Copper mediated carbometalation reactions†

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Since the first discovery of carbocupration of alkynes in the 1970s a tremendous amount of research has been carried out in this field. The exceptionally high selectivities obtained attribute to the great synthetic value of carbocupration reactions. This tutorial review will present the most important features of carbocupration of alkynes and highlight the most relevant reviews. Then a comprehensive review of copper mediated carbometalation of cyclopropenes will follow. The latter method has received much attention over the last decade as it allows the highly selective construction of poly-substituted cyclopropanes which can be transformed into acyclic derivatives bearing one or multiple tertiary or quaternary carbon stereocenters.

1. Introduction

In the last 50 years an enormous number of stereoselective reactions have appeared. Literally thousands of chiral catalysts, auxiliaries and reagents were invented and are used today with great success. In this context, the stereoselective synthesis of double bonds, which initially received little attention since the historical Wittig- or Julia olefination reactions, became a vibrant field of research.1 The most recent contributions of stereoselective cross-metathesis and transition metal catalysed cross coupling reactions are noteworthy with the latter being dependent isomerically on pure and configurationally stable alkenyl coupling partners.2–4 With the advent of copper mediated carbometalation reactions of alkynes, it soon became obvious that this reaction constitutes one of the best approaches to generate polysubstituted olefins with a defined substitution pattern. In particular, tri- and tetra-substituted alkenes can be prepared with very high stereochemical purity.5 More recent research focuses on translating this exceptional selectivity from the two dimensional space (E/Z-selectivity) into the three dimensional space (creation of stereocenters) by replacing the alkyne substrates with alkenes.6 The copper mediated carbometalation reaction of strained alkenes provides stereoselective access to cyclic- and acyclic stereocenters, which are challenging to synthesize using traditional methods.

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Given many reviews on carbometalation of alkynes, this review will only provide a short introduction on carbocupration of alkynes. The main intention in the short overview below is to guide the readers to the most relevant reviews with a special emphasis on the basic principles governing the copper-mediated carbometalation reactions. Then a comprehensive review of copper mediated carbometalation of strained alkenes, mainly cyclopropenes, will follow. The nucleophilic species is shown in blue whereas the electrophile is given in red (abbreviated E⁺). If the electrophile is a proton, the hydrogen atom in the product is not shown for the sake of clarity.

2. Short overview of the carbocupration reactions of alkynes

Gilman cuprates (R₂CuLi), formed by the reaction of two equivalents of organolithium reagent with one equivalent of organocopper salt, react with monosubstituted alkynes through deprotonation in tetrahydrofuran. In 1971 Normant and Bourgain found that a less basic ethyl copper reagent, prepared from an equimolar amount of EtMgBr and CuBr, reacted with n-hexyne in low-polar solvents such as diethyl ether by regio- and stereospecific syn-addition to give, after treatment with iodine, pure (E)-iodoolefin 2 (Scheme 1). Throughout this review, the [Cu] notation will be used to indicate a copper metal bearing ligands.

Three key features of this reaction are remarkable: (a) carbometalation is faster than deprotonation; (b) the excellent regio- and cis-stereoselectivity; and (c) synthetic usefulness of the in situ generated copper reagent 1 which can be trapped by a large variety of electrophiles including transmetalation reactions with various metals to catalyse subsequent cross-coupling reactions. Importantly many factors such as the presence of salts, nature of solvents, temperature as well as steric effects of the organometallic species and alkyne can greatly influence the outcome of the reaction. It is noteworthy that the most simple alkyne, acetylene, does not show deprotonation with Gilman cuprates but carbocupration reactions. The carbometalation of acetylene was intensively investigated and its usefulness culminated by the total synthesis of many natural products containing Z-alkenes. Comprehensive reviews by Alexakis, Normant and Lipshutz as well as a more recent book chapter by Chemla and Ferreira discuss in detail the diverse effects of carbocupration reactions with various types of alkynes. Yorimitsu also reviewed the transition-metal catalysed carbomagnesiation and carboxinzation of alkynes. The directional effect of heteroatoms attached to the alkyne on the carbometalation reaction is particularly interesting and therefore has stimulated much research efforts in the last few decades. While dialkylsubstituted alkynes typically react inefficiently with organocupper reagents, mono- or disubstituted alkynes bearing heteroatoms or functional groups may react smoothly. Alkynes substituted with a directing (or alternatively blocking group) X in general afford the linear (or the branched product). Branched and linear refer to the hydrolysed products where R² = H. Bold bonds were used to emphasize the linear or branched shape of the alkenes. Hetero-substituted alkynes lead to the linear or branched product depending on the nature of X. Both cases shown in Scheme 2 were covered by several excellent reviews.

