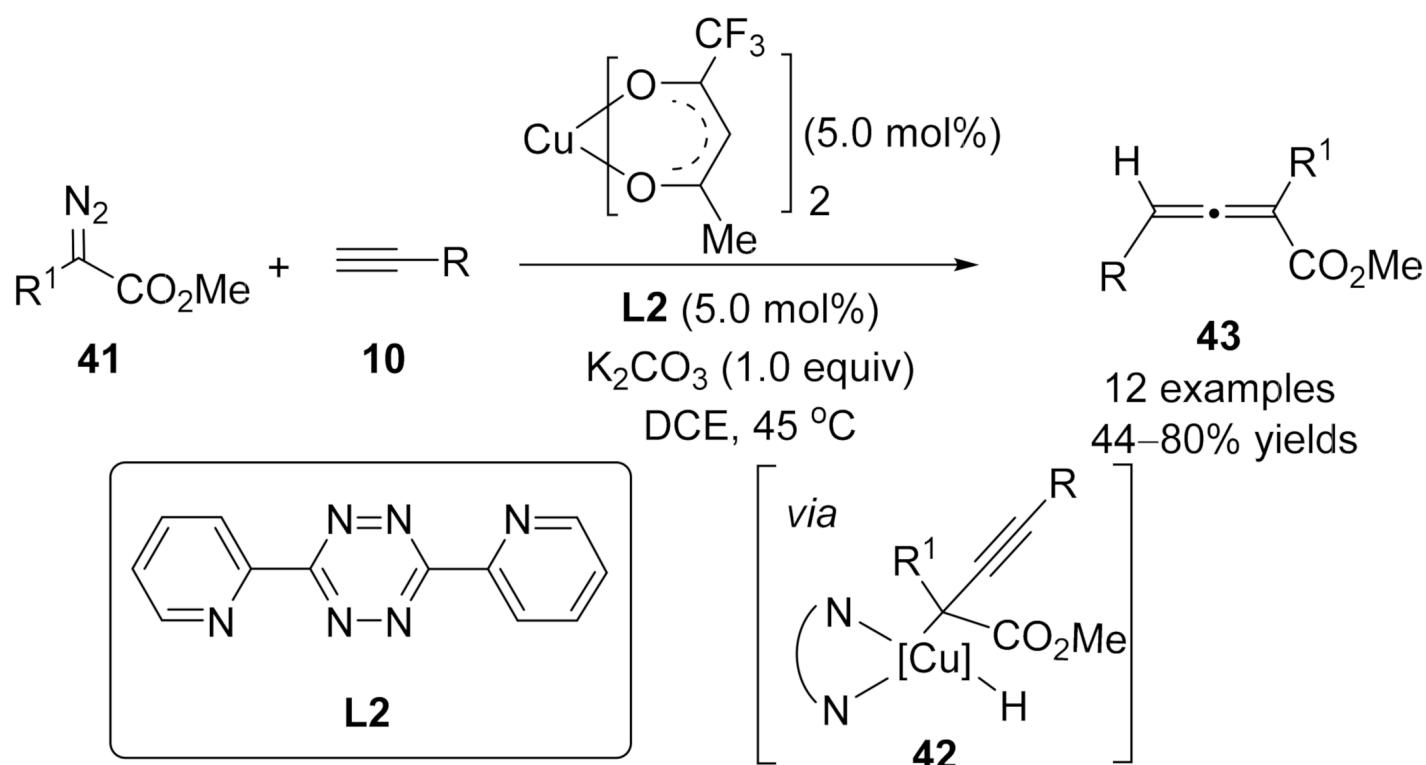


Scheme 7. Copper-catalyzed alkynylation terminated by β -elimination. (a) Coupling reaction of *N*-tosylhydrazone with alkyne. (b) Coupling reaction of diazoacetate with alkyne. (c) Coupling reaction of α -diazo phosphonate with alkyne.

2. Allenylation

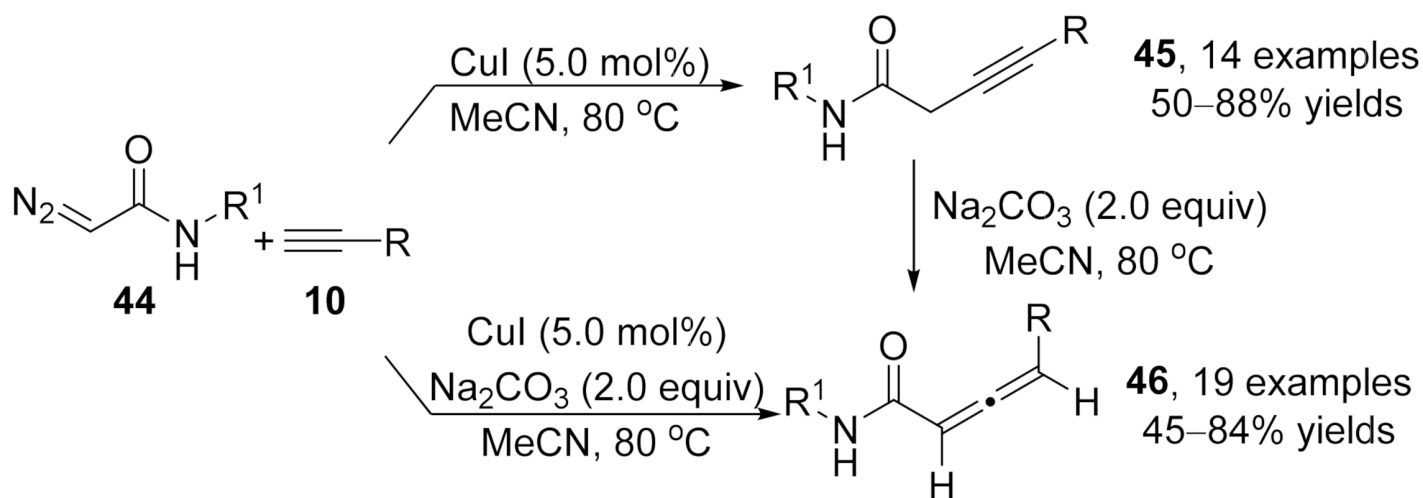
2.1. Allenylation Terminated by Protonation

As a complementary to Fu's method for the selective synthesis of alkynoates ^[10], in 2011, Fox's group reported a selective coupling reaction of α -substituted- α -diazoesters **41** with terminal alkynes **10** to the syntheses of allenates in the presence of Cu(II)(trifluoroacetylacetonate)₂ and 3,6-di(2-pyridyl)-s-tetrazine **L2** in DCE ^[11]. As a result, allenates **43** were obtained as the main products with slight traces of the alkynoates. Key to the development of this selective method was the recognition of an adventitious base, potassium carbonate, which improved the selectivity of isomerization to form the allenate products (Scheme 8).



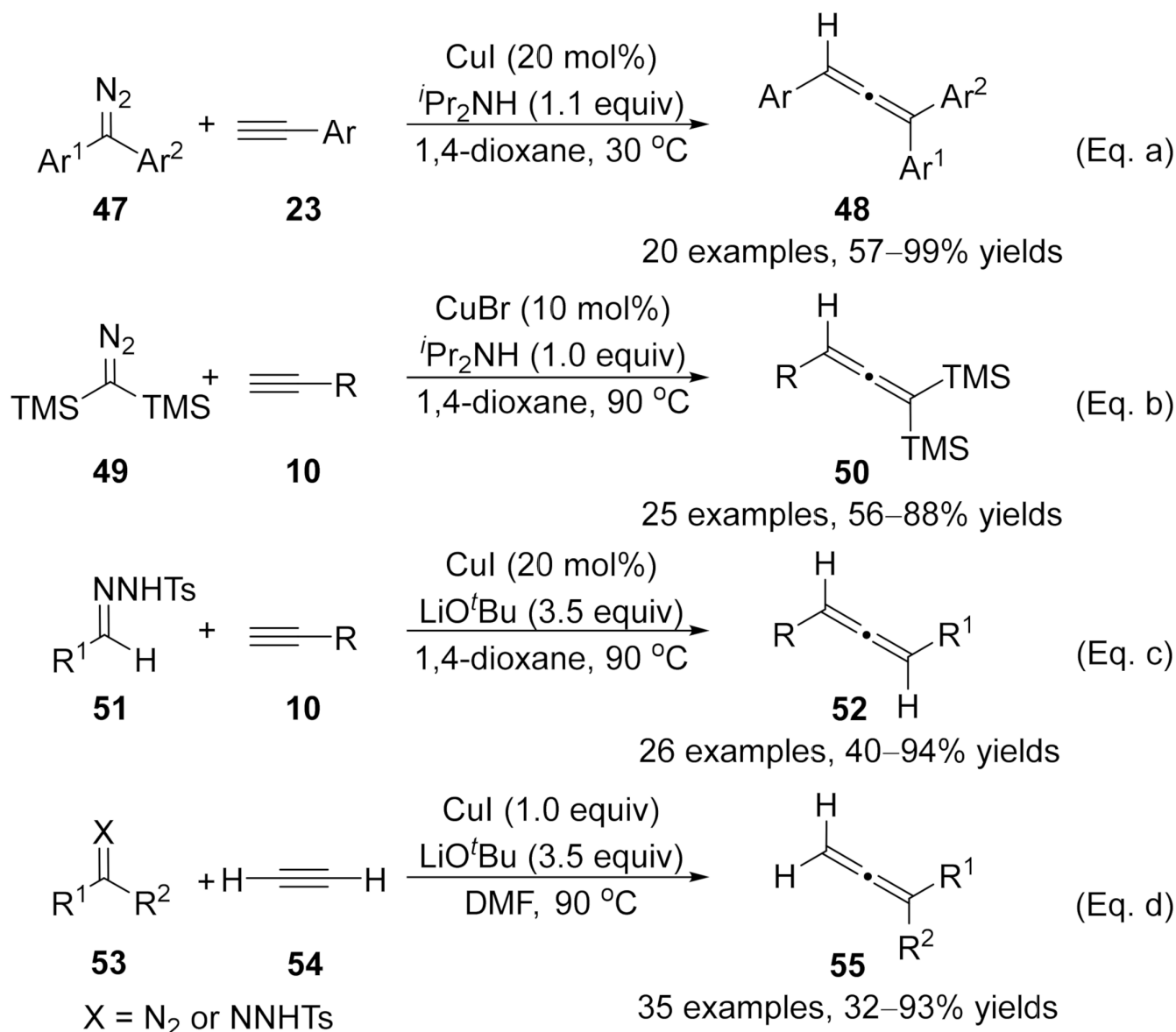
Scheme 8. Cross-coupling reaction of copper carbene with terminal alkynes for the synthesis of allenates.

Later in 2013, an efficient copper-catalyzed cross-coupling between diazoacetamides **44** and terminal alkynes **10** under ligand-free conditions was developed by Sun [12]. This method provided a practical method for the assembly of substituted 3-butynamides **45** and dienamides **46**. Interestingly, when sodium carbonate was added to the reaction mixture, the allenic compounds **46** were obtained as the main products. However, alkyne products **45** were generated as the major products in the absence of this base (Scheme 9). Moreover, the alkynoate compounds **45** could be smoothly converted into the isomeric allenes **46** in the presence of sodium carbonate without assistance of the copper catalyst.



Scheme 9. Copper-catalyzed divergent cross-coupling reaction of diazoacetamides with terminal alkynes.

