Transition metal-catalyzed \([2 + 2 + 2]\) cycloaddition of nitrogen-linked 1,6-diynes: a straightforward route to fused pyrrolidine systems

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Transition metal-catalyzed \([2 + 2 + 2]\) cycloadditions of nitrogen-linked 1,6-diynes with unsaturated motifs, such as alkynes, alkenes, nitriles, ketenes and isocyanates, have recently attracted more attention from synthetic organic chemists because of their high efficiency in the construction of numerous pyrrolidine based systems. Utilizing different chiral transition metal catalysts to access challenging chiral skeletons has been extensively explored in recent years. We hope to highlight the power of \([2 + 2 + 2]\) cycloaddition chemistry to access a variety of fused pyrrolidine structures from inexpensive and easily available nitrogen-linked 1,6-diynes. The literature has been surveyed until the end of 2016.

1. Introduction

The construction of carbo- and heterocyclic skeletons remains a fertile active area of research because of their ubiquitous occurrence in natural products and pharmaceutical agents.\(^{2}\) Transition metal-catalyzed \([2 + 2 + 2]\) cycloadditions of unsaturated substrates, such as alkenes, alkynes, nitriles, imines, and isocyanates, provide a straightforward route to the synthesis of highly substituted carbo- and heterocyclic compounds in one step with ideal atom economy (all the atoms present in the starting materials are present in the product).\(^{3}\) In addition, enantioselective cycloaddition using chiral transition metal catalysts is a powerful synthetic tool for the construction of chiral cyclic frameworks.\(^{4}\) The recent growth in \([2 + 2 + 2]\) cycloaddition reactions has led to a large number of excellent reviews on this chemistry.\(^{3,4}\) However, most of them focused on the general aspects of the field and organization of the reactions based on starting materials and desired products (to give the

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opportunity to compare the efficacy of procedures) are usually ignored. The literature review reveals that alkynes are the most popular substrates in this chemistry and 1,6-diyne is the most common type of them. Despite great popularity of these substrates in [2 + 2 + 2] cycloaddition reactions, no review in the literature is available covering different aspects (scope, limitations, advances, and so forth) of 1,6-diyne in [2 + 2 + 2] cycloaddition chemistry. Thus, considering the lack of such a review in the literature and in continuation of our interest on N-propargylamines chemistry, here we will focus exclusively on transition metal-catalyzed [2 + 2 + 2] cycloaddition of nitrogen-linked 1,6-diyne. The review is organized based on the desired products (e.g. isoindolines, 2,3-dihydro-1H-pyrrolo[3,4-c]pyridine, 2,3-dihydropyranoo[4,3-c]pyrrol-6(1H)-ones, and 2,3-dihydro-pyrrolo[3,4-c]pyridin-6-imines), type of reactions (intermolecular and intramolecular), and metal catalysts (e.g. rhodium, ruthenium, cobalt, and nickel). With this review, we aim to inspire researchers to conduct more research in [2 + 2 + 2] cycloaddition of nitrogen-linked 1,6-diyne so that synthesis of important fused pyrrolidine-based natural products and pharmaceuticals through this reaction can become a reality.

2. Isoindolines

2.1. Isoindoline derivatives from nitrogen-linked 1,6-diyne and monoalkynes

2.1.1. Rhodium. Among all the available catalysts, rhodium complexes have proved the most popular and efficient reagents for the [2 + 2 + 2] cycloaddition reaction of nitrogen-linked 1,6-diyne with alkynes. This ability of rhodium complexes discovered by T. Kim nearly 20 years ago.\(^6\) In general, Rh-catalyzed [2 + 2 + 2] cycloaddition reactions have been performed in common organic solvents, water, and biphasic systems at temperatures ranging from room temp. upwards. The first example of isoindoline synthesis through this strategy in biphasic system has been reported by Kinoshita, Shirokubo, and Oshima in 2003, when N-tosylated diyne 1 underwent a cycloaddition reaction with propargyl alcohol 2 in the presence of the [RhCl(cod)]\(_2/\)tppts (tris(m-sulfonatophenyl)-phosphine)/NaCl/HCl/H\(_2\)O/Et\(_2\)O system (Scheme 1a).\(^7\) Later, Wu's group studied the scope of alkynes in this reaction. Thus, a variety of mono and disubstituted isoindolines were synthesized via selective [2 + 2 + 2] cyclization of 1 with both terminal and internal alkynes 4 using the [Rh(cod)]\(_2/\)H\(_2\)O/THF system at room temperature. The results demonstrated that the substrates with terminal alkynne unite gave higher yields than those with internal unites, and ary lacetyl ene derivatives gave higher yields than alkyl acetyl enes (Scheme 1b).\(^8\) With the objective of designing a greener procedure to isoindolines through [2 + 2 + 2] cycloaddition strategy, Tsai and co-workers were able to demonstrate that a series of 5-substituted isoindolines could be obtained from the reaction of 1 with terminal alkynes 6 employing [RhCl(cod)]\(_2/\)cationic 2,2'-bipyridyl 7 combination as the catalytic system in the most environmentally benign solvent, water (Scheme 1c).\(^9\)