Scheme 3 illustrates several examples for the directing effect of functionalized alkynes (a) or α-hetero-substituted alkynes (b). The first example shows the effect of a chelating tertiary metal alcoholate to direct the copper into the α-position to afford the linear product 3 (eqn (1), Scheme 3). When the alcohol is slightly further away and protected by a TMS group, it acts as a blocking group providing the normal branched product 4 (eqn (2)). Eqn (3) shows the exceptional but highly important case of primary propargylic alcohol which undergoes unexpected trans allyl-addition to provide the branched product 5 (eqn (3)). However, when the propargylic alcohol is transformed into a leaving group (OLG, e.g. OAc) then propargyl displacement is usually obtained providing the corresponding allene 6 (eqn (4)). It is noteworthy that carbocupration reactions with methyleopper or cuprate reagents usually suffer from low efficiency. Negishi was able to solve this important problem by developing a well-known zirconium-catalysed carboalumination reaction. Eqn (5) shows that the directing effect of a functional group outweighs the effect of heteroatom substitution. The carbonyl group of oxazolidinone 7 directs the copper, in a low polar solvent, into the α-position and overwrites the tendency of the nitrogen substituent to direct the substituent into the z-position. The highly regioselective reaction affords chiral alkenyl copper reagent 8 as a reactive intermediate which was selectively oxidized with t-BuOOH into the β,β’-stereodefined trisubstituted copperenolate 9 as a single isomer. When treated with a proton source, imide 10 was obtained with high diastereoselectivity.

Allenies, which are less reactive than alkynes, can also undergo carbocupration reactions. The presence of three possible centers where carbometalation may proceed makes the regioselective carbocupration much more difficult. A recent enantioselective example with a strongly biased allene is shown in Scheme 4. Mechanistic studies by Nakamura concerning the carbometalation of acetylenic carbonyl derivatives showed that in
this case one cannot distinguish anymore between 1,4-addition reaction and carbometalation reaction because both reaction pathways proceed via the same intermediate. Based on the pioneering work of Iwai and Konotsune for the addition of organocopper reagents to acetylenic acids, several research groups investigated the E/Z selectivity of these reactions in more detail. It was shown that at low temperatures the cis-addition product could be obtained with very high selectivity (Scheme 5).

### 3. Carbocupration reactions of cyclopropenes

#### 3.1 Introduction

Before introducing the topic of copper mediated carbometalation of cyclopropenes it should be noted that several excellent reviews have been published concerning the synthesis and reactivity of cyclopropenes and alkyldene cyclopropanes. In particular recent reviews of Nakamura, Rubin and Gevorgyan, Fox, Marek cover the carbometalation reactions of cyclopropenes. Several reviews have appeared concerning the strain energy and geometry of cyclopropenes. Cyclopropenes show similar reactivity compared to alkynes. For example the heat of hydrogenation for the conversion of cyclopropene to cyclopropane is ~54 kcal mol⁻¹ and is considerably larger than that for the conversion of acetylene to ethylene (~42 kcal mol⁻¹) (Scheme 6). The true nature of the strain in these compounds is very complex as theoretical studies by Wiberg, Borden, Kass and Bach have shown. However, this simplified comparison suggests the following two principles: (a) reactions which are possible with alkynes should also be applicable to cyclopropenes. (b) Reactions with cyclopropenes, if carried out under identical conditions, should require less harsh conditions compared to alkynes due to the release of additional 12 kcal mol⁻¹ of strain. A closer look into the literature reveals that these two principles apply in many cases. Scheme 6 describes the copper catalysed formal hydroboration and the rhodium catalyzed hydroformylation reaction which were carried out on alkynes and cyclopropenes under very similar conditions. Even though reaction conditions are not identical, both reactions suggest that the reaction with cyclopropenes can be carried out at lower temperatures, lower catalyst loading or reduced reaction times. In both cases the reaction with cyclopropenes creates a new stereogenic center. Tortosa and Rubin were able to find chiral copper or rhodium complexes which afforded the corresponding enantioenriched cyclopropanes.