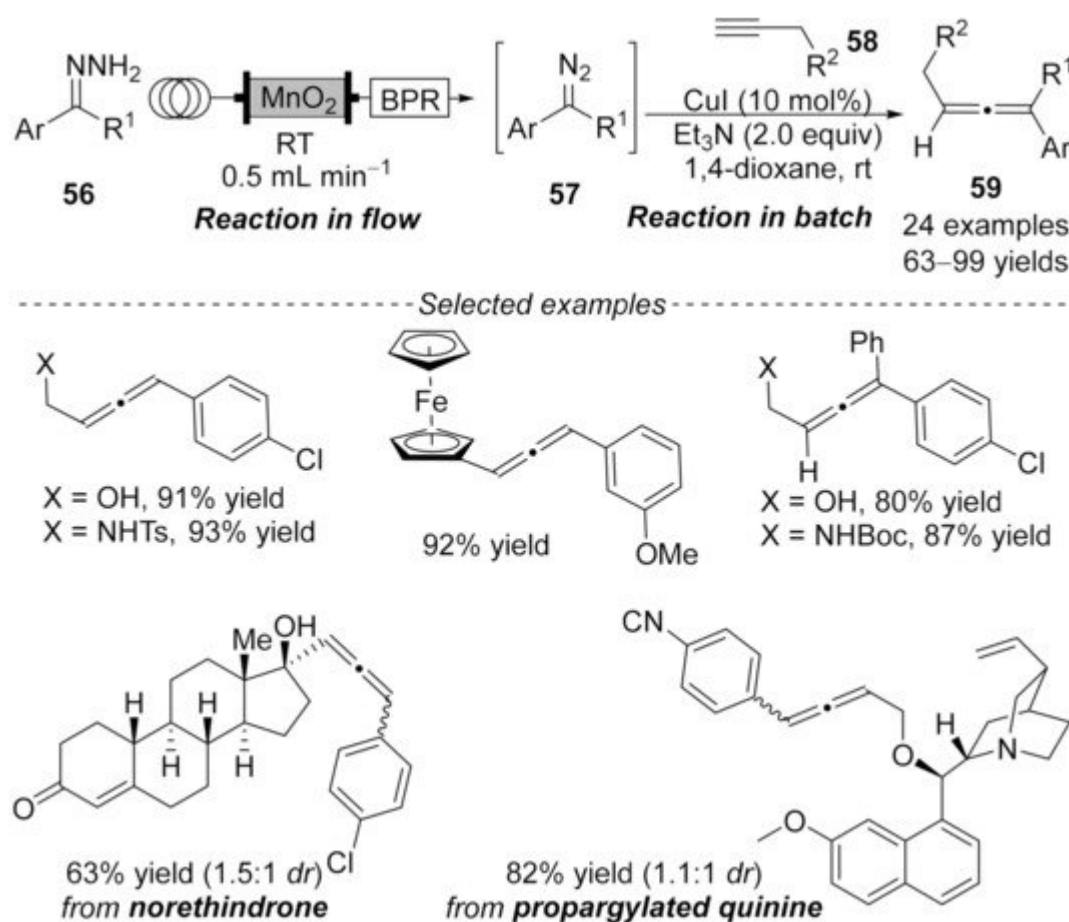
In the same year, Wang and co-workers developed a series of synthetic methods to form substituted allenes under optimized conditions in the presence of copper(I) complexes ^{[13][14][15][16]}. Different types of substituted diazo compounds and *N*-tosylhydrazones were employed as the carbene precursors for the coupling with various terminal alkynes, delivering the allenic derivatives in good yields with a wide range of functional group tolerance ([Scheme 11](#)). Notably, ethyne **54** was also a compatible substrate for this reaction, which leads to a new synthetic method for the synthesis of terminal allenes **55** in moderate to excellent yields^[16]. However, using one equivalent amount of CuI with DMF as the solvent is critical to the success of this transformation [[Scheme 10](#), Eq. (d)].



Scheme 10. Copper-catalyzed cross-coupling reaction for the syntheses of substituted allenes. (a) Cross-coupling reaction of diaryl diazo compound with alkyne. (b) Cross-coupling reaction of diazo compound **49** with alkyne. (c) Cross-coupling reaction of *N*-tosylhydrazone **51** with alkyne. (d) Cross-coupling reaction of **53** with acetylene.

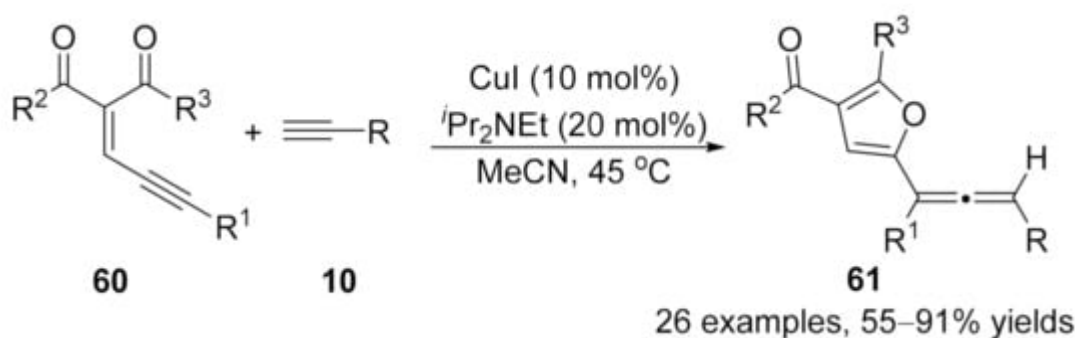
In 2015, a copper-catalyzed coupling reaction between flow-generated unstabilized diazo compounds and terminal alkynes was reported by Ley's group, providing a practical method for the synthesis of di- and tri-substituted

allenes **59** in high yields under mild conditions [17]. The unstable diazo compounds **57** were generated in situ from hydrazones **56** through oxidation with activated MnO_2 . Then, the above solution was injected directly into the other reaction mixture, which contained terminal alkynes **58**, base, and CuI catalyst. The reaction delivered the allene products **59** in good to excellent yields. To highlight the selectivity and functional group compatibility of this protocol, norethindrone and propargylated quinine were successfully applied to the optimal reaction conditions, generating the corresponding products in 63% and 82% yields, respectively (Scheme 11).



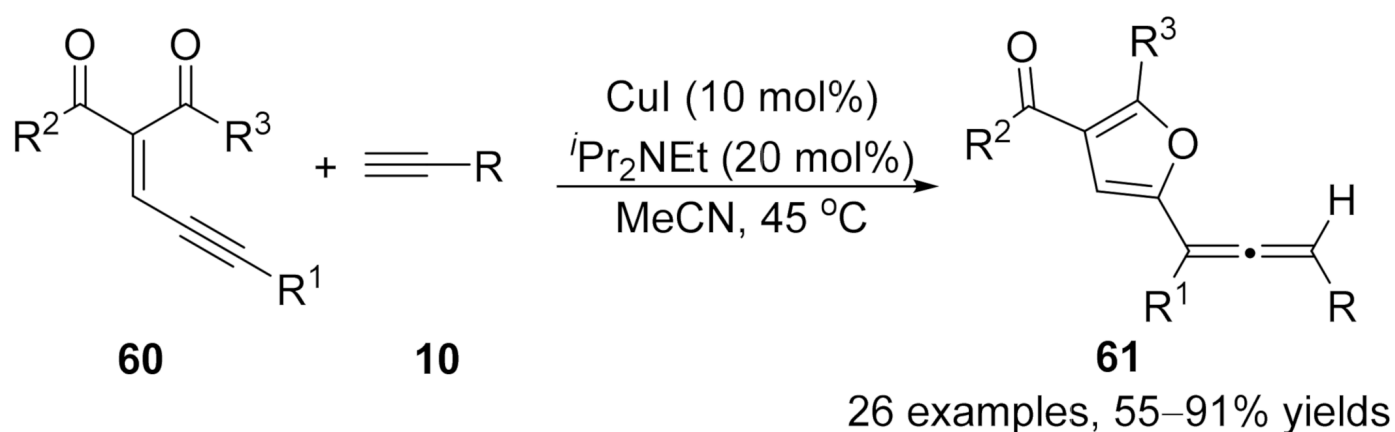
Scheme 11. Copper-catalyzed cross-coupling reaction via flowing chemistry.

In addition to the diazo compounds, conjugated eneyne ketones **60** were introduced as carbene precursors by Wang and co-workers in 2016 in a copper-catalyzed cross-coupling reaction with terminal alkynes [18]. This reaction afforded trisubstituted allenes **61** in high yields with broad functional group tolerance under mild reaction conditions (Scheme 12).



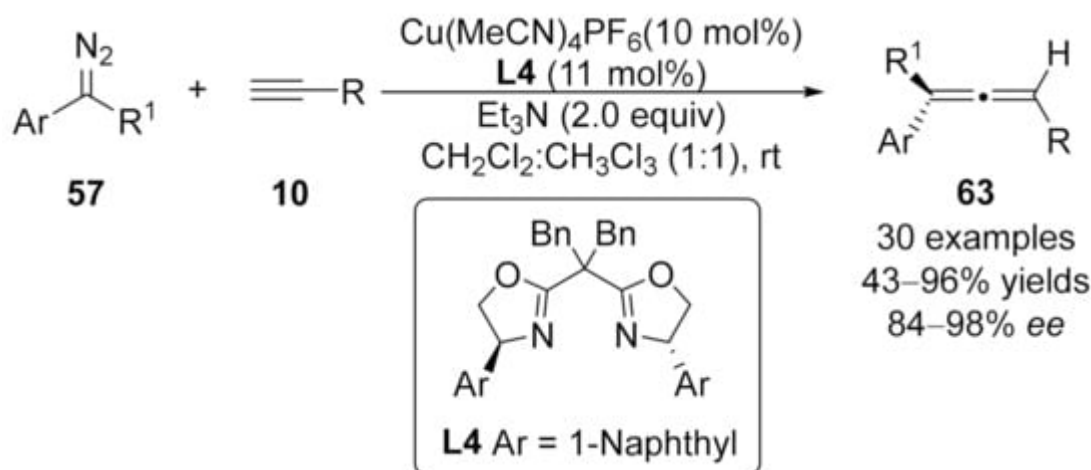
Scheme 12. Copper-catalyzed cross-coupling reaction of conjugated enyne ketones with terminal alkynes.

In 2015, Feng and Liu contributed an asymmetric cross-coupling of α -diazoesters **15** with terminal alkynes **10** using chiral Cu(I) /guanidine complex as the catalyst [19]. Notably, no additional base was necessary for this transformation, providing optically active 2,4-disubstituted allenoates **62** under mild reaction conditions in good to high yields (up to 99 %) with good to excellent enantioselectivities (Scheme 13).



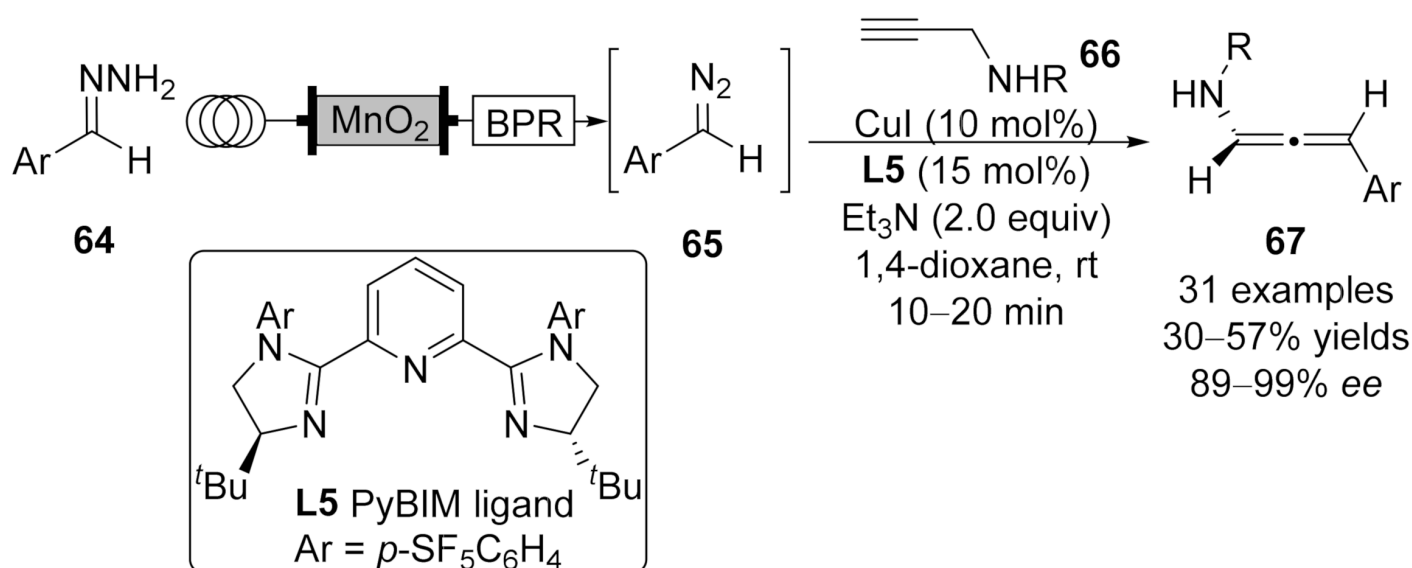
Scheme 13. Asymmetric cross-coupling reaction catalyzed by copper/guanidine complex.