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In addition to terminal diyne 1, internal diynes were also reacted with alkynes giving highly substituted isoindoline derivatives. In 2007, Clayden and Moran investigated the reaction of symmetric internal dipropargyl amines 9 with phenyl acetylenes 10 to 4,5,7-trisubstituted N-protected isoindolines 11. After studying a number of solvents, such as heptane, d-chloroform, dichloromethane, toluene, and THF, and catalyst, such as CpCo(CO)2, Rh(cod)2BF4, Pd(PPh3)4, and Pd/C, the system DCM/Rh(cod)2BF4/rac-BINAP at room temperature was found to be optimum with respect to the yield of product isolated (Scheme 2a). The authors also examined the cyclization of 9 with 2-ethoxyethyltrimethylsilane 12 under the standard reaction conditions. However, poor results were obtained (Scheme 2b).10

Roglans’s group improved the efficiency of these reactions by using rhodium complexes of N-heterocyclic carbenes Rh(cod)2BF4/rac-BINAP at room temperature. The reaction was performed in DCM with varied ligands and substrates. The reaction was found to be effective with a variety of substrates, producing high yields of products.

Scheme 1 Rh-catalyzed isoindoline synthesis from terminal dipropargyl amine and alkynes.

Scheme 2 (a) Synthesis of three substituted isoindolines 11 described by Clayden; (b) synthesis of four substituted isoindolines 13 developed by Clayden.
due to the difficulty associated with the presence of paramagnetic species which made impossible the following by NMR. Recently, the groups of A. Roglans were able to detect key intermediates in Rh(I)-catalyzed [2 + 2 + 2] cycloaddition of dipropargyl amines with alkynes by electrospray ionization mass spectrometry (ESI-MS) and DFT calculations. The authors proposed the mechanistic pathway shown in Scheme 5 that starts with the coordination of the rhodium(i) catalyst to the dipropargyl amine 1, leading to the formation of intermediate A, which undergoes oxidative addition–cyclization to furnish the rhodacycloheptadiene complex B. Its coordination to monoalkyne 6 yields complex C which undergoes monoyne insertion to give the rhodacyclo[3.2.0]heptatriene intermediate D and then the cycloheptatriene intermediate E. Its reductive elimination yields the complex F, which finally furnishes the final product 8 and regenerates the Rh(i) catalytic species.\(^\text{13}\)

2.1.1.2. Ruthenium. The regioselective Ru(II)-catalyzed [2 + 2 + 2] cycloaddition of dipropargyl amines with alkynes was first reported by Itoh and co-workers. They showed that unsymmetrical dipropargyl amine 21 underwent chemo- and regio-selective [2 + 2 + 2] cycloaddition with hex-1-yne 22 in the presence of [Cp*Ru(cod)Cl] in 1,2-dichloroethane and gave the desired product 23 in good yield and excellent regioselectivity (Scheme 6). According to the mechanistic pathway proposed by the authors, the steric repulsion between bulky Cp* ligand and butyl group in the ruthenacycloheptatriene intermediate A* is responsible for the regioselectivity (Scheme 7).\(^\text{16}\)

Inspired by this work, Witulski and co-workers compared the Ru-mediated [2 + 2 + 2] cycloaddition reaction of unsymmetrical dipropargyl amine 21 with a series of terminal alkynes 6 with Rh-mediated reaction of the same set of compounds. The results showed that the regioselectivity of the products were drastically changed by changing the catalysts. When [RuCl2(-NCHPPh)(PCy3)] was applied as the catalyst, 4,6-disubstituted isoindolines (meta) were obtained as major product, whereas when [RhCl(PPh3)3] is used the yield of 4,6-disubstituted

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**Fig. 1** Chemical structures of 14 and 15.

[\text{RhCl(IPr)(cod)}] 14 and [\text{RhCl(IMes)(cod)}] 15 as catalyst in ethanol or toluene (Fig. 1).\(^\text{11}\)

The possibility of solid-support [2 + 2 + 2] cycloaddition reaction of dipropargyl amines with alkynes in the presence of Wilkinson’s catalyst was first realized by Sun and co-workers, who synthesized a series of NH-free isoindolines 17 from resin-bound dipropargyl amines 16 and terminal alkynes 6 in good yields (Scheme 3). It should be mentioned that the internal alkynes do not give good yields under this reaction conditions.\(^\text{12}\) Later, the group of Deiters improved the efficiency of this protocol using [\text{RhCl(PPh3)3}] as the catalyst.\(^\text{13}\)

Along this line, recently Roglans and co-workers developed a beautiful rhodium–NHC hybrid silica material, which efficiency catalyzed both inter and intramolecular [2 + 2 + 2] cycloaddition of nitrogen-linked 1,6-diynes. This catalyst was prepared in two steps from silylated imidazolium salt 18: treatment of imidazolium salt 18 with [\text{Rh}[\mu-OEt][\text{cod}]]\(_2\) in anhydrous dichloromethane at room temperature for 4 hours led to the desired monosilylated rhodium complex 19 in a 91% isolated yield. The rhodium–NHC hybrid silica catalyst 20 was then obtained in high yield by grafting of 19 to the mesostructured silica SBA-15 in refluxing anhydrous toluene under N\(_2\) atmosphere over the course of 24 hours (Scheme 4).\(^\text{14}\)

The documented mechanism for Rh-catalyzed [2 + 2 + 2] cycloaddition reaction of dipropargyl amines with alkynes in intermolecular as well as intramolecular fashion is rare, possibly

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**Scheme 3** Solid-phase synthesis of isoindolines 17.