Given the similarity of cyclopropenes and alkynes, carbocupration of cyclopropenes should be a feasible reaction. Furthermore, in analogy to alkynes a functional group incorporated in the
Cyclopropenyl scaffold should be able to direct the carbo-
metalation reaction selectively from one face. The obtained
metalated cyclopropane would be a useful reagent, which could
undergo many subsequent reactions: trapping with electro-
philes, copper- and palladium catalysed carbon–carbon bond
forming reactions, oxidation and many others (Scheme 7).
All these assumptions based on the analogy with alkynes
have shown to be true and the herein described overview of
copper mediated carbometalation of cyclopropenes will illustrate
very well the similarity of cyclopropenes and alkynes. Never-
thless, it is important to stress that this comparison cannot
always be applied. Cyclopropenes, due to their highly strained
character, are more prone to decomposition and side reactions
as compared to their alkylene counterparts, especially when
exposed to strong Lewis acids. The following tutorial review
first discusses the addition of stoichiometric amounts of organo-
copper reagents to cyclopropenes. Next, the copper catalysed
carbomagnesiation and carbozincation reaction will be described.
Finally, carbometalation reactions followed by elimination reac-
tions, oxidation and cyclopropane ring opening will be presented.
The review will not cover carbometalation which does not involve
copper in stoichiometric or catalytic amounts. Copper mediated
addition reactions with non-carbon nucleophiles are also not
included in this comprehensive account. These types of addi-
tion reactions with cyclopropenes are reviewed elsewhere.
It is important to mention that when describing syn- and anti-
addition to cyclopropenes, we refer to the facial selectivity as
regards to a functional group (or any group) present in the
cyclopropenyl substrate. The words diastereo-selectivity and
facial-selectivity also refer to the syn- or anti-selectivity. In all
the described examples the addition of the metal–carbon bond
across the cyclopropene double bond proceeds in an exclusive
cis-fashion and will therefore not be mentioned anymore.
The regioselectivity of the addition reactions is explained in
Scheme 9 and refers to the way the carbon–metal bond is added
across the double bond. All the different types of selectivities
are depicted in Fig. 1.

3.2 Carbocupration reactions with organocuprate or
organocopper reagents
In 1985, Negishi reported the first carbocupration of cyclo-
propenes (Scheme 8). As previously discussed for the reac-
tions displayed in Scheme 6, he noticed that carbometalation of
cyclopropenes proceeds indeed faster compared to the corres-
ponding alkynes. In a competition experiment of alkyne 13 and
cyclopropene 11 only cyclopropane 12 and unreacted alkyne 13

Scheme 6 Comparison: alkynes and cyclopropenes.

Scheme 7 Carbocupration of cyclopropenes.

Scheme 8 First reported carbocupration reaction of cyclopropene by Negishi.
were obtained. This implies that cyclopropene 11 reacted much faster compared to alkyne 13. The high regioselectivity can be explained by the $\alpha$-effect of silicon, stabilizing the intermediate tertiary copper reagent $\alpha$ to the TMS-group.

The preference for the most stabilized cyclopropyl metal reagent is a general rule that dictates the regioselectivity of the carbometalation reactions of cyclopropenes. Two cases have to be distinguished: (1) monosubstituted cyclopropene (A, Scheme 9) leads to product B exclusively due to the greater stability of the secondary compared to the tertiary organometallic reagent C. However, C may also be observed in some cases when R is an anion stabilizing group (e.g. phenyl as shown in Scheme 21). (2) For disubstituted cyclopropenes of type D mixtures of cyclopropyl metal compounds E and F are observed. However, if substituent R is an anion stabilizing group (e.g. TMS, Scheme 8) then product F becomes major or even an exclusive product of the reaction.

In 1988 Nakamura and co-workers examined for the first time the carbocupration of cyclopropene ketals. They found that cyclopropene ketals, when treated with an organocuprate or heterocuprate, efficiently underwent a carbocupration reaction to afford cyclopropyl copper species 14 which was reported to be stable at temperature lower than $-50 \, ^\circ C$. Subsequent reactions with electrophiles, or palladium catalysed cross-coupling reactions gave the corresponding cyclopropane products in good yields. Palladium catalysed cross coupling reaction with (E)-1-iodohex-1-ene afforded 15, which immediately underwent divinylcyclopropane–cycloheptadiene rearrangement to afford cycloheptadienone ketal 16 in 67% yield. In general, the cyclopropyl anion is configurationally stable unless the anion is substituted by an electron withdrawing group (Scheme 10).