In 2016, Wang and co-workers reported a highly enantioselective copper-catalyzed cross-coupling of aryldiazoalkanes **57** with terminal alkynes **10** [20]. By utilizing chiral Cu(I) /bisoxazoline ligand **L4**, this reaction delivered a series of trisubstituted allenes **63** in moderate to high yields (up to 96%) with excellent enantioselectivities (up to 98% ee). Unlike the previous works using CuI as the catalyst, $\text{Cu}(\text{MeCN})_4\text{PF}_6$ complex was used as the optimal metal catalyst to enable high reactivity and stereoselectivity in this reaction (Scheme 14).



Scheme 14. Asymmetric cross-coupling reaction catalyzed by copper/Box complex.

One year later, Ley's group reported their continuous flow strategy for the asymmetric coupling reaction of unstabilized diazo compounds **65** with propargyl amines **66** in the presence of chiral Cu(I)/PyBIM complex [21]. This method generated the amino-substituted chiral allenates **67** in moderate yields (up to 57%) with high enantioselectivities (up to 96% ee) in a fast reaction rate (10–20 min) with a variety of functional group compatibility (Scheme 15).

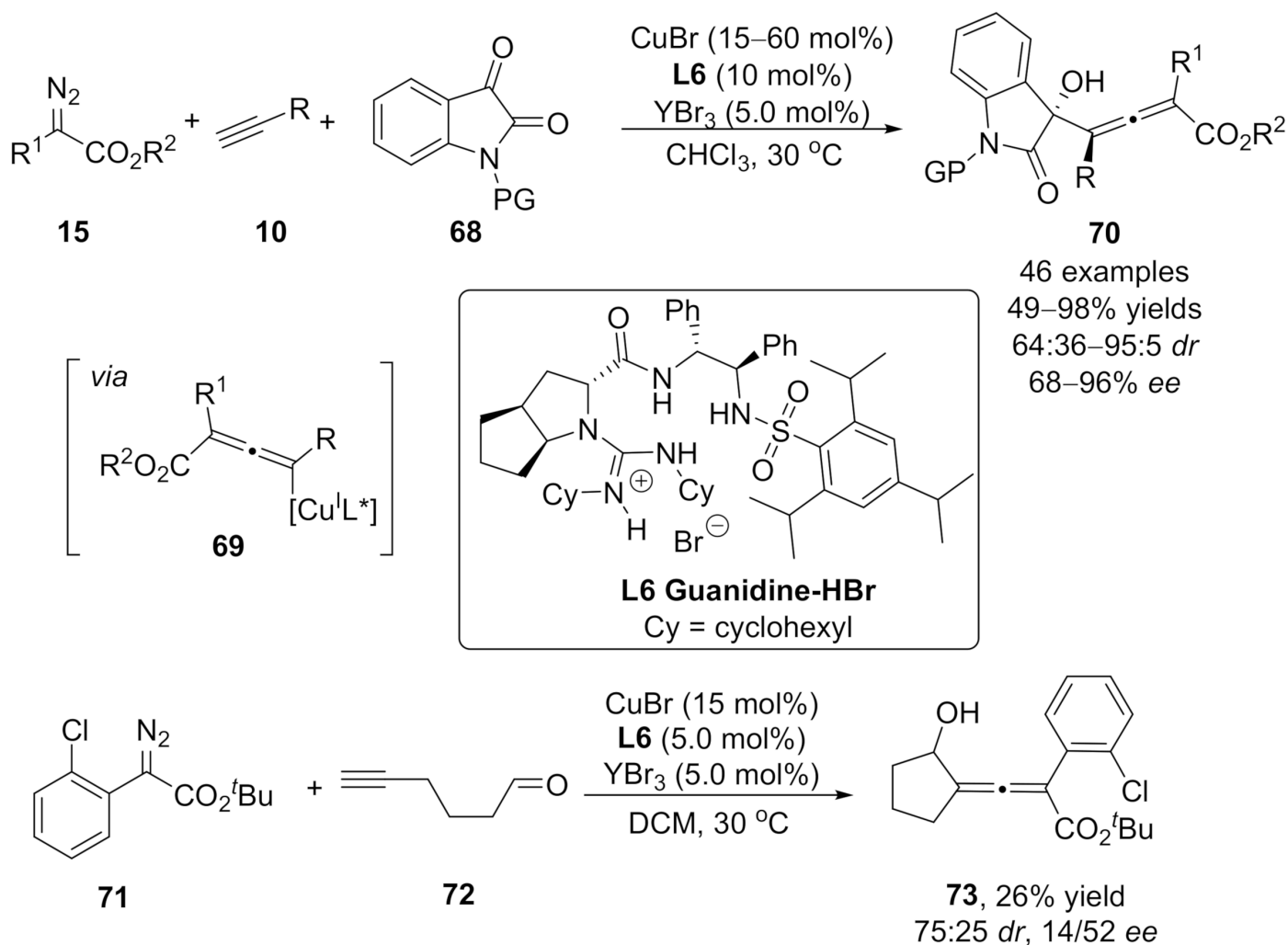


Scheme 15. Asymmetric cross-coupling reaction catalyzed by copper/PyBIM complex through flowing chemistry.

2.2. Allenylation Terminated by Electrophilic Addition

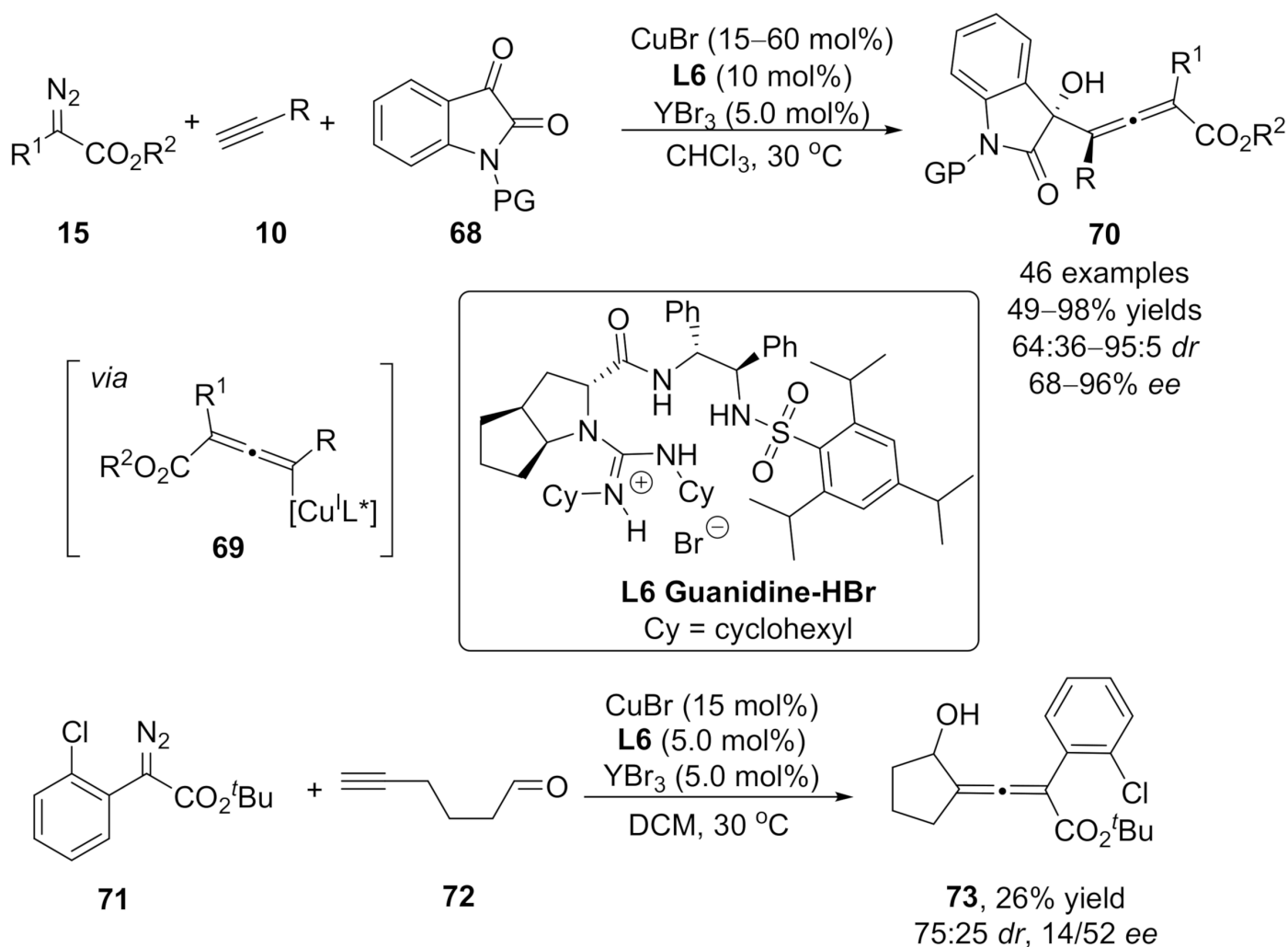
In 2018, a one-pot copper-catalyzed asymmetric three-component reaction of diazoesters **15** with terminal alkynes **10** and isatins **68** was reported by Liu's group [22]. Axially chiral *tetra*-substituted allenates **70** bearing a stereogenic center were obtained under a chiral Cu(I)/guanidinium salt/YBr₃ catalytic system with high diastereo- and enantioselectivities. The aldol type addition of allenolate-copper intermediates **69** with isatins **68** has been

proposed as the key step in this reaction. Moreover, convincing experimental evidence for the formation of allenolate-copper intermediate **69** was provided through the synthesis of chiral allenolate, which was generated from the C–H insertion reaction of α -diazooester with alkyne. Researchers found that additional acids improved the catalyst efficiency of the chiral copper complex. The intramolecular nucleophilic trapping reaction of allenolate-copper intermediate with embedded aldehyde species was also successful, generating the cyclic allenolate product **73**, albeit the yield and stereoselectivity were moderate ([Scheme 16](#)).



Scheme 16. Asymmetric three-component reaction for the synthesis of tetra-substituted allenates.

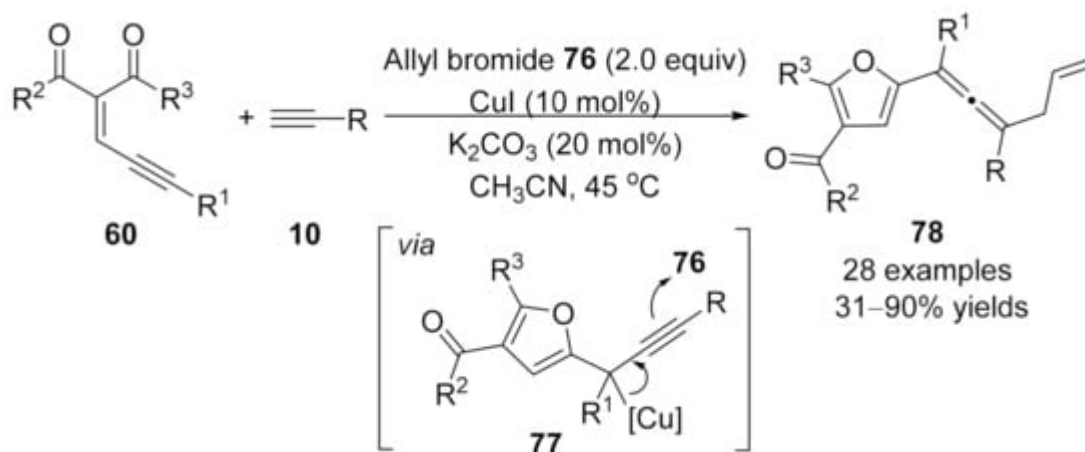
Recently, Sun's group has realized an enantioselective intramolecular nucleophilic aldol addition of in situ formed allenolate-copper intermediate with aldehyde using chiral Cu(II)/Box complex [\[23\]](#). Distinct from the previous version with copper(I) catalysts, this protocol used copper(II) salt as an optimal catalyst in this asymmetric cross-coupling reaction. The *tetra*-substituted allenates **75** containing both central and axial chiralities have been obtained in moderate to good yields with good to excellent stereoselectivities. ([Scheme 17](#)).



Scheme 17. Copper-catalyzed asymmetric allenylation terminated by aldol addition.

2.3. Allenylation Terminated by Allylation

In 2016, Wang and co-workers realized the synthesis of allyl-substituted allenes through trapping of allenolate-copper intermediate with allyl bromide through a nucleophilic substitution process ^[18] In this reaction, conjugated eneyne ketones **60** have been used as the carbene source. Mechanistically, the copper-(2-furyl) carbene intermediate was generated in situ from eneyne in the presence of CuI, followed by a migratory insertion process to afford nucleophilic alkynoate copper intermediate **77** that was trapped by allyl halide **76** (Scheme 18). In this method, the choice of the base was pivotal for the reaction outcomes when K₂CO₃ was employed as the base, affording 2-furyl substituted allenes **78** in generally good yields.