**Scheme 4** Synthesis of rhodium–NHC hybrid silica material 20.
isoindolines are decreased in favor of the 4,5-disubstituted isoindolines (ortho) (Table 1).¹⁷

Deiters groups have applied microwave heating for ruthenium-catalyzed [2 + 2 + 2] cycloaddition of polymer supported dipropargyl amines with both terminal and internal alkynes, generally reporting improved yields in shorter reaction times, with no effect on the regioselectivities.¹⁸ More recently it was found that the use of commercially available ruthenium trichloride allowed to catalyze this process at loadings as low as 2 mol% under solvent-free conditions. This system has several advantages such as environmentally friendly processes (Green Chemistry), good to high yields and number of functional groups tolerance (Scheme 8). This procedure was applied on the gram-scale without sacrificing the yield or outcome of the methodology.¹⁹ Other applications of Ru catalysts in [2 + 2 + 2] cycloaddition of di(3-iodopropargyl)amine with alkynes to afford 4,7-diodoisoindoline derivatives are also known.²⁰

Yamamoto and co-workers were able to demonstrate that a series of regioisomeric isoindolinones 28 and 29 be efficiently synthesized from the cyclization of amide-tethered diynes 27 with terminal alkynes 6 by treatment with [Cp*Ru(cod)Cl] in DCM. The results showed that the regiochemical effect of the reaction was strongly dependent on the substitution pattern of 27. Thus, when R¹ = R², the cyclization afford isomer 28 as major product, whereas when R¹ ≠ R², isomers 29 are regioselectively formed in higher yields than isomer 28 (Scheme 9).²¹ The regioselectivity of this process also depended on the solvent employed in the reaction. The use of cyclopentyl methyl ether (CPME) as the
solvent afforded better regioselectivity and a higher yield than DCM for isomers 28.

2.1.1.3. Molybdenum. Mori and co-workers showed that the reaction of dipropargyl amine 9a with terminal alkynes 4 in the presence of Mo(CO)6/p-ClPhOH as the catalytic system afforded highly substituted isoindolines 30 via formal [2 + 2 + 2] cycloaddition together with 3,4-diethylidenepyrrolidine 31 through a metathesis-type reaction. They found that the use of a large amount of alkyne 4 greatly favored the trimerization. However, pyrrolidine derivative 31 has always obtained as a by-product (Table 2). The author proposed mechanism for this transformation is depicted in Scheme 10.

2.1.1.4. Iridium. One of the latest catalysts to join the [2 + 2 + 2] cycloaddition story is the iridium(III) complex. In 2012, Ratovelomanana-Vidal et al. have shown that the Ir(m) complex $\text{[Ir(H)[rac-BINAP]}_2(\mu_1,\mu_1)]$ effectively catalyzes the $[2 + 2 + 2]$ cycloaddition of dipropargyl amines 32 to alkynes 4 via iridium-cycloheptadiene intermediates. These reactions were run in isopropyl alcohol, tolerated various functional groups, and generally provided mono, di, tri, and tetrasubstituted isoindolines 33 in good yields and high regioselectivity (Scheme 11).

2.1.1.5. Titanium. In 2014, Y. Six and co-workers reported an example of isoindoline preparation through a titanium-mediated intermolecular $[2 + 2 + 2]$ cycloaddition strategy. They found that dipropargyl amine 1 underwent cyclization with 1,2-diphenylethylene 34 in the presence of Ti(OiPr)$_4$/n-BuLi as the catalytic system in THF at 0°C. The corresponding 5,6-diphenylisoindoline 36 was obtained in yield of 49%. According to the author proposed mechanism, the key step of the reaction involves generation of titanacycloprenne intermediate 35 by reaction of in situ generated complex A with 1,2-diphenylethylene 34 (Scheme 12).

2.1.2. Achiral isoindolines via intramolecular $[2 + 2 + 2]$ cycloadditions. Isoindoline derivatives can also be synthesized by completely intramolecular fashion of $[2 + 2 + 2]$ cycloaddition reaction of three alkyne functionalities. The possibility of this reaction using triyne 37 in the presence of a Mo(CO)$_6$/p-ClPhOH catalyst was first realized by Mori and co-workers, who
synthesized highly substituted isoindolines 38 in moderate yields (Scheme 13).

A method involving the microwave-assisted intramolecular cyclization of N-(prop-2-ynyl)octa-2,7-diyn-1-amine 39 was also developed, leading to the formation of 6,7,8-trihydrocyclopenta[e]isoindoline 40 in 48% yield (Scheme 14). The reaction was performed under metal-free conditions using focused microwave heating in DMF.