Two years later, in 1990, Nakamura reported the stereo- and regioselective carbocupration of cyclopropenes (Scheme 11). Protection of the cyclopropenone with a chiral ketal moiety transmitted successfully the stereoinformation during the carbocupration reaction. Interestingly, 1-substituted cyclopropene afforded the opposite diastereomer compared to the unsubstituted cyclopropene. In the case of monosubstituted cyclopropenes the reaction is also highly regio- and stereoselective: the more stable secondary cyclopropyl reagent is obtained exclusively (see Scheme 9 for explanation). The extension of the nucleophile scope to vinylic cyanocuprates and Normant cuprates ($R_2CuMgBr$) is noteworthy.

In 2007, we reported that 1-hydroxymethyl substituted cyclopropenes have the ability of direct addition of Gilman type cuprates ($R_2CuLi$) with high facial control. The high selectivity is rather surprising as the Gilman cuprates (Scheme 12) provide the opposite diastereoisomer than the Normant type cuprates ($R_2MgX$), formed from the Grignard reagent in the presence of
catalytic amounts of copper, (Scheme 23). Although the origin of this diastereodivergency is still unknown, it is interesting to note that both diastereoisomers can be obtained at will and in a good ratio by simply changing the nature of the organometallic species involved in the carbometalation reaction.

When treating cyclopropene esters with lower order cyano-cuprates \( \text{R}_2\text{CuCNLi} \) an unusual anti-addition was observed (Scheme 13). It was hypothesized that chelation does not take place, probably due to the low availability of free lithium cations in lower order cyanocuprates \( \text{R}_2\text{CuCNLi} \). Hence, no efficient chelation takes place and the ester group instead of directing the cuprate to the syn-face of the cyclopropene blocks this face and the cuprate adds from the less-hindered face, anti to the ester group.

In this special case the reaction was very sensitive to stoichiometry. The occurrence of small amounts of free alkyllithium in the presence of \( \text{RCuCNLi} \) was able to reverse the facial selectivity back to the expected syn-selectivity which is probably due to the better availability of lithium ions for chelation (Scheme 14).

syn-Selectivity was also observed for the addition of organocopper reagents (\( \text{RCuLiX} \) or \( \text{RCuMgX}_2 \)) and classical Gilman cuprate (\( \text{Me}_2\text{CuLi} \)). The depicted vinyl sulfones 16 and 17 were obtained from the corresponding alkenyl iodide 18 via an addition-elimination sequence or from the alkyne 19 via addition-hydrolysis (Scheme 15).

Furthermore, the substrate scope was extended to homologous esters and amides (Scheme 16). Despite the greater flexibility of the directing group, very high diastereoselectivity was observed with organocopper reagents. The copper-catalysed carboxymagnesiolation worked equally well.
In 2015, it was shown that lower cyano cuprates can also undergo \textit{syn}-addition with cyclopropenes bearing a weak ether-coordinating group in the 3-position (Scheme 17).\textsuperscript{64} The dramatic influence of an additional substituent at the 3-position on the facial selectivity was also reported by Fox in the context of iron-catalysed carbomagnesiation of cyclopropenes.\textsuperscript{65} This reversal of selectivity compared to the previous reaction depicted in Scheme 13 could be rationalized by the additional substituent at the 3-position of the cyclopropene. This additional substituent exerts a strong steric effect and now the weak coordination of the ether group is sufficient enough to direct the incoming organometallic reagent. The cyclopropyl copper intermediate was \textit{in situ} oxidized with retention of configuration with t-BuOOLi. It should be noted that oxygen cannot be used for this type of oxidation because organocopper reagents dimerize when exposed to oxygen.\textsuperscript{8} t-BuOOLi was found by Whitesides and Boche to be an excellent oxidizing reagent for organolithium and copper reagents.\textsuperscript{66,67} After aqueous work-up highly-substituted cyclopropanols were obtained in good yields and excellent diastereoselectivities. The starting material is easily obtained in enantioenriched form by enantioselective cyclopropenation of alkynes with methyl aryldiazoacetates in the presence of a chiral rhodium catalyst.\textsuperscript{68} Two examples were prepared in their enantioenriched form (Scheme 17).