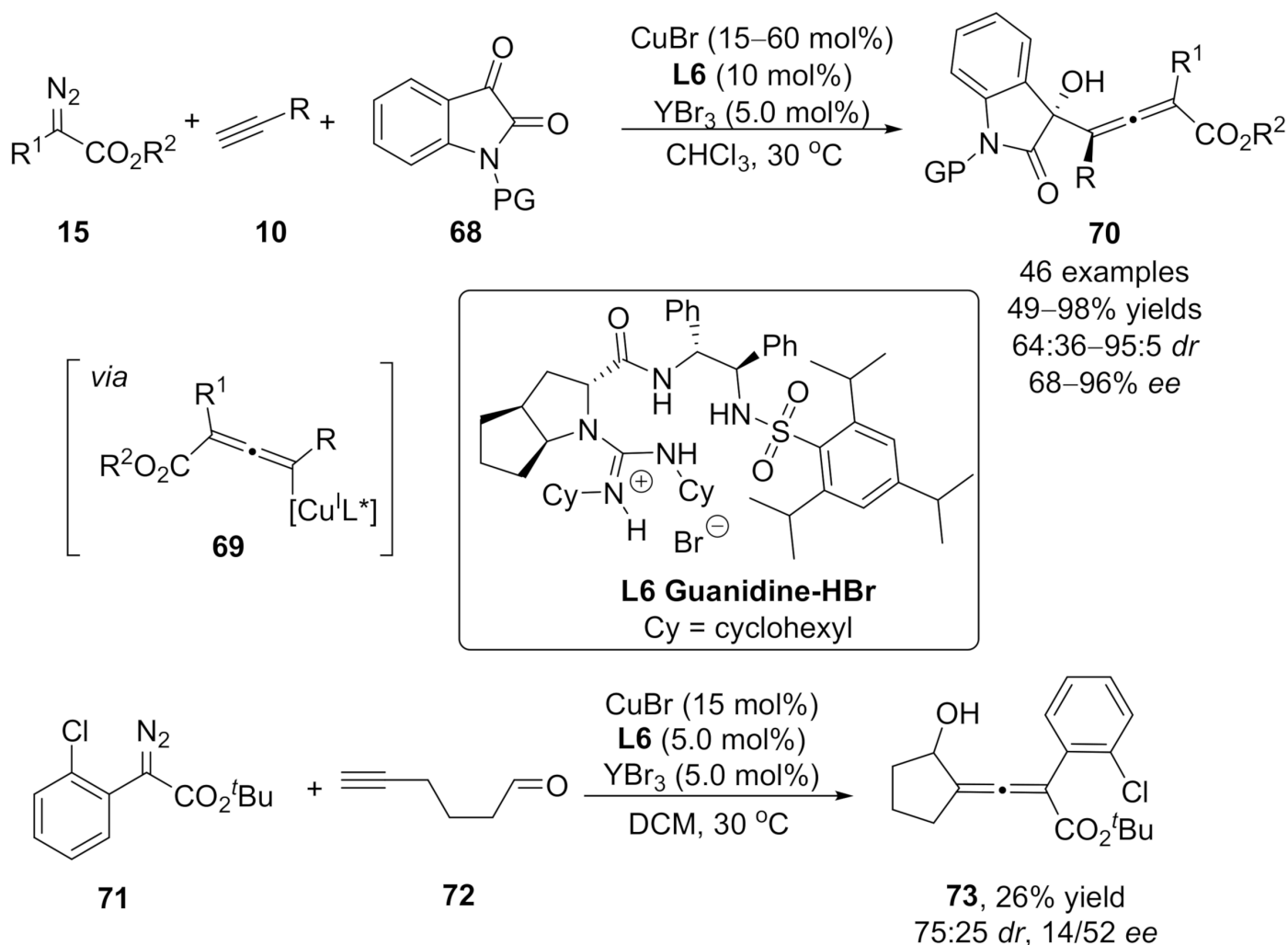
Scheme 18. Copper-catalyzed allenylation terminated by allylation.

2.4 Cascade Transformations Involving Allenylation Process

Nucleophilic addition with allenic ester or its isomeric compound is generally used synthetic strategy for the expeditious construction of highly functionalized carbocycle or heterocycle structures [24][25][26][27][28][29][30][31][32]. Thus, a variety of inter- or intramolecular cascade reactions have been developed through different nucleophilic addition processes of the allene derivatives generated from cross-coupling between alkynes and copper carbenes.

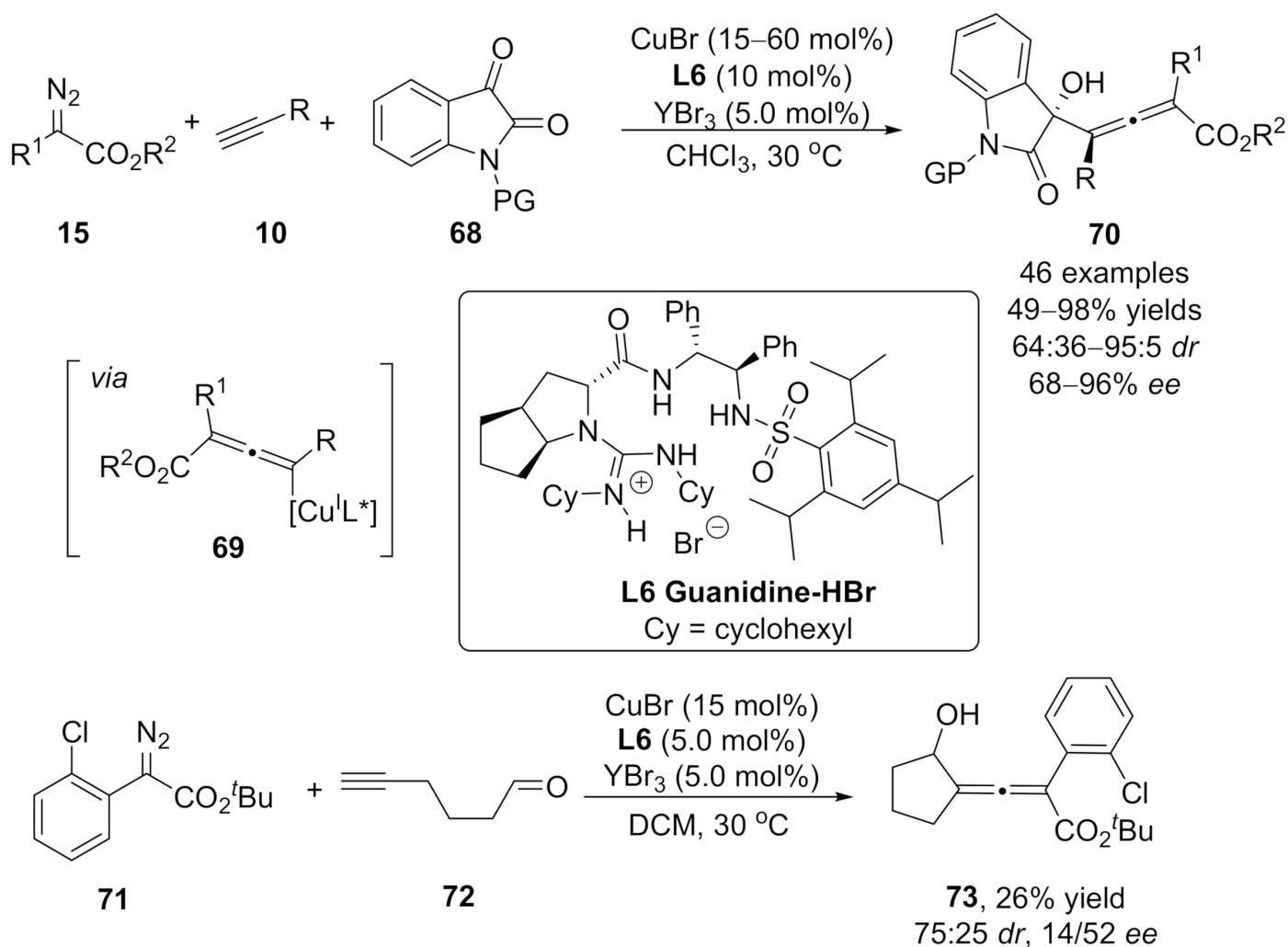
In 2011, a one-pot synthesis of phenanthrenes **81** via ligand-free CuBr₂-catalyzed coupling reaction/intramolecular cyclization of terminal alkynes **23** with N-tosylhydrazones **79** derived from o-formyl biphenyls was developed by Wang and co-workers [33]. Allene intermediates **85** were initially generated in this cascade reaction through a cross-coupling reaction of N-tosylhydrazones **79** with terminal alkynes **23**, followed by a 6π-cyclization and isomerization

deliver the phenanthrene products **81** with broad functional group compatibility (Scheme 19).



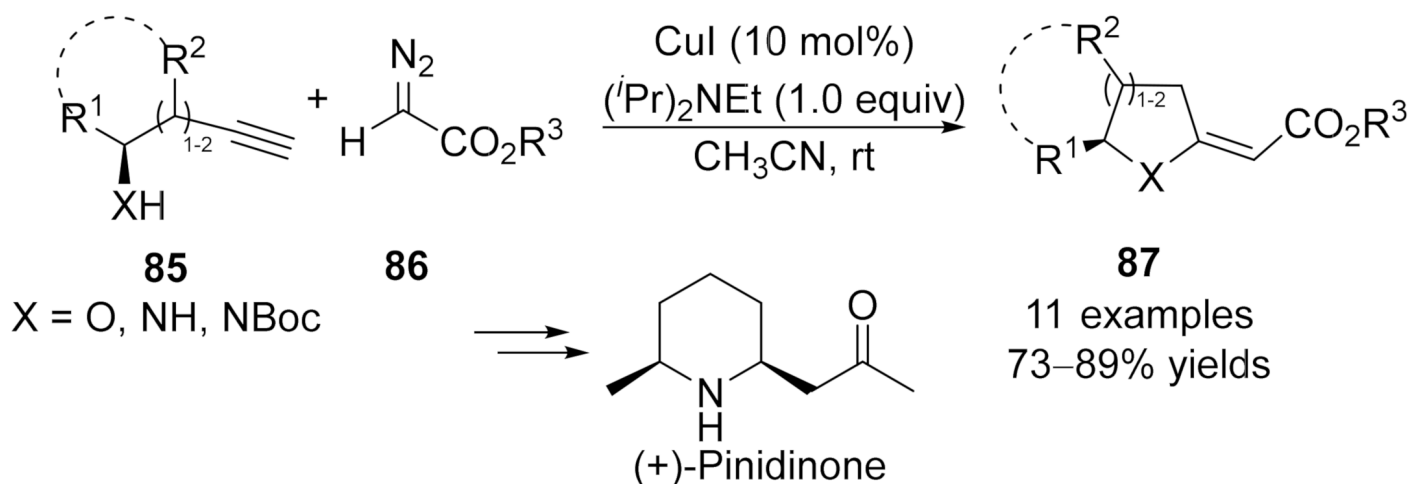
Scheme 19. Copper-catalyzed allenylation followed by 6 π -cyclization.

Later in the same year, instead of using *o*-aryl substituted *N*-tosylhydrazones, *o*-hydroxy- or *o*-amino-substituted *N*-tosylhydrazones were introduced by the same group as carbene precursors in an analogous cascade transformation, a ligand-free CuBr-catalyzed coupling reaction/intramolecular cyclization sequence, achieving the synthesis of benzofuran or indole derivatives **84** in moderate to excellent yields [34]. The initially formed allene intermediates **83** were trapped through a nucleophilic addition by the embedded *o*-hydroxy- or *o*-amino group to afford the cyclized products **84** (Scheme 20).



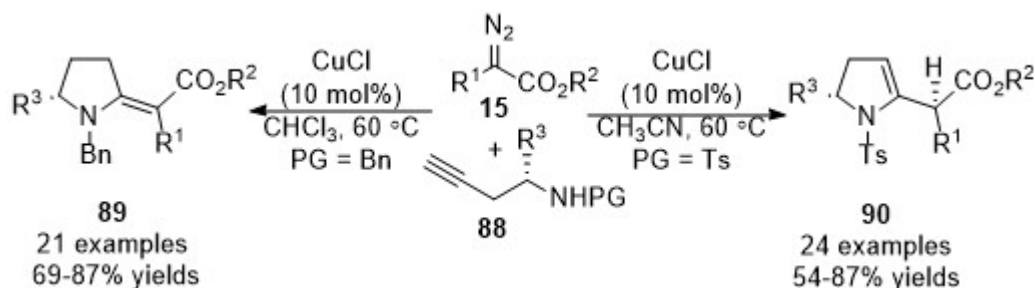
Scheme 20. Copper-catalyzed allenylation followed nucleophilic addition.