Recently, an excellent method for the generation of furoisoindoline 42a and pyrrolo-isooindoline 51 from triynes 41a and 41b have been reported by Roglans and co-workers using hybrid silica material 20 as a catalyst in refluxing ethanol.
Along this line, Tanaka’s group reported the enantioselective synthesis of axially chiral biaryl diphosphonates 47 and dicarboxylate 49 by cationic rhodium(i)/Segphos complex catalyzed [2 + 2 + 2] cycloaddition of internal di(but-2-ynyl)amines 9a (and 45) with a phosphonate-ester and ester-substituted 1,3-butenadiyne 48, respectively. The desired double-annulation products were obtained in moderate to good yields and outstanding enantioselective excesses (Scheme 17). It is noted that the use of electron-deficient phosphonate- ester-substituted 1,3-butenadiyne as a cycloaddition partner is crucial for the success of this reaction.11

In 2009, Shibata and co-workers were able to synthesize the chiral tetrathienylene 51 by the consecutive inter- and intramolecular cycloadditions of two N-(3-(2 ethynylphenyl)prop-2-ynyl)-3-phenylprop-2-yn-1-aminos 50 using [Rh(cod)2]BF4 as the catalyst and Cy-BINAP as the chiral ligand (Scheme 18).12 Asymmetric Ir-catalyzed [2 + 2 + 2] cycloaddition of dipropargyl amines in both inter- and intramolecular fashions to isoindoline derivatives has also been well documented in the literature by the same research team.13-15

2.2. Isoindoline derivatives from nitrogen-linked 1,6-diynes and alkenes

2.2.1. Achiral isoindolines via intermolecular [2 + 2 + 2] cycloadditions

2.2.1.1. Rhodium. Rh-catalyzed intermolecular [2 + 2 + 2] cyclization of nitrogen-linked 1,6-diynes with olefins is a straightforward route to the isoindoline cores. One of the earliest reports of the applicability of these reactions, has been reported by Hara, Hirano, and Tanaka in 2008. They showed that intermolecular [2 + 2 + 2] cycloaddition of internal dipropargyl amine 9a with enol ethers 52 in the presence of [Rh(cod)2]BF4 as the catalyst and ras-BINAP as a biphosphine ligand in DCM led to the corresponding di- and trisubstituted isoindolines 53 in good to excellent yields (Scheme 19a). The mechanistic course of this reaction sequence is shown in Scheme 19b, and involves the initial formation of the rhodacyclopentadiene A from the reaction of diyne 9a with rhodium. Insertion of 52 into one of the Rh–C bonds gives intermediate C through intermediate D. Finally, reductive elimination of rhodium affords the observed isoindolines 53 (Scheme 19b).16,17

They also showed that treatment of diyne 9a with vinylene carbonate 54 under the aforementioned rhodium catalyst gave the corresponding 4,7-dimethyl-2-tosilisoindolin-5-ol 55 in yield of 82% (Scheme 20).18 More recently, the same authors...
extended their methodology to cyclization of dipropargyl amines with 2,3-dihydropyran to 2-(isoindolin-6-yl)ethanol derivatives.

To develop a new protocol for the synthesis of isoindoline derivatives via \([2 + 2 + 2]\) cycloaddition reaction, Tong and co-workers have investigated the cyclization of dipropargyl amines with potassium \((Z)-(2-	ext{bromovinyl})\)trifluoroborates in the presence of \(\text{Rh(OH)}(\text{cod})/\text{PPh}_3/\text{CsF}\) system in dioxane/water, and moderate to good yields of desired products were obtained (Scheme 21). The reaction scope appears that the diynes with internal alkene units are more reactive than those with terminal alkene units. A plausible catalytic cycle is depicted in Scheme 22.

### 2.2.1.2. Cobalt.

The utility of \(\text{CoI}_2(\text{PPh}_3)_2/\text{Zn}\) system in the \([2 + 2 + 2]\) cycloaddition of dipropargyl amines has been described by Cheng et al. They showed that \(N\)-propargyl amine 1

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**Scheme 16** Ni(0)-catalyzed enantioselective \([2 + 2 + 2]\) cycloadditions of dipropargyl amines 43 with acetylene.

**Scheme 17** (a) Formation of axially chiral biaryl diphosphonates 47 through \(\text{Rh(I)}\)-catalyzed \([2 + 2 + 2]\) cycloaddition; (b) synthesis of axially chiral biaryl dicarboxylate 49 from 9a and ester-substituted 1,3-butadiyne 48.

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**Scheme 18** Shibata’s synthesis of chiral tetraphenylenes 51.
in the treatment with allenes 59 in the presence of Co(I){PPh$_3$)$_2$ and zinc powder in 1,2-dichloroethane at 80°C, underwent a [2 + 2 + 2] cycloaddition to afford cyclohexadiene intermediates A, that after isomerization led to the 5-substituted isoindolines 60 in good yields (Scheme 23). It is noted that in the absence of zinc powder, no reaction occurred. To the best of our awareness this is the only example of isoindolines synthesis from dipropargyl amines and allenes reported so far.