### 3.3 Copper catalysed carbomagnesiation

Interestingly, early studies by Lukina, Rudavshevskaya and Nesmeyanova showed that cyclopropenes react with alkyl Grignard reagents to give mainly the carbometalated product.\textsuperscript{69} Alkenyl- and allylmagnesium bromides also react cleanly with cyclopropenes to afford the desired products in moderate to good yields.\textsuperscript{70} Lehmkull utilized the potential of this chemistry for a one-pot synthesis of \textit{cis}-chrysanthemic acid (eqn (1), Scheme 18).\textsuperscript{70}

The depicted example illustrates the increased reactivity of cyclopropenes compared to disubstituted alkynes which under similar conditions do not undergo carbomagnesiation with Grignard reagents.\textsuperscript{8} Richey found that, in analogy to propargylic alcohols, the alcohol function of 3-(hydroxymethyl)cyclopropenes was able to direct the carbomagnesiation of allyl Grignard reagents with high diastereoselectivity (eqn (2), Scheme 18).\textsuperscript{71} The new carbon substituent adds to the more substituted site of the cyclopropene to avoid the generation of the less-stabilized tertiary organomagnesium reagent. Therefore, the overall process is completely regio- and stereoselective.

Other Grignard reagents employed by Richey such as alkyl-, benzyl- and aryl magnesium halides failed to react.\textsuperscript{71} With the desire to expand the scope of potential nucleophiles, Fox in 2002 reported the replacement of the hydroxyl group by a methoxymethyl acetal (MOM)-group which is well known for its strong chelating properties. After the optimization of the reaction conditions he found that the replacement of ethereal solvents by pentane afforded the addition of copper catalysed allyl- and vinyl-magnesium halide with high diastereoselectivity and good yield (Scheme 19).\textsuperscript{72}

Under these conditions, many nucleophiles could be added to 3-(hydroxymethyl)cyclopropene. Previously unsuccessful alkyl-,
alkenyl- and even alkynyl Grignard reagents underwent carbo-
metalation in good yields and excellent facial selectivity
(Scheme 20).72 Fox and co-workers also developed a cost-efficient
synthesis method providing enantiomerically pure cyclopropene
substrates on gram scale.73 After the copper-catalysed carbo-
magnesiation reaction the corresponding cyclopropanes were
obtained with high facial selectivity and complete preservation
of enantiomeric purity. Two examples were provided in their
study (20 and 21, Scheme 20).

One of the last challenging issues that remained was the
introduction of an aryl nucleophile. Standard conditions shown
in Scheme 20 afforded less than 1% of product when PhMgBr
was used as a nucleophile. After much experimentation Fox and
co-workers found that super stoichiometric amounts of electron
rich phosphine ligand P(n-Bu)3 allowed for the efficient copper-
catalysed carbo-
magnesiation with aryl Grignard reagents
(Scheme 21).74 High syn-selectivity was observed in all cases,
especially for 3,3′-disubstituted cyclopropanes providing the
corresponding products with excellent selectivity. The facial
selectivity only dropped when sterically encumbered aryl-
nucleophiles were used. Reaction with ortho-tolylmagnesium
bromide gave a 60:40 mixture of syn- and anti-diastereomers,
whereas phenylmagnesium bromide and meta-tolylmagnesium
chloride provided the syn-diastereomer with high selectivity.

As explained in Scheme 9 the regioselectivity is determined by
the stability of the intermediate cyclopropylmetal species. In
general, the organic substituent is added to the more substituted
carbon. Only in the case of phenyl-substituted cyclopropanes the
regioselectivity may be significantly lower. Following the guide-
lines provided in Scheme 9 one can easily rationalize the eroded
regioselectivity for product 22 as 23 benefits from benzylic
stabilization. The low selectivity suggests that a tertiary benzylic
carbanion is almost as stable as a secondary carbanion. For the
formation of product 24 the situation is rather simple as both
regioisomers of addition are tertiary cyclopropylmagnesium
products but only one of them benefits from benzylic stabi-
lization. Hence, only product 24 is formed.