In 2011, a similar catalytic strategy was developed by Balakishan's group. They reported a simple procedure for synthesizing aza- and oxacycles via a copper-catalyzed coupling reaction of functionalized terminal alkynes **85** with diazoesters **86** [35]. Initially, the allene intermediates were formed in the presence of CuI , followed by an intramolecular aza- or oxa-Michael cycloaddition and isomerization to generate the cyclized five- or six-membered products **87** in generally good yields (Scheme 21).



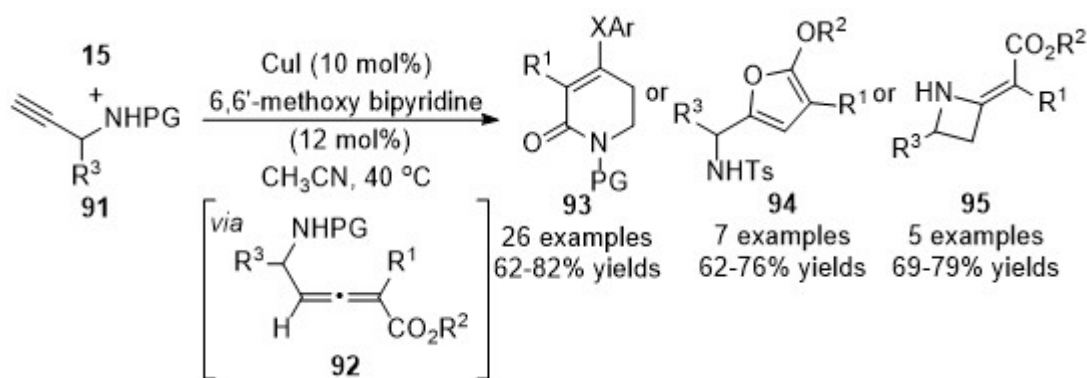
Scheme 21. Copper-catalyzed allenylation followed by cyclization.

In 2015, a stereo-divergent synthesis of five-membered heterocycles was developed by Sun's group [36]. The proposed reaction mechanism involves trapping *in situ* formed allene intermediates, yielding 2-methylenes **89** (when PG = Bn) and 2,3-dihydropyrroles **90** (when PG = Ts) in good yields with broad functional group tolerance under mild conditions. Control experimental results show that *N*-benzyl amino alkynes were more likely to form 2-methylenespyrroles derivatives **89** through 5-*exo-dig* cycloaddition, while 2,3-dihydropyrroles **90** generated from *N*-tosylamino alkynes through 5-*endo-dig* cycloaddition would be more favorable (Scheme 22).



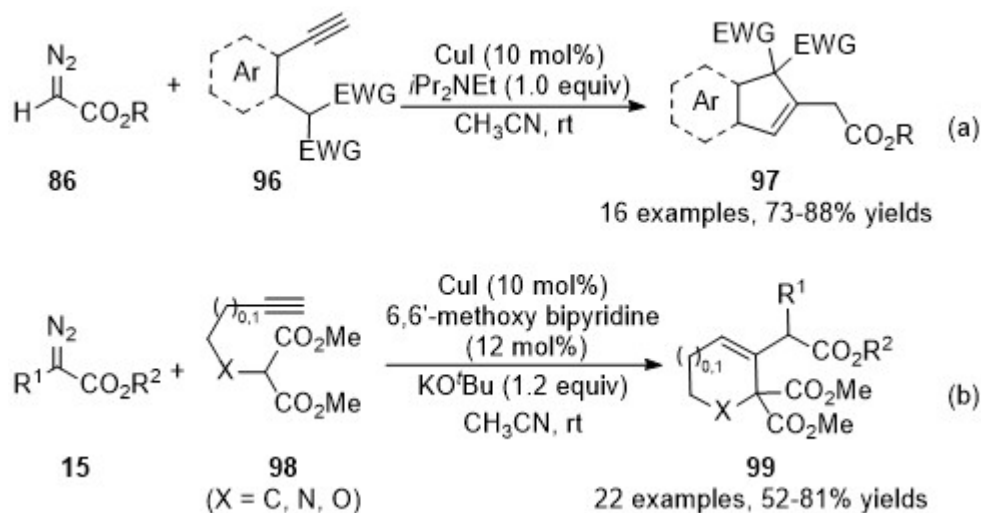
Scheme 22. Copper-catalyzed allenylation followed by divergent annulation.

In 2018, Sun and co-workers expanded the above chemistry to synthesize the four- to six-membered heterocycles with *N*-substituted prop-2-yn-1-amines **91** and diazoacetates **15** [37]. Generated allenoid species **92** have been proven as the key intermediates for the subsequent diverse annulations under optimized conditions toward functionalized heterocycle in moderate to good yields. Treatment of allenoidates **92** with sodium phenolates led to six-membered products **93**; silver nitrate and triethylamine yielded five-membered products **94**; what's more, four-membered products **95** were generated under lithium *tert*-butoxide conditions (Scheme 23).



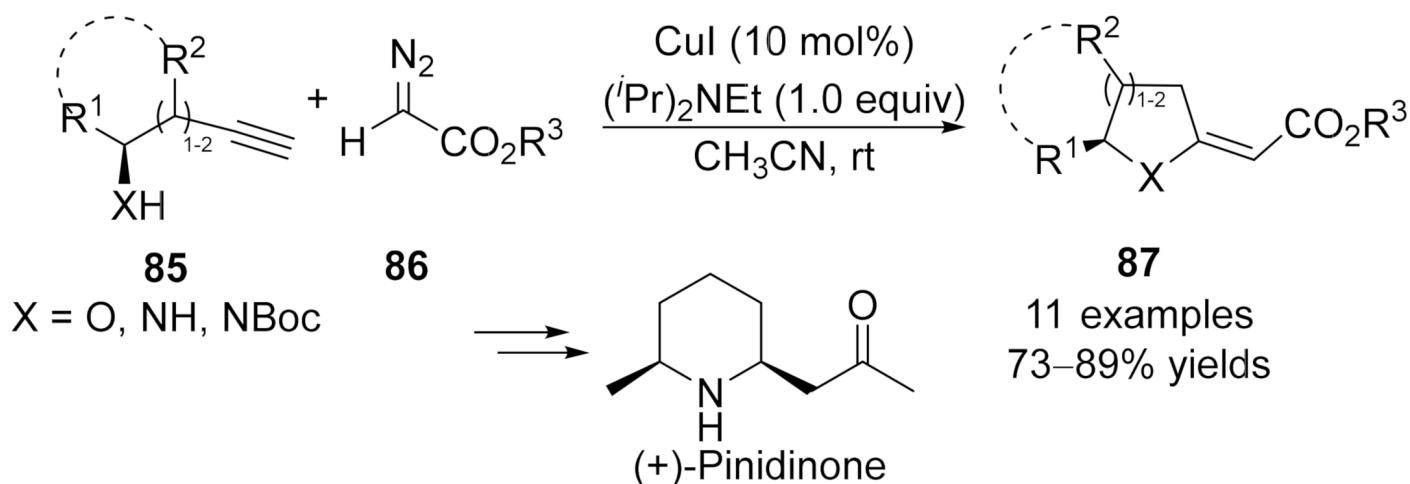
Scheme 23. Divergent synthesis of four- to six-membered heterocycles involving an allenylation process.

In addition to the cyclization through addition with a heteroatom, carbon-based nucleophilic species could also be served as the nucleophile to addition with these allenes, forming the C-C bond instead of the C-X bond [38][39]. In 2015, Kumaraswamy's group developed a copper catalyzed cross-coupling reaction/intramolecular Michael addition cascade reaction [40], achieving the formation of indene and dihydronaphthalene derivatives **97** in good yields with broad functional group tolerance (Scheme 24a). Later in 2017, Sun's group reported an analogous approach toward five- or six-membered carbo-/heterocycles with diazo compounds **15** and alkyne-substituted malonates **98** [41]. In this reaction, the ligand significantly enhanced the reaction yields and inhibited the Conia-ene side reaction. As a result, the polyfunctionalized cyclohexenes, tetrahydropyridines, and dihydropyrans have been prepared in moderate to high yields under mild reaction conditions (Scheme 24b).



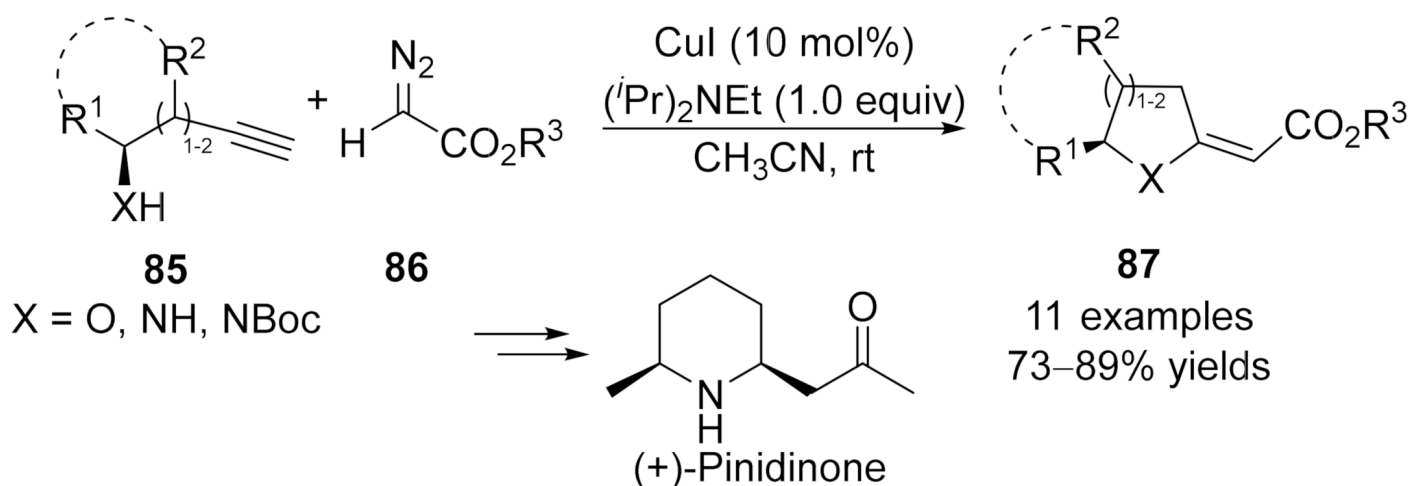
Scheme 24. Copper-catalyzed allenylation followed by Michael addition.

In 2015, a Cu(I)-catalyzed denitrogenative annulation reaction of pyridotriazoles **100** with terminal alkynes **10** was developed by Gevorgyan's group [42]. Initially, α -pyridyl copper carbenes were generated from pyridotriazoles **100** in the presence of the copper catalyst, followed by a cross-coupling reaction with terminal alkyne to form either propargylic or allenic intermediates **101**, which were terminated by copper-catalyzed cycloisomerization to furnish the indolizines **102** in moderate to excellent yields (Scheme 25).



Scheme 25. Copper-catalyzed allenylation followed by cycloisomerization.