2.2.2. Achiral isoindolines via intramolecular [2 + 2 + 2] cycloadditions. The reported methods for the synthesis of isoindoline cores via intramolecular cyclization of dipropargyl amines with olefins are very scarce and to the best of our knowledge there is only one example for such reactions. A Rh-catalyzed polycyclization of liner diyne-enones 61 has been developed for construction of fused isoindolines 62 in two steps under microwave irradiation. The polycyclization takes place through Rh-catalyzed intramolecular [2 + 2 + 2] cycloaddition of vinyl ketone with the diyne followed by aromatization to give observed products in moderate yields (Scheme 24a). Co/Mn-catalyzed intramolecular [2 + 2 + 2] cycloaddition of enediyynes 63, provided unsaturated isoindolines 64 instead (Scheme 24b).

2.2.3. Chiral isoindolines. Shibata and co-workers were the first to use a chiral rhodium complex-catalyzed intermolecular [2 + 2 + 2] cycloaddition of dipropargyl amines with olefins, to generate chiral isoindoline derivatives. In their investigations they reacted different unsymmetrical dipropargyl amines 65 with norbornene 66 in the presence of Rh–difluorophos complex,
and obtained tetracyclic products 67 in moderate to high yields with good to excellent enantioselectivity. As shown in Table 3 the reaction was more efficient for diynes possessing an ester functionality at their alkyne terminus.44

In another interesting report, Roglans and co-workers utilized rhodium complex A in the enantioselective intramolecular [2+2+2] cycloaddition of enediyynic macrocycles 68 to prepare fused tetracycles 69 (Scheme 25).45 Interestingly, using PNSO (N-phosphino tert-butylsulfamides)/Rh complex A combination as catalyst, the reaction afforded 69 in opposite enantioselectivity to that obtained with complex A.46

2.3. Isoindoline derivatives from nitrogen-linked 1,6-diyenes and arynes

The intermolecular [2+2+2] cocyclotrimerization of arynes with diynes represents an interesting method to prepare naphthalene ring systems. The use of this protocol in the synthesis of benzene fused isoindolines has been scarcely studied; in fact, only one example of such a reaction was reported in the literature by Hsieh and Cheng. They observed that treatment of dipropargyl amine 1 with 2-(trimethylsilyl)phenyl trifluoromethanesulfonate 70 (as benzylene precursor) in the presence of NiBr2(dppe)/CsF/Zn system gave 2,3-dihydro-1H-benzof[f]isoindole 71 in yield of 74% (Scheme 26).47

3. 2,3-Dihydro-1H-pyrrolo[3,4-c]pyridine derivatives from nitrogen-linked 1,6-diyenes and nitriles

3.1. Achiral 2,3-dihydro-1H-pyrrolo[3,4-c]pyridines via intermolecular [2+2+2] cycloadditions

3.1.1. Cobalt. After the pioneering work of Chiusoli and Costa, in 1989,48,49 who showed that dicobalt octacarbonyl complex could catalyze intermolecular [2+2+2] cycloaddition of dipropargyl amines with nitriles to give 2,3-dihydro-1H-pyrrolo[3,4-c]pyridines, this methodology has developed into a general powerful tool for the synthesis of a wide range of 2,3-
dihydro-1H-pyrrolo[3,4-c]pyridine derivatives. Thus, numerous cobalt-based catalytic system have been developed (Table 1). These include: CpCo(PPh3)2,50 dppe/CoCl2–6H2O/Zn,51 [CpCo(-CO)(fumarate)],52 [CpCo(cod)],53 and (1-neomenthylindenyl)Co(cod).54 An important study on this interesting field was carried out by Young and Deiters, in which solid-supported N-trityl dipropargyl amine72 were transformed into 2,3-dihydro-1H-pyrrolo[3,4-c]pyridine74 by the treatment with nitriles73 in the presence of CpCo(CO)2 under microwave irradiation (300 W) for 10 min, and then deprotection of trityl group with trifluoroacetic acid. The reaction tolerates both aromatic and aliphatic nitriles and gave final product in excellent yields (Scheme 27).55 DFT calculations suggest that this transformation takes place through cobaltacyclopentadiene intermediate.56

3.1.2. Rhodium. A preliminary example of Rh-catalyzed [2 + 2 + 2] cycloaddition of dipropargyl amines with nitriles has been reported by Tanaka et al., in 2006. They showed that treatment of diyne 9a with ethyl cyanoformate75 in the presence of [Rh(cod)2]BF4/BINAP system afforded the corresponding 2,3-dihydro-1H-pyrrolo[3,4-c]pyridine76 in a 99% isolated yield (Scheme 28a).57 Shortly afterwards, the same group has investigated the cycloaddition of 9a with perfluoroalkylacetylene77 under the aforementioned rhodium catalyst but using a higher catalyst loading (Scheme 28b).58 The authors also reported an example of pyrrolidine-fused pyridine80 preparation from 9a and diethyl phosphorocyanidate79 (Scheme 28c).59 Recently, Wan and co-workers described a general and efficient synthesis of 4,6,7-trisubstituted 2,3-dihydro-1H-pyrrolo[3,4-c]pyridines83 by the [2 + 2 + 2] cycloaddition of corresponding dipropargyl amines81 with ethyl 2-(hydroxyimino)acetate82 employing a Rh(NBD)2BF4/MeO-biphep combination as the catalytic system in refluxing ethanol. This process tolerated various sensitive functional groups and generally provided 83 in good to excellent yields (Scheme 29). However, dipropargyl amines derived from 4-bromobenzene-sulfonamide don’t work well under this reaction conditions. The results also demonstrate that the reaction do not give good yields with aromatic aldoximes, and aliphatic aldoximes failed to participate in the reaction.60 The mechanism proposed by the authors to explain