As already described in Scheme 12, a secondary alcohol in the
1-position of the cyclopropane can lead to facial differentiation.
The organocuprate reagent generated from n-BuMgBr and copper
iodide (Normant cuprates) with high selectivity was added
(Scheme 22).61 Only when the secondary alcohol was substituted
with alkyl and aryl-substituents the diastereoselectivity was
moderate. The facial selectivity can be rationalized by consider-
ing the most stable conformer (minimization of allylic strain).

The copper-catalysed carbo-
magnesiation of cyclopropenyl
esters was also investigated. Methyl magnesium bromide in

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**Scheme 20**  High diastereoselectivities with CH2OH directing group in 3-position of cyclopropane.

**Scheme 21**  Extension of scope: aryl nucleophiles.
the presence of catalytic amounts of copper iodide gave the desired products in excellent syn-selectivity (Scheme 23). Interestingly, even in a more coordinating solvent such as THF, the syn-chelated adducts are formed with the same diastereomeric ratios. Different copper salts such as CuBr and CuCN could be used with equal efficiency. The ester moiety was not attacked by the Grignard reagents at low temperatures. In contrast, lower order cyanocuprates RCuCNLi afforded the anti-product with high selectivity (490:10; Scheme 13). It is important to note that the Cu-catalysed carbomagnesiation of cyclopropenyl ester has to be carried out at temperature below −20 °C. If the reaction is warmed up to 0 °C, a stereospecific fragmentation reaction occurs (Scheme 36).63

In 2015, Endo showed that catalytic amounts of copper salt also allowed high yielding reactions with unfunctionalized cyclopropenes (Scheme 24).75 A great number of copper catalysed carbomagnesiation reactions of aryl- and alkyl-3,3-disubstituted cyclopropenes were reported. The copper-catalysed carbomagnesiation with various Grignard reagents resulted in high yields of the corresponding cyclopropanes. Only the addition of encumbered Grignard reagents such as t-BuMgBr and isopropenyl Grignard reagent did not afford the desired products. Unsymmetrical cyclopropenes resulted in a mixture of unassigned diastereomers 25 and 26.

### 3.4 Copper catalysed carbozincation

In 2009, Fox showed that the addition of dialkyl- and diaryl-zinc reagents to cyclopropenyl esters was more efficient in the presence of catalytic amounts of copper. The use of toluene as a solvent was crucial to afford high facial selectivity (Scheme 25).76 All products were obtained in good to excellent diastereoselectivity. Fox and co-workers were also able to generate non-commercial diorganozinc reagents from Grignard reagents and ZnCl2 which successfully afforded the corresponding products 31 and 32. If cyclopropane is substituted with an alkyl substituent at the 3-position (R 1 H) improved diastereoselectivity was found (compare diastereoselectivity of 27 and 28). Surprisingly, a decrease of diastereoselectivity was found for the addition of Ph 2Zn leading to compound 29 when substituted at the 3-position (30).

In 2012, Fox reported an extension of his previous work. Several diarylzinc reagents were prepared in situ from the corresponding aryl iodides by a iodine–magnesium–zinc exchange (Scheme 26).77
An efficient protocol was developed which includes the removal of magnesium salts by dioxane, which was shown to be crucial for the diastereoselectivity of the reaction. Importantly, a wide range of functionality (nitrile, ester, nitro, alkoxy, CF₃) was tolerated. When nitro groups were present, the iodine–magnesium exchange had to be done with PhMgBr instead of the “turbo Grignard (i-PrMgBr)”.

Related cyclopropenes such as chiral and non-chiral acyloxazolidinones were also shown to direct the carbozincation with high facial and enantioselectivity (Scheme 27).

Very recently, in 2015, Endo showed that the addition of hydrazine ligand L₁ greatly improved the yield of the copper catalysed carbozincation of unfunctionalized cyclopropenes (Scheme 28). The obtained cyclopropylzinc reagents were usually hydrolysed but also functionalization with iodine or allylbromide was reported. Compound 34 was observed as a single spot by TLC but could not be isolated as a pure product.