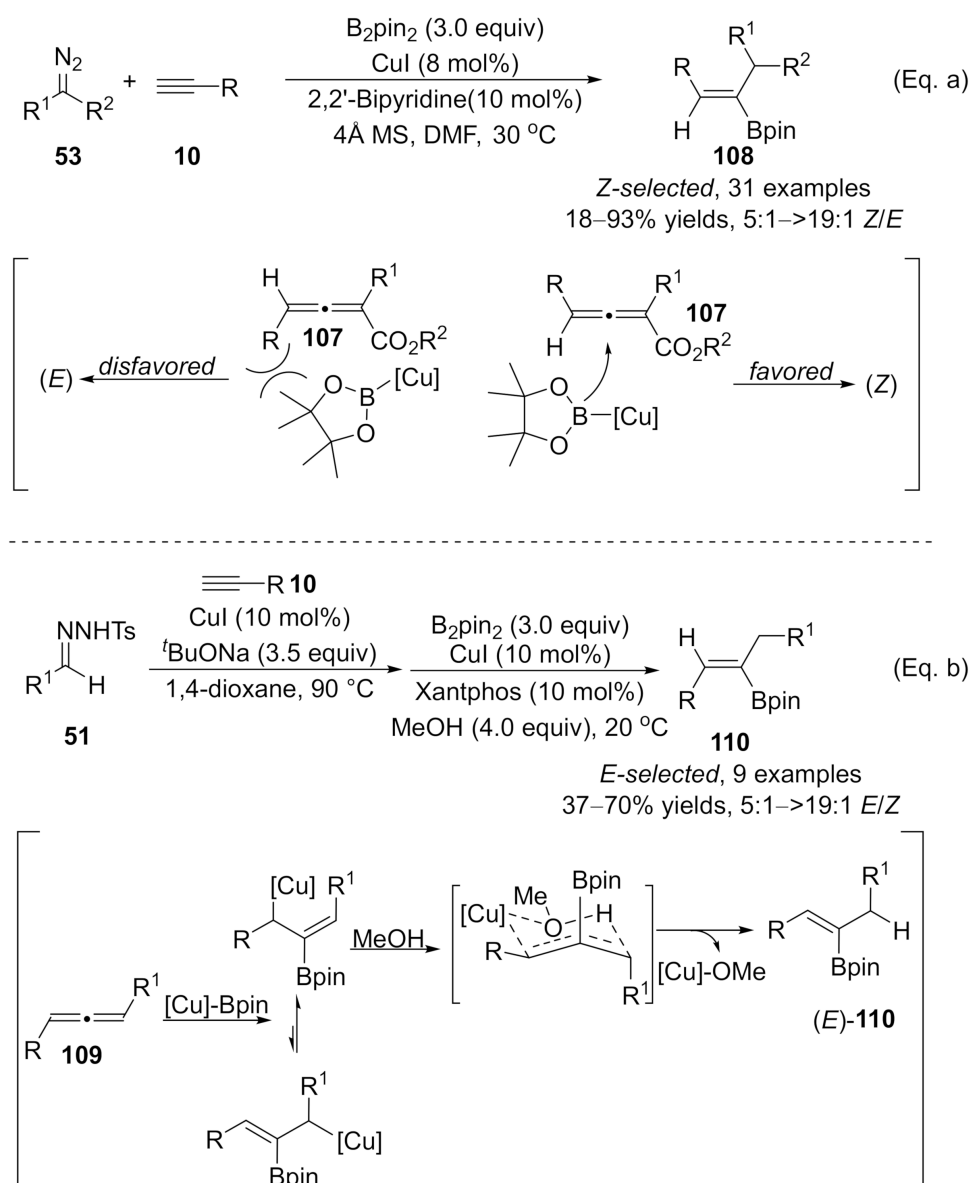
In 2018, Wang and co-workers reported a copper-catalyzed geminal difunctionalization reaction of terminal alkynes [43]. The key step in this cascade reaction is trapping the *in situ* generated allenic species **105** with a sulfonyl anion to form the carbon-sulfur bond, providing a variety of vinyl sulfones **106** in good yields with excellent stereoselectivities under mild reaction conditions. It was noted that the excellent stereoselectivities might be due to the influence of steric hindrance, and no ligand and additive were required in this transformation (Scheme 26).



Scheme 26. Copper-catalyzed allenylation followed by sulfonylation.

Recently, Sun and co-workers demonstrated a copper-catalyzed three-component reaction of terminal alkynes with diazo compounds and B_2pin_2 for the synthesis of trisubstituted alkenylboronates [44]. Copper catalysts played dual roles in these alkynes' difunctionalizations. Initially, copper catalyzed the cross-coupling to form an allenic intermediate, followed by a copper-catalyzed stereoselective boration reaction with B_2pin_2 . When diazo compounds **53** were used as carbene precursors, the steric interaction forced the boron group to attack the β -carbon from the opposite side of the γ -phenyl group on the allenic species **107**, leading to the favored (*Z*)-isomers **108** as major products. Whereas, in the case with *N*-tosylhydrones **51** as carbene precursors, the addition of Cu-Bpin complex to

corresponding allenic species **109** provided allyl copper intermediate, which was more favored to form a six-membered ring transition state with the association of MeOH, finally furnishing the more thermodynamically stable (*E*)-products **110** (Scheme 27).



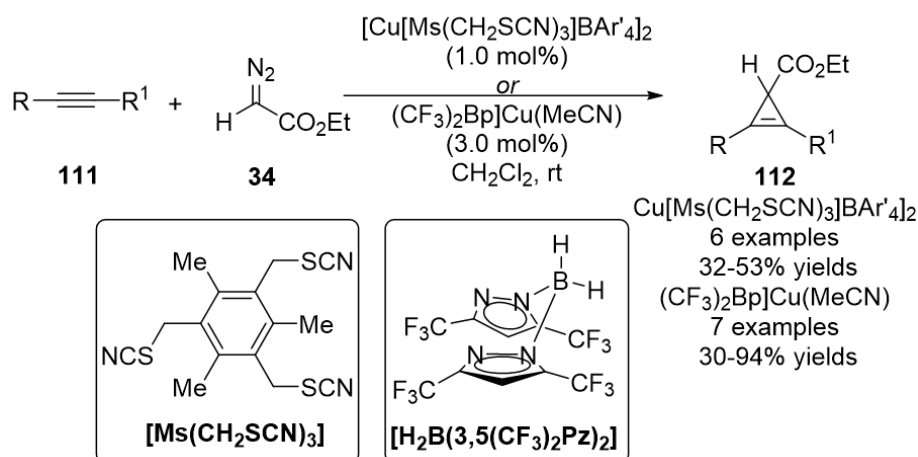
Scheme 27. Copper-catalyzed allenylation followed by boroalkylation.

3. Copper Carbene Intermediate Addition onto C-C Triple Bond

3.1 Cyclopropanation

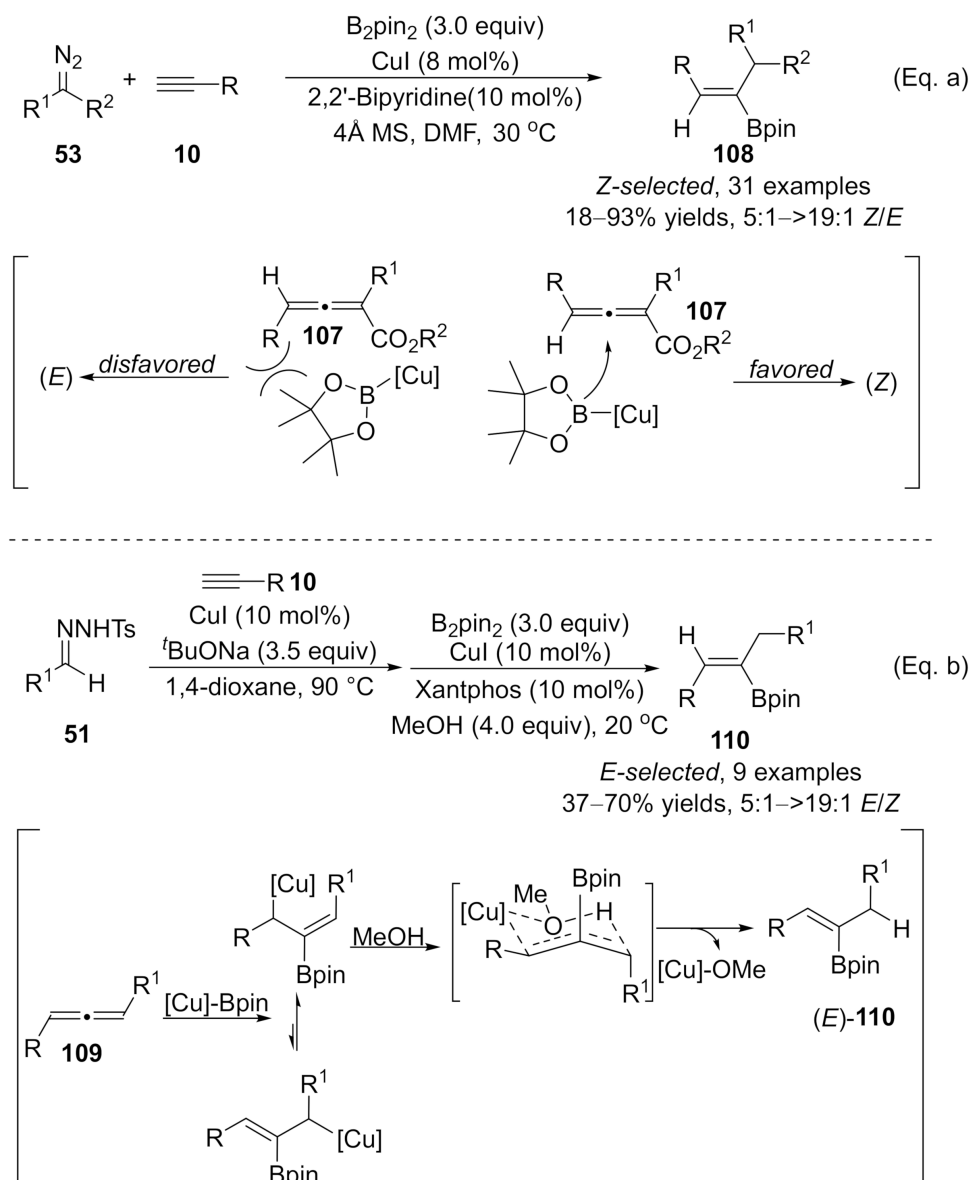
Cyclopropanation is a well-known reaction of metal carbene intermediate with alkynes. This widely used reaction could be catalyzed by rhodium [45][46][47][48][49], cobalt [50], gold [51], silver [52][53] and many others [54][55][56]. Herein, selected examples related to copper catalysis will be discussed.

In 2010, a new tridentate coordination copper complex, $\text{Cu}[\text{Ms}(\text{CH}_2\text{SCN})_3]\text{BAR}'_4$ (BAR'_4 = tetra(3,5-bis(trifluoromethylphenyl)borate), was designed by Miguel and co-workers by using $[\text{Cu}(\text{OTf})_2]\cdot\text{C}_6\text{H}_6$ and an alkylthiocyanate ligand [57]. This catalyst promoted the cyclopropenation of ethyl diazoacetate **34** (EDA) with a wide range of internal alkynes **111**, providing cyclopropenes **112** in moderate yields (Scheme 28). The same cyclopropenation work was achieved by Dias, unique bis(pyrazolyl)borate ligand supported $[(\text{CF}_3)_2\text{Bp}]\text{Cu}(\text{NCMe})$ catalyst was used, yielding cyclopropene products in moderate to high yields [58].



Scheme 28. Copper/alkylthiocyanate complex catalyzed cyclopropenation.

In 2016, a Cu(I)/*N*-heterocyclic carbene (CuNHC) complex catalyzed cyclopropenation of internal alkynylsilanes **113** with diazoacetate **15** was reported by Coleman's group [59]. A series of 1,2,3-trisubstituted and 1,2,3,3-tetrasubstituted cyclopropenylsilane compounds **114** were isolated in moderate to good yields (Scheme 29). An interesting regioselective and chemodivergent reaction pathway occurred furnished a *tetra*-substituted furan through an intramolecular cyclopropene ring-opening transformation in the case of electron-rich diazoacetate.