### Table 3 Rh-catalyzed asymmetric cycloaddition of nitrogen-linked 1,6-diyenes 65 with norbornene 66

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<td>69</td>
</tr>
<tr>
<td>3</td>
<td>Ph</td>
<td>COMe</td>
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<td>95</td>
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<tr>
<td>4</td>
<td>Ph</td>
<td>CH&lt;sub&gt;2&lt;/sub&gt;OMe</td>
<td>70</td>
<td>69</td>
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<tr>
<td>5</td>
<td>4-OMe-Ph</td>
<td>CO&lt;sub&gt;2&lt;/sub&gt;Me</td>
<td>90</td>
<td>95</td>
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<td>6</td>
<td>4-(CO&lt;sub&gt;2&lt;/sub&gt;Et)-Ph</td>
<td>CO&lt;sub&gt;2&lt;/sub&gt;Me</td>
<td>93</td>
<td>96</td>
</tr>
</tbody>
</table>

<sup>a</sup> 5,5'-Bis(diphenylphosphino)-2,2',2',2'-tetrafluoro-4,4'-bi-1,3-benzodioxole.

![Scheme 25](image_url)  
**Scheme 25** Rh(Ⅰ)-catalyzed [2 + 2 + 2] cycloisomerization of enediyne macrocycles 68.

![Scheme 26](image_url)  
**Scheme 26** Single example of a Ni-catalyzed [2 + 2 + 2] cycloaddition of dipropargyl amine 1 with 2-(trimethylsilyl)phenyl trifluoromethanesulfonate 70.
this reaction is based on the formation of hydroxylpyridine derivatives A as intermediates from the starting dipropargyl amines and oximes, followed by their spontaneous dehydration.

### 3.1.3. Ruthenium

Although, ruthenium catalyzed [2 + 2 + 2] cycloadditions of nitrogen-linked 1,6-diyne with nitriles are extremely efficient, they has received scarce attention. The most commonly used Ru-complex in general, and particularly in this chemistry, are commercially available Cp*Ru(cod)Cl, though only a few synthetic applications were reported (Scheme 30).

### 3.1.4. Iron

In 2011, B. Wan and co-workers demonstrated for the first time the usefulness of iron catalysts for the construction of 2,3-dihydro-1H-pyrrolo[3,4-c]pyridines. Thus, in the presence of FeI2/dppp/Zn as catalytic system in anhydrous tetrahydrofuran at room temperature, cycloaddition of dipropargyl amines 87 with nitriles 88 furnished corresponding

### Scheme 27

Microwave-mediated solid-supported formation of 2,3-dihydro-1H-pyrrolo[3,4-c]pyridine derivatives 74.

### Scheme 28

Rh-catalyzed [2 + 2 + 2] cycloaddition of nitrogen-linked 1,6-diyne 9a with nitriles developed by Tanaka.

### Scheme 29

Rhodium-catalyzed [2 + 2 + 2] cycloaddition of 1,6-diynes 81 and ethyl 2-(hydroxyimino)acetate 82 developed by Wan.

### Scheme 30

Ruthenium catalyzed [2 + 2 + 2] cycloadditions of dipropargyl amines 84 with nitriles 85.
pyrrolidine-fused pyridines 89 via azaferracyclopentadiene intermediates A (Scheme 31). The results showed that propargyl amines with internal alkyne unite gave higher yields than those with terminal unite, and aryl nitriles are slightly more reactive than alkyl nitriles. Shortly after, the same author showed that their catalytic system could effectively catalyze the cycloaddition of dipropargyl amines with cyanamides.

Along this line, Louie and co-workers reported an elegant regioselective iron-catalyzed [2 + 2 + 2] cycloaddition between unsymmetrical dipropargyl amines 90 and dimethylcyanamide 91 employing 5 mol% FeCl2 as catalyst, 10 mol% bis-(imino) pyridine as ligand, and 10 mol% Zn dust as reductant in benzene. Although both of possible regioisomers of the pyrrolidine-fused pyridines 92 and 93 are produced, their combined yields are good. Interestingly, in all cases the isomer that the larger alkyne substituent is placed next to the pyridine nitrogen has highest yield (Scheme 32a). In sharp contrast, the same reaction using Wan’s catalytic system shows a strong preference to place the smaller substituent next to the pyridine nitrogen (Scheme 32b). The author proposed mechanism to explain this different reaction patterns is depicted in Fig. 2.

3.1.5. Nickel. Nickel catalyzed [2 + 2 + 2] cycloaddition of dipropargyl amines with nitriles has been scarcely studied; in fact, only three examples of such a reaction were reported in the literature (Scheme 33). In 2005, the group of J. Louie presented an example of Ni-catalyzed [2 + 2 + 2] cycloaddition between

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**Scheme 31**  Cycloaddition of 1,6-diynes 87 with nitriles 88 catalyzed by FeI2/dppp/Zn system.