Very recently, the authors reported the first asymmetric copper catalysed carbozincation of cyclopropenes (Scheme 29). Following the pioneering work by Nakamura and Lautens they were able to identify two chiral copper catalysts which efficiently furnished highly enantioenriched polysubstituted cyclopropanes. Several features of the method are noteworthy: (1) the combination of highly encumbered phosphine ligand (R)-DTBMe-SEGPhos with Cu(CH₃CN)₄PF₆ or CuTC (copper(I) thiophene-2-carboxylate) completely suppressed dimeric side products which appeared otherwise in significant amounts (typically 15–20%). (2) The developed method provided the products with excellent...
diastereoselectivity (in general > 95:5). The carbozincation is directed by the substituents in the 3-position and occurs on the less hindered face of the cyclopropene. In the absence of the ligand low diastereoselectivities were observed. Hence, the developed method also provides an efficient and highly diastereoselective route to racemic cyclopropanes. Only in the case where the two substituents in the 3-position lack steric bias, the diastereoselectivity decreases.

(3) The enantiomeric ratio of the obtained products is in general high (> 90:10) and in some cases even excellent (> 99:1). It is noteworthy that diaryl- and dialkyl-substituted cyclopropanes are also obtained in high optical purity. (4) The in situ generated cyclopropylzinc reagent is configurationally stable and can be further functionalized under mild conditions with a great variety of electrophiles and coupling partners. Importantly palladium-catalysed cross coupling reactions with commercially available alkenyl- and aryl bromides allows for the rapid synthesis of a library of cyclopropanes from a single cyclopropylzinc precursor.

3.5 Carbocupration followed by elimination

If one takes into consideration the similar chemistry of cyclopropanes and alkynes, the propargylic displacement shown in Scheme 3 (eqn (4)) should also apply to cyclopropanes (Scheme 30).

Indeed, in 2006 Fox and Marek reported simultaneously the diastereoselective synthesis of methylenecyclopropane (MCP) and alkylidenecyclopropane derivatives (ACP) respectively from cyclopropanes.82,83 Fox showed that cyclopropanes substituted with suitable leaving groups such as methyl, MEM- and SEM ethers gave a mixture of addition (35) and addition–elimination (36) products (Scheme 31). Further investigations showed that the presence of copper salts was not necessary to induce the elimination reaction. Changing the Grignard counter ion from chloride to bromide resulted in the exclusive formation of the addition–elimination product 36. The change in regioselectivity to form the less substituted product contrasted all prior studies on the copper-catalysed carbomagnesiation reaction of 1-alkylcyclopropanes, where the normal course of reactivity is for the nucleophile to add to the more substituted carbon-center of the double bond (Scheme 20). Fox proposed the simultaneous chelation of magnesium by the hydroxymethyl and the OMEM group (eqn (2)). Then, similar to propargylic substitution the organic substituent is delivered while the leaving group is expelled. A large excess of the respective Grignard reagent is required for efficient reactions (eqn (5) and (6)). Poorly coordinating groups such as trityl ethers afforded mainly the addition product 37 (eqn (3)). Noteworthy is that the cyclopropene
precursors are readily available in the enantioenriched form and therefore this route can provide access to the enantiomerically enriched MCP.

In 2013, Fox described the synthesis of highly substituted MCP from 1,2-disubstituted 3-(hydroxymethyl) cyclopropenes bearing a methoxy substituent as the leaving group (Scheme 32). Copper-catalysed displacement reactions with Grignard reagents proceeding in good yield and high diastereoselectivity. When substituent R1 was introduced the diastereoselectivity increased dramatically (38 and 39). Interestingly, iron catalysis furnished the anti-addition product whereas copper-catalysis afforded the syn-addition product. In parallel, it has been shown that when 1-(1-hydroxyalkyl)-cycloprop-1-ene was treated with copper salt and alkyl Grignard reagents, alkylidenecyclopropanes were obtained in good yield, high diastereoselectivity and E/Z selectivity (Scheme 34). Without the addition of copper salt, the reaction did not proceed. The absolute configuration of the starting cyclopropenylcarbinols and the final alkylidenecyclopropanes imply an overall syn SN2’ displacement of the alcohol moiety. The reaction proceeds by a four step procedure (Scheme 33): (1) deprotonation of the allylic alcohol by the Grignard reagent and subsequent chelation of the organocopper reagent. (2) Copper-catalysed carbomagnesiation reaction of the most stable conformer. (3) Transmetalation from copper to magnesium (4) subsequent β-elimination. The transmetalation from copper to magnesium (step 3) is crucial for efficient elimination reactions. If copper is used stoichiometrically, the reaction stalls at intermediate 40 and can be trapped by electrophiles (Scheme 22). This is in accordance with the previous reports which show that the carbon–copper bond is less prone to β-elimination compared to the carbon–magnesium bond. It should be noted that highly enantioenriched cyclopropenes can easily be obtained by Sharpless kinetic resolution or enzyme-mediated acylation of the racemic starting material. In a full account the generality of this method was demonstrated by providing a great number of examples (Scheme 34).