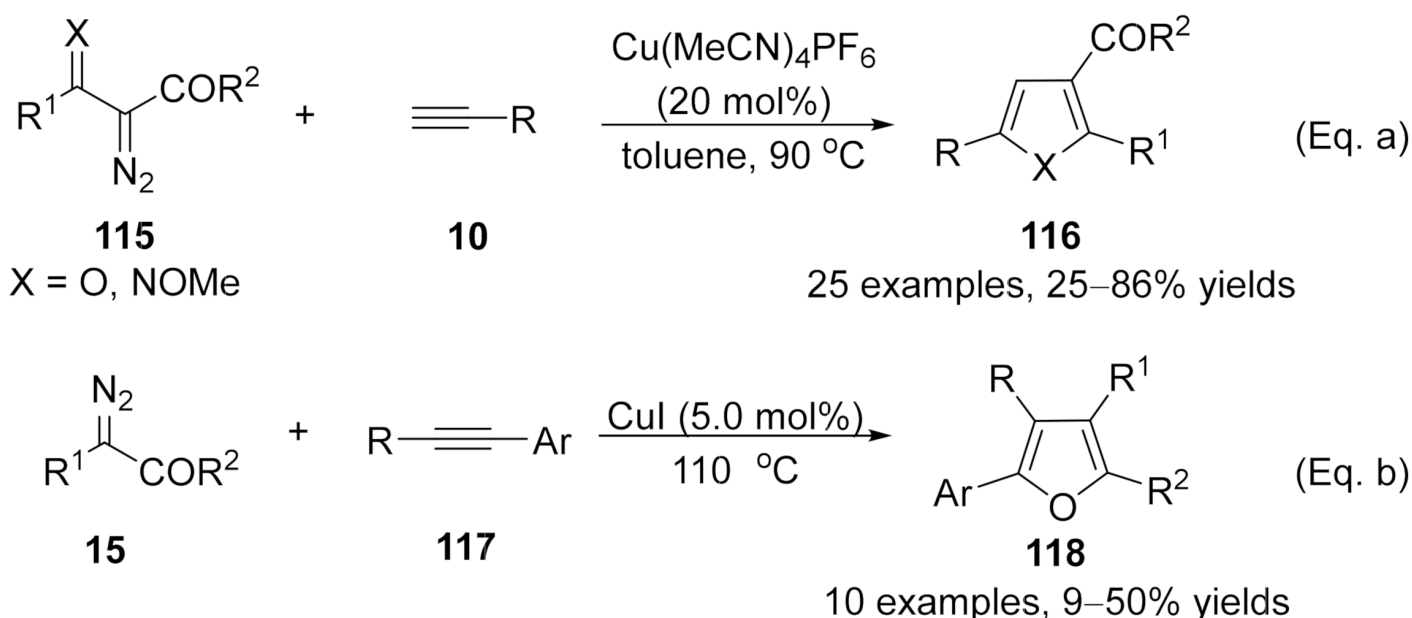
Scheme 29. Copper/N-heterocyclic carbene (CuNHC) complex catalyzed cyclopropanation.

3.2 Cascade Reaction Involving Carbene/Alkyne Metathesis Process

Carbene/alkyne metathesis (CAM) refers to the processes where a metal carbene reacts with an alkyne, generating a new vinyl metal carbene intermediate, which was difficult to access with other carbene precursors [60] [61] [62]. This *in situ* generated vinyl metal carbene intermediate could be involved in typical metal carbene reactions, such as [3+2]-cycloaddition [63], cyclopropanation [64] [65] [66], C-H bond insertion [67] [68] [69] [70], and others [71] [72] [73].

It's a general protocol for the synthesis of furan derivatives through transition metal-catalyzed formal [3+2] cycloaddition of α -diazocarbonyl compounds with alkynes [74] [75] [76] [77]. However, the cases under copper carbenes mediated were limited. In 2014, Wang's group developed a copper-catalyzed formal [3+2] cycloaddition reaction of terminal alkynes with β -keto α -diazoesters **115** (X = O), offering an operationally simple and applicable method for the synthesis of trisubstituted furans **116** (X = O) with a wide substrate scope (Scheme 30a) [78]. This reaction has

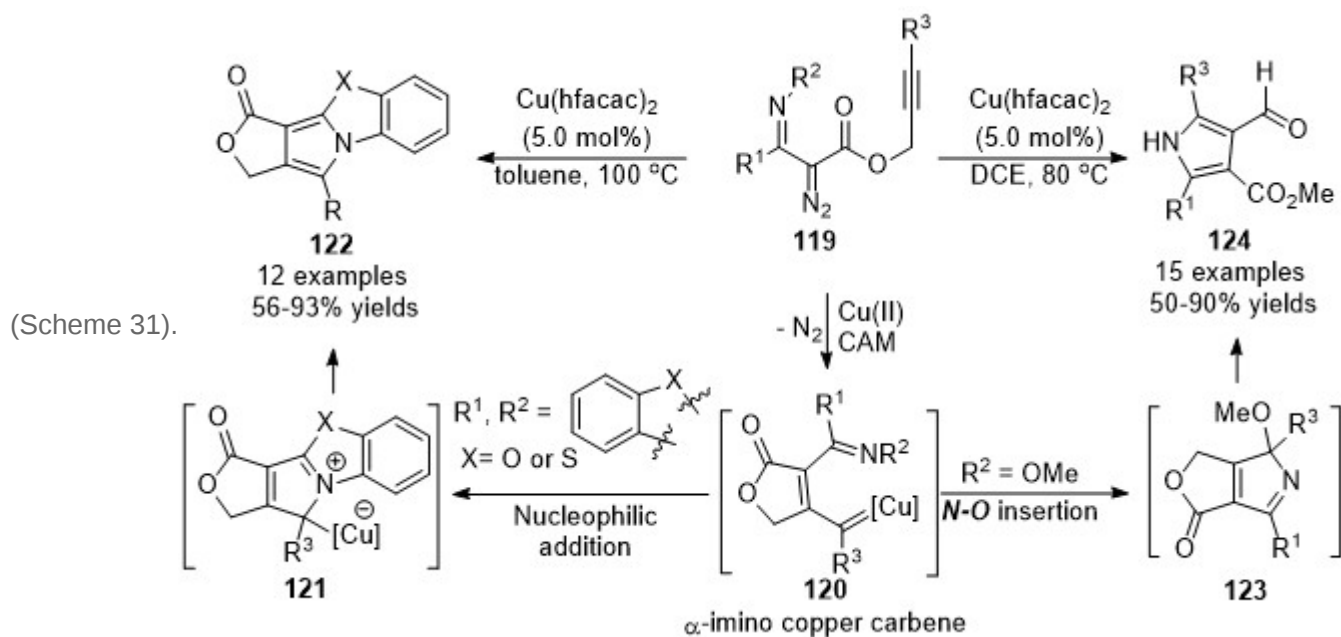
also been applied to ethyl (*E*)-2-diazo-3-(methoxyimino)butanoate **115** (X = NOME) for the synthesis of 2,3,5-trisubstituted *N*-methoxypyrroles (X = NOME). Later in 2016, a Cu(I)-catalyzed cycloaddition of diazoacetates **15** with electron-rich internal aryl alkynes **117** was discovered by Coleman and co-workers [79]. Tetra-substituted furans **118** were generated in moderate isolated yields with high chemoselectivities and regioselectivities (Scheme 30b).



Scheme 30. Copper-catalyzed carbene/alkyne metathesis for the synthesis of furan derivatives.

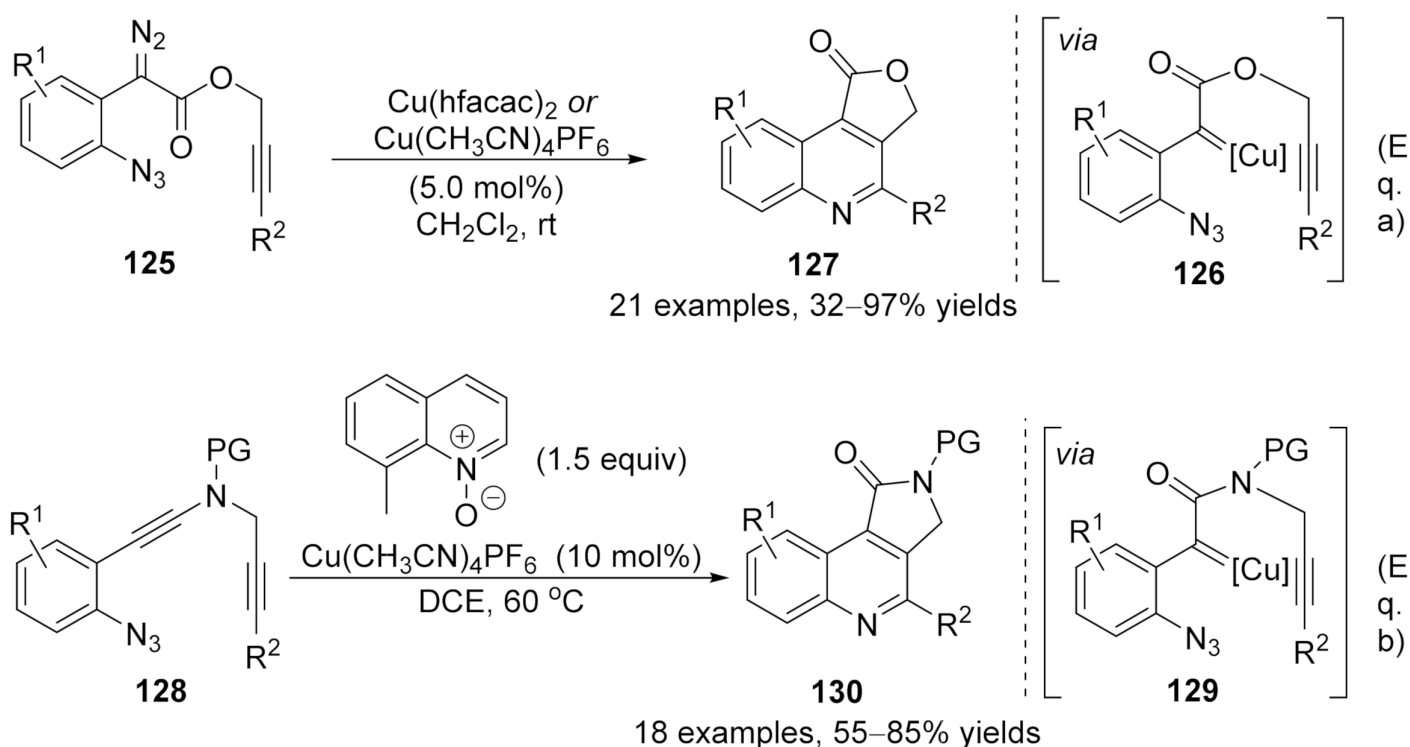
In 2016, Xu's group developed a chemo-divergent copper-catalyzed cascade reaction of alkynyl-tethered α -iminodiazooacetates **119**, providing polycyclic and multi-substituted pyrroles in high yields with a broad substrate scope [80]. Especially, the *tetra*-substituted 3-formylpyrroles **124**, which were difficult to access by alternate approaches. Mechanistic studies indicated that the α -imino carbene **120** is the key common intermediate in this divergent reaction, which was generated by metal-catalyzed carbene/alkyne metathesis of the alkynyl-tethered diazo compounds **121**. When R¹, R² was imbedded with an aromatic ring, polycyclic pyrroles **122** were formed as the major products through a [3+2]-cyclization and aromatization process. Whereas, substrates with a methoxy group on the nitrogen (R² = OMe), the carbene intermediate underwent an *N*–O insertion/alkoxy

migration/alcoholysis sequence, giving the 3-formylpyrrole products **124** in generally good to excellent yields



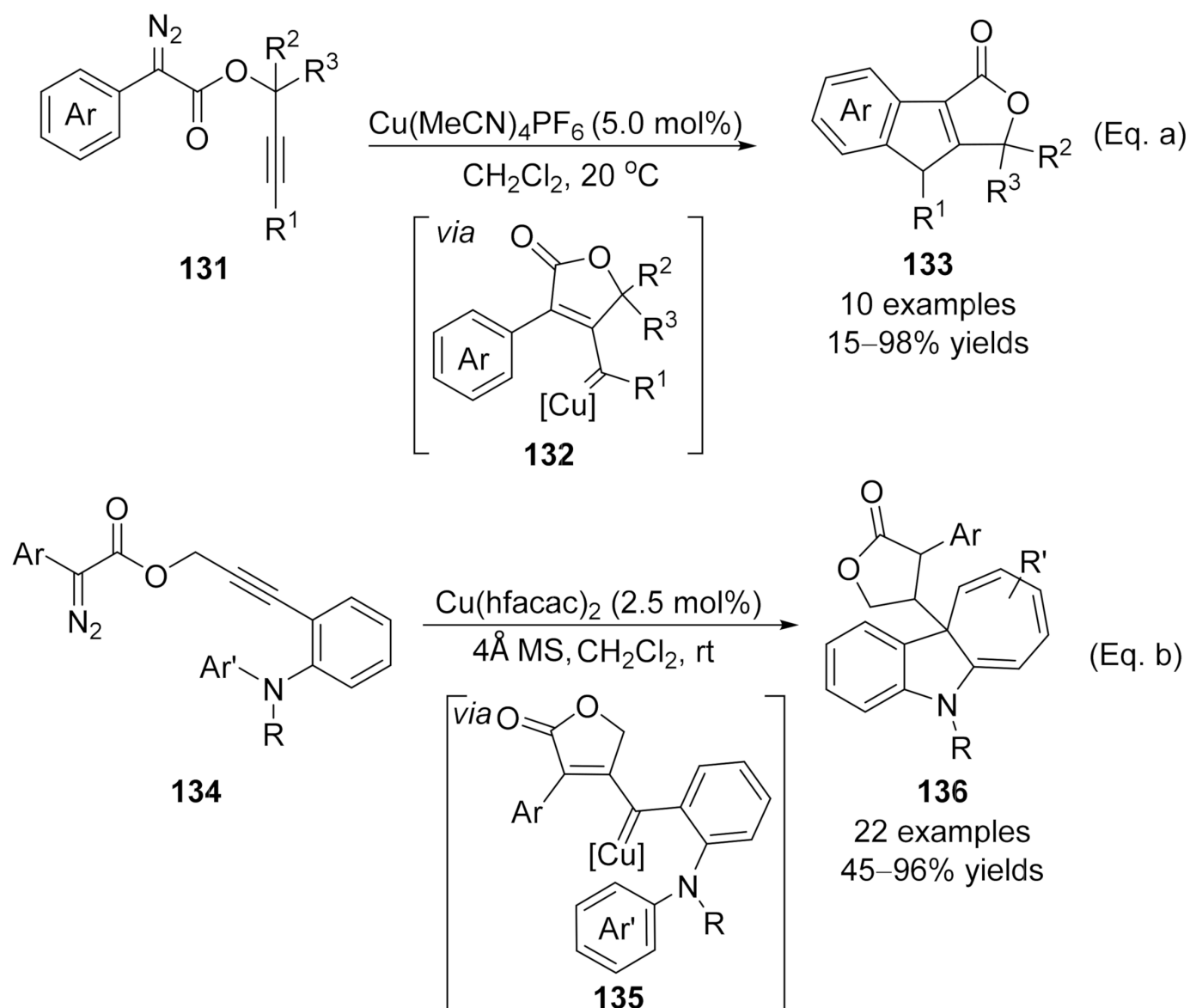
Scheme 31. Copper-catalyzed carbene/alkyne metathesis for the synthesis of pyrroles.