**Scheme 32**  (a) Iron-catalyzed cycloaddition reaction of diynes 90 and dimethylcyanamide 91 developed by Louie; (b) iron-catalyzed cycloaddition reaction of diyne 90a and dimethylcyanamide 91 developed by Wan.

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**Fig. 2**  Mechanistic proposal for the reactions in Scheme 32.
internal dipropargyl amine 9a and benzonitrile employing 3 mol% of Ni(cod)$_2$ as catalyst and 6 mol% of SIPr as ligand in toluene at room temperature. The corresponding pyrrolidine-fused pyridines 94 was obtained in yield of 78%. Subsequently, the same group reinvestigated this reaction using Ni(acac)$_2$/SIPr·HCl/ nBuLi/benzene system but lower yield than previously reported was obtained. Ni-catalyzed cycloaddition of 9a and pyrrolidine-1-carbonitrile, was also reported by the same authors.

3.1.6. Iridium. The [2 + 2 + 2] cycloaddition of dipropargyl amines with nitriles can also be carried out efficiently by iridium(i) complex of [Ir(cod)Cl]$_2$ (Scheme 34). However, the number of reported examples are very scarce and there is an urgent need to study the scope and limitations of this chemistry.

3.2. Achiral 2,3-dihydro-1H-pyrrolo[3,4-c]pyridines via intramolecular [2 + 2 + 2] cycloadditions

To develop a new protocol for the construction of 2,3-dihydro-1H-pyrrolo[3,4-c]pyridines via [2 + 2 + 2] cycloaddition reaction, Cheng and co-workers have investigated the purely intramolecular cycloaddition of dipropargyl amine-nitriles 101 in the presence of Co$_4$(dppe)/Zn system in refluxing acetonitrile, and
good yields of polycyclic products 102 was obtained (Scheme 35a). The intramolecular cycloaddition of dipropargyl amine-nitriles was also performed under microwave-assisted conditions. For example, 2-(pyridin-2-yl)pyridine 103 was prepared in yield of 88% via Rh(I)-catalyzed double cocyclization of 102 under microwave irradiation at 90 °C (Scheme 35b).

### 3.3. Chiral 2,3-dihydro-1H-pyrrolo[3,4-c]pyridines

There has been a recent increase in the use of chiral substrates or ligands in the [2 + 2 + 2] cycloaddition of dipropargyl amines with nitriles for the construction of chiral cycloadducts. In an elegant study, Shibata’s group disclosed the catalytic enantioselective synthesis of planar-chiral macrocyclic tripodal cage compounds 105 via rhodium(i)(R)-Cy-BINAP complex-catalyzed intramolecular cycloaddition of 2-aminophenol-tethered dipropargyl amine-nitriles 104 (Scheme 36). Under optimized conditions, cycloadducts 105 were obtained in high yields and excellent enantiomeric excess up to 99% ee. However, very poor enantioselectivities were achieved with diyne-nitriles possessing very long ansa chains \( n = 11 \). To solve the problem, the authors replaced Cy-BINAP with BINAP. Along this line, Tanaka and co-workers reported an example of regio- and enantioselective synthesis of axially chiral 7-(2-chlorophenyl)-pyrrolidinopyridine 107 by cationic rhodium(I)/S-BINAP complex catalyzed atroposelective [2+2+2] cycloaddition of unsymmetrical dipropargyl amine 106 with ethyl cyanoformate 85a (Scheme 37a) and Tan’s group performed Rh-catalyzed [2 + 2 + 2] cycloaddition of chiral tert-butylsulfanimidated N-propargylamines 108 with 85a to prepare chiral pyrrolidine-fused pyridine 109 (Scheme 37b).

### 4. 2,3-Dihydro-1H-pyrrolo[3,4-c]pyridin-6(5H)-one derivatives from nitrogen-linked 1,6-diynes and isocyanates

#### 4.1. Achiral 2,3-dihydro-1H-pyrrolo[3,4-c]pyridin-6(5H)-ones

The first mention of the synthesis of 2,3-dihydro-1H-pyrrolo[3,4-c]pyridin-6(5H)-ones through a ruthenium(II)-catalyzed [2 + 2 + 2] cycloaddition between dipropargyl amines and isocyanates can be found in a 2001 paper by Itoh, although only one example was described (Scheme 38a). Louie and co-workers have subsequently shown that Ni(cod)2/SIPr system can also efficiently catalyze this reaction (Scheme 38b). Later, the group of Tanaka extended the scope of isocyanates to benzyl and alkyl isocyanates employing a cationic rhodium(i) catalytic system (Scheme 38c) and Takeuchi’s group improved the efficiency of this reaction in terms of yield and reaction time by performing the process in refluxing 1,2-dichloropropane using [Ir(cod)Cl]2 as the catalyst (Scheme 38d).