It is important to note that the counter ion of the Grignard reagent is not an innocent spectator in such displacement reactions. In some cases, the stereochemistry of β-elimination reactions (syn versus anti) significantly improved when changing from alkyl magnesium bromides to iodides. In general arylmagnesium halides reacted with lower selectivities compared to alkylmagnesium halides.
3.6 Carbocupration followed by ring opening

The similar reactivity of cyclopropene and alkyne is especially striking when cyclopropane ring-opening reactions are involved. A vinylmetal reagent generated from the carbometalation of an alkyne is well known to react with a great variety of electrophiles (eqn (1), Scheme 35). In analogy, cyclopropyl metals can be trapped with electrophiles to provide cyclopropane products. However, cyclopropanes substituted by an electron withdrawing group (EWG) possess an “internal electrophile” and can undergo ring-opening to provide acyclic products which resemble very much the products obtained via alkyne-chemistry (eqn (2)).89 The same holds true for the oxidized metal species. Alkyne chemistry and cyclopropene chemistry provide aldehydes with an alpha all-carbon quaternary stereocenter (eqn (3) and (4)).90 Obviously, if the electron-withdrawing group is replaced by a suitable leaving group, elimination occurs and an alkene is obtained (eqn (5)).

We could demonstrate the feasibility of all three pathways. First, the ring opening of cyclopropyl-magnesium species containing an ester group as an electron withdrawing group was observed upon warming the reaction to 0 °C (Scheme 36).63 The formation of the alkene is highly selective due to the pre-organized position of the substituents in the cyclopropane scaffold (eqn (2), Scheme 35).

Next, the oxidation-ring opening sequence was shown to efficiently provide the corresponding acyclic aldehydes (Scheme 37).63

Scheme 35  Cyclopropane ring-opening reactions.

Scheme 36  Thermally induced ring-opening reaction.

Scheme 37  Diastereodivergent carbometalation reaction allows for the synthesis of both enantiomers of aldehydes 42.

Scheme 38  Carbometalation–oxidation–elimination sequence.
The diastereodivergent carbometalation reaction of cyclopropenyl esters (Schemes 13–15) was cleverly exploited to access both enantiomers of the corresponding aldehyde (S)-42 and (R)-42 starting from the same cyclopropenyl ester 41 (Scheme 37). It is noteworthy that in the case of copper-catalysed carbomagnesiolation, molecular oxygen provided clean oxidation without leading to dimerized products which are typically observed with organocopper reagents.

Finally, the addition–oxidation–elimination sequence described in eqn (5), Scheme 35 was realized (Scheme 38). Acetate was found to be the best leaving group. As previously discussed for Scheme 17, substituent R2 is necessary to block one face of the cyclopropane allowing the high observed diastereoselectivity. Two aldehydes were prepared in their enantioenriched form starting from enantioenriched cyclopropenes. It has been noticed that the corresponding aldehydes are prone to decomposition. Very likely oxidation of the aldehyde into the corresponding acid occurred. It is well known that aldehydes with a high degree of substitution are prone to aerobic oxidation.91

4. Conclusions

Copper mediated carbometalation remains one of the best methods to construct carbon–carbon bonds with high selectivity. The product, itself an organometallic reagent, can enter a manifold of subsequent reactions such as electrophile trapping but also oxidation and carbon–carbon bond cleavage. Most of the past research focused on the highly selective construction of E/Z alkenes. Nowadays the high syn-selectivity and the possibility of employing substrate controlled carbometalation have opened new possibilities for the highly stereoselective construction of tertiary and quaternary stereocenters. Finally, the first highly enantioselective copper-catalysed carbometalation was shown which is the most recent way to synthesize non-functionalized enantioenriched cyclopropanes.

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Notes and references