At the same time, Xu and co-workers have also developed a copper-catalyzed carbene/alkyne metathesis cascade reaction with alkyne-tethered diazo compounds **125** [81]. This transformation provided rapid access for the construction of multi-substituted 4-carboxyl quinoline derivatives **127** in high to excellent yields. In this cascade reaction, one $\text{C}=\text{N}$ and one $\text{C}=\text{C}$ bond were formed with the assistance of the copper catalyst under mild reaction conditions (Scheme 32a). Later in 2017, Ye's group reported an analogous protocol by using ynamides **128** as carbene precursor [82]. In this work, a copper carbene was generated in situ through a catalytic oxidation process in the presence of quinoline *N*-oxide, followed by a CAM process and terminated by carbene reaction with an embedded azide group, providing a wide range of pyrrolo[3,4-*c*]quinolin-1-ones **130** in good yields. Those works represented practical methods for the dual functionalization of alkynes (Scheme 32b).



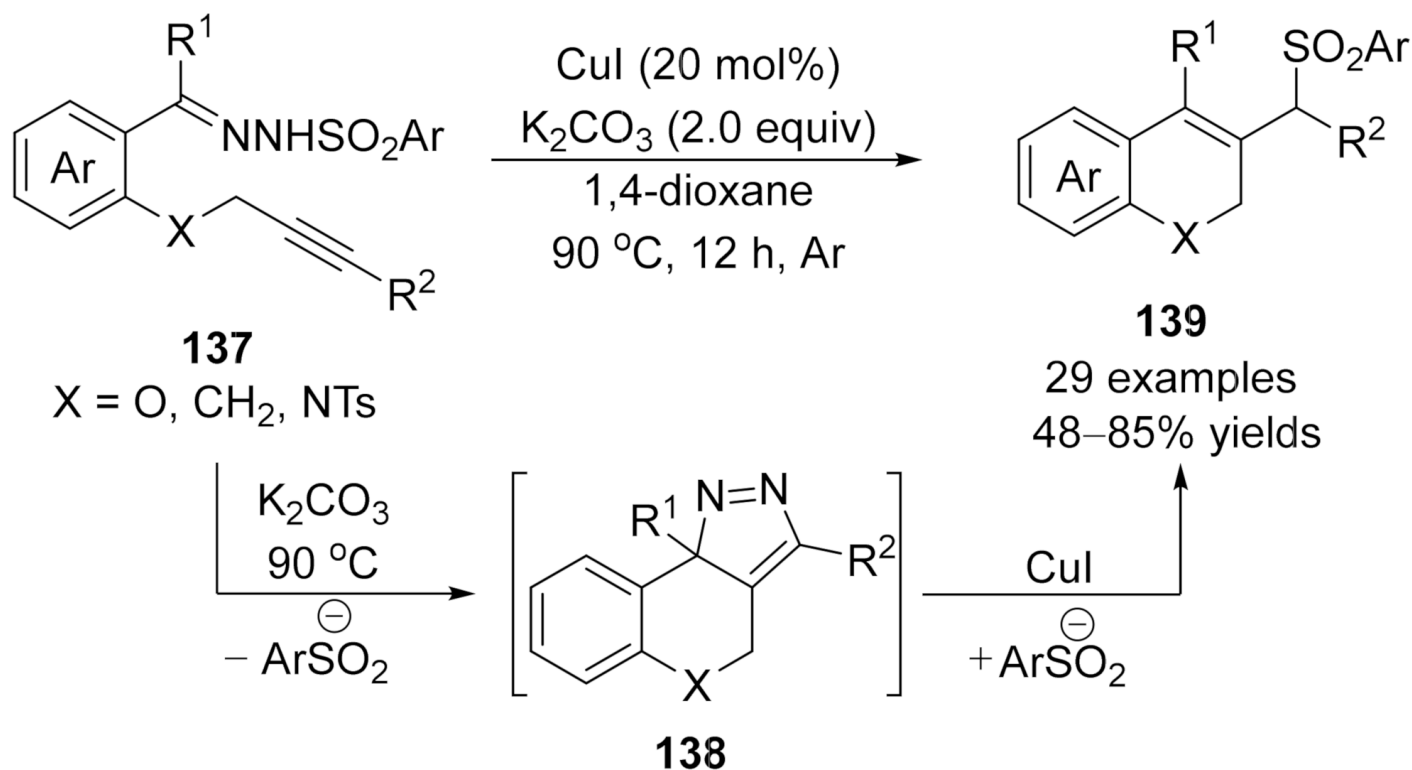
Scheme 32. Copper-catalyzed carbene/alkyne metathesis for the synthesis of quinolines.

In addition to the nucleophilic addition of the *in situ* formed copper carbene intermediates, electrophilic aromatic substitution or C(sp²)–H bond functionalization is another useful terminating transformation for the direct construction of polycyclic fused frameworks. In 2017, Doyle's group reported a copper-catalyzed intramolecular cascade reaction of diazo compounds **131**, this transformation went through a CAM process followed by a carbene C(sp²)–H bond functionalization cascade, yielding the fused indeno-furanone derivatives **133** in excellent yields under mild reaction conditions (Scheme 33a) [83]. Instead of terminating reaction in C–H functionalization, a selective Buchner insertion reaction occurred as the terminating step in Xu's work when the *ortho*-aniline substituted propargyl diazoacetates **134** were employed, selectively affording the dihydrocyclohepta[*b*]indole derivatives **136** in moderate to high yields (Scheme 33b). Notably, this reaction described a rare example of the Buchner reaction with donor/donor type metal carbene species [84].



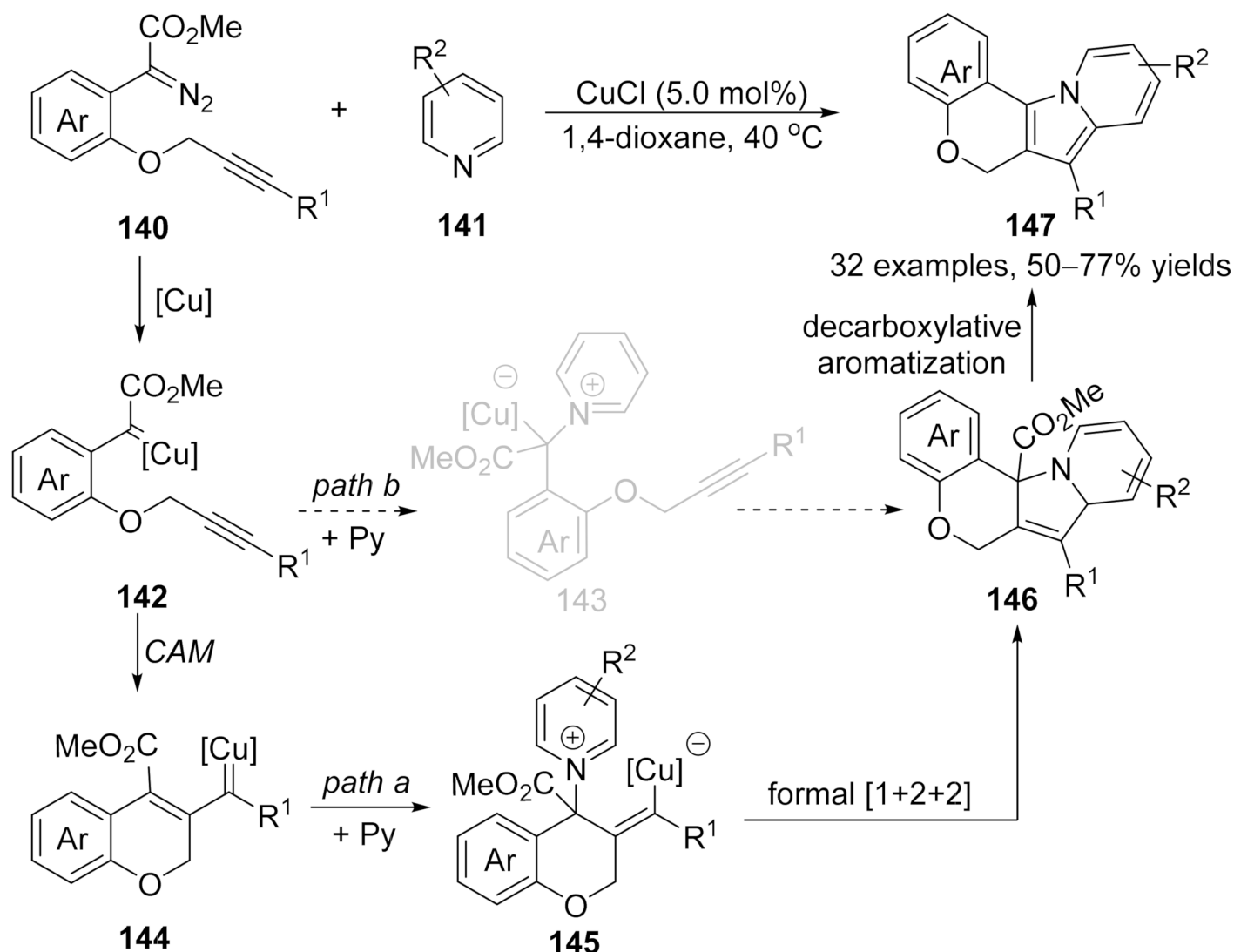
Scheme 33. Copper-catalyzed carbene/alkyne metathesis for the synthesis of tri-cyclic molecules.

In 2018, Xu and co-workers developed an intermolecular copper-catalyzed formal CAM process ^[85], which underwent a copper promoted [3+2] cycloaddition/dinitrogen exclusion/nucleophilic addition process, providing a direct and effective access to 2*H*-chromene derivatives **139** in generally good to high yields. Mechanistic studies indicated that the 3*H*-pyrazole **138** is the key intermediate in this cascade transformation, and this critical intermediate was isolated and confirmed by single-crystal X-ray diffraction and spectroscopy analysis for the first time (Scheme 34).



Scheme 34. Copper-catalyzed formal carbene/alkyne metathesis for the synthesis of 2H-chromene derivatives.

Based on a similar protocol, a copper-catalyzed formal [1+2+2]-annulation of alkyne-tethered diazo compounds **140** with pyridines **141** has been reported by Xu's group recently [86]. In contrast to the previously reported cascade reaction that was terminated the copper carbene intermediate on the carbonic center, a vinylogous addition of vinyl carbene intermediate with pyridine derivatives was occurred in this reaction, followed by an intramolecular annulation to form cycloadducts **146**, which underwent a decarboxylative aromatization process to form the desired polycyclic fused indolizine derivatives **147** in good to high yields (Scheme 35, path a), although direct formal [3+2]-cycloaddition via pyridinium ylide pathway could not be ruled out so far (Scheme 35, path b).



Scheme 35. Copper-catalyzed formal carbene/alkyne metathesis for the synthesis of polycyclic indolizines.

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