Recently, Pérez-Castells et al. described the preparation of 2,3-dihydro-1H-pyrrolo[3,4-c]pyridin-6(5H)-ones 117 in moderate yields via [2 + 2 + 2] cycloaddition of isocyanates 116 with both symmetrical and unsymmetrical dipropargyl amines 115 in the...
presence of catalytic amounts of ruthenium–alkylidene complex 118 (Scheme 39). The reaction of unsymmetrical dipropargyl amines exhibits remarkable regioselectivity, and the substituent is located exclusively next to the 2-pyridone nitrogen. The proposed mechanism is outlined in Fig. 3.

4.2. Chiral 2,3-dihydro-1H-pyrrolo[3,4-c]pyridin-6(5H)-ones

In 2008, Tanaka and co-workers reported an example of axially chiral 2,3-dihydro-1H-pyrrolo[3,4-c]pyridin-6(5H)-one preparation through a cationic rhodium(I)-catalyzed [2+2+2] cycloaddition strategy. They showed that dipropargyl amine 9a underwent cyclization with 2-methoxybenzene-isocyanate 116 in the presence of [Rh(cod)₂]BF₄/R'-BINAP as catalytic system in DCM at room temperature and gave the corresponding pyrrolidine-fused pyridinone 120 in good yield (70%), although in low enantiomeric excess (34%) (Scheme 40, conditions A). Five years later, the group of Aubert reinvestigated this reaction employing [Rh(cod)Cl]₂/(R)-BINAP/AgBF₄ combination as the catalytic system. Under these conditions, the cycloadduct 120 was obtained in good enantiomeric excess, although in low yield (Scheme 40, conditions B). This drawback was significantly improved by replacing of (R)-BINAP with (R)-DM-BINAP (Scheme 40, conditions C).

5. 2,3-Dihydropyrano[4,3-c]pyrrol-6(1H)-ones from nitrogen-linked 1,6-diienes and carbon dioxide

An interesting and rare example for the synthesis of 2,3-dihydropyrano[4,3-c]pyrrol-6(1H)-ones was reported by Saegusa et al. in 1988. Thus, in refluxing benzene, Ni(0)-catalyzed [2 + 2 + 2] cycloaddition between dipropargyl amine 121 and carbon dioxide, furnished -dihydropyrano[4,3-c]pyrrol-6(1H)-one 122 in yield of 78% (Scheme 41a). The mechanism proposed by the authors to explain this reaction is depicted in Scheme 42. 26 years later, Tanaka and co-workers reinvestigated this reaction employing [Rh(cod)₂]BF₄/[R]-BINAP/DCE system, but the corresponding 2-pyranone 123 was isolated in only 14% yield (Scheme 41b).

6. 2,3-Dihydro-pyrrolo[3,4-c]pyrindin-6-imines from nitrogen-linked 1,6-diienes and carboxiimides

Young and Deiters have also developed a novel method for the preparation of 2,3-dihydro-pyrrolo[3,4-c]pyrindin-6-imines 125 involving Co(n)-catalyzed [2 + 2 + 2] cycloaddition of resin-bound N-trityl dipropargyl amine 72 with carboxiimides 124 under microwave irradiation conditions, followed by deprotection of trityl group with trifluoroacetic acid (Scheme 43a). The authors found that microwave irradiation play a crucial role for the success of the reaction. When the same...
The cyclotrimerization was conducted without microwave irradiation, no product formation was observed, even after a prolonged reaction time of 24 h at 110 °C. In a closely related investigation, Tanaka and co-workers reported a cationic rhodium(I) catalytic system to get access to 2,3-dihydro-pyrrolo[3,4-c]pyridin-6-imines \(^{127}\) (Scheme 43b). \(^{92}\)

Ketenes are one of the latest coupling partners to join the [2 + 2 + 2] cycloaddition story of dipropargyl amines. In 2011, Louie et al. successfully prepared 2,3-dihydro-1H-isoindol-5-ones \(^{129}\) utilizing cycloaddition of dipropargyl amine \(^{9a}\) and 1.2 equivalents of ketenes \(^{128}\) in the presence of Ni(cod)\(_2\)/DPPB system in moderate yields (Scheme 44). \(^{93}\) Interestingly, under this reaction conditions no pyran product was observed, it means...
that exclusively the C=C bonds in the ketenes participated in the reactions.

8. Outlook and conclusion

Transition metal catalyzed [2 + 2 + 2] cycloaddition of nitrogen-linked 1,6-dynes with unsaturated motifs, have drawn much interest from organic chemists because of their high efficiency in the construction of various pyrrolidine-fused systems. This method offers the opportunity to synthesis of complex frameworks, which may be difficult or impossible by other means. It also offers the possibility for the rapid synthesis of potential drugs from cheap and easily available dipropargyl amines in one step with ideal atom economy.

Despite the significant achievements during the past few years in this field, many challenges still remain to be overcome: (a) most of the [2 + 2 + 2] cycloadditions using nitrogen-linked 1,6-dyne reagents are limited to the use of expensive transition metal catalysts. Thus the exploration of cheaper transition metal catalysts (such as Fe and Cu catalysts) are highly desirable in terms of cost and availability; (b) generally, the number of reported examples are narrow and there is an urgent need to study the scope and limitations of reagents; (c) and finally, the mechanism of the reactions remains elusive, though some mechanistic courses were proposed based on DFT calculations.

Conflicts of interest

There are no conflicts to declare.

References


