

EDGE ARTICLE

View Article Online
View Journal | View IssueCite this: *Chem. Sci.*, 2021, 12, 435

All publication charges for this article have been paid for by the Royal Society of Chemistry

Received 11th September 2020
Accepted 30th October 2020

DOI: 10.1039/d0sc05036a

rsc.li/chemical-science

Metal–ligand–Lewis acid multi-cooperative catalysis: a step forward in the Conia-ene reaction†

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An original multi-cooperative catalytic approach was developed by combining metal–ligand cooperation and Lewis acid activation. The [(SCS)Pd]₂ complex featuring a non-innocent indenediide-based ligand was found to be a very efficient and versatile catalyst for the Conia-ene reaction, when associated with Mg(OTf)₂. The reaction operates at low catalytic loadings under mild conditions with HFIP as a co-solvent. It works with a variety of substrates, including those bearing internal alkynes. It displays complete 5-*exo* vs. 6-*endo* regio-selectivity. In addition, except for the highly congested ^tBu-substituent, the reaction occurs with high *Z* vs. *E* stereo-selectivity, making it synthetically useful and complementary to known catalysts.

Introduction

Major breakthroughs have been made in homogeneous catalysis thanks to cooperative approaches.¹ Concomitant and synergistic participation of several active sites, as in enzymes, is very attractive to achieve chemical transformations under milder conditions and reduce wastes. Thanks to cooperativity, not only the efficiency and selectivity of catalysts can be improved, but the diversity and complexity of the obtainable chemical compounds can also be expanded. In this context, metal–ligand cooperativity (MLC) is clearly prominent.² Here, a ligand participates directly in substrate activation, in addition to modulating the stereoelectronic properties of the metal center. Complexes of the group 7–9 metals featuring non-innocent pincer ligands, in particular amido and pyridine-based scaffolds, occupy a forefront position in this area.³ Tremendous achievements in hydrogenation and dehydrogenation reactions have been reported with these catalysts.⁴ Extending the scope of transformations⁵ and developing the MLC catalytic approach further are currently a major endeavor, along with the diversification of complexes displaying MLC.

Our group introduced a few years ago an SCS pincer framework featuring an electron-rich indenediide back-bone.⁶ The paucity of MLC behavior reported for Pd and Pt complexes,⁷ despite the prominent role of these metals in homogeneous

catalysis, led us to study complexes of type **I** (Fig. 1). The electron-rich indenediide motif was shown to be non-innocent and to open a MLC pathway for Pd and Pt-catalyzed C–O and C–N bond formation. The (SCS) Pd,Pt complexes proved to very efficiently promote the cycloisomerization of alkynoic acids and *N*-tosyl alkynylamides, giving access in high yield and selectivity to a wide variety of alkylidene lactones and lactams, including seven-membered rings.⁸ Of note, indepth mechanistic studies substantiated the participation of proton shuttling in addition to MLC.⁹ This drove us to use catechols as hydrogen bond donor (HBD) additives. The activity and selectivity of the (SCS) Pd,Pt complexes were thereby spectacularly improved.⁹ The synergy between MLC and proton shuttling is a very appealing approach to boost catalytic performance. It has been proposed for many hydrogenation/dehydrogenation reactions involving MLC¹⁰ and recently unequivocally demonstrated by Hazari *et al.*¹¹ Its

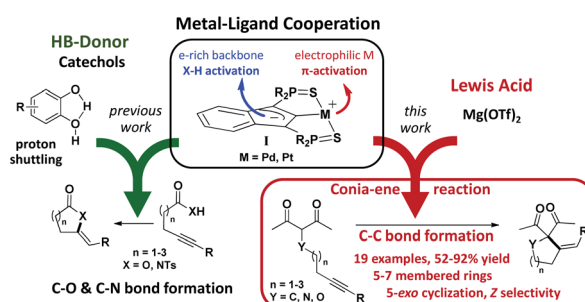


Fig. 1 Multi-cooperative catalysis combining metal–ligand cooperation (MLC, indenediide-based (SCS)M pincer complexes) with hydrogen-bond donors (HBD, previous work) or Lewis acid activation (LA, this work) to promote cycloisomerization reactions (C–O, C–N and C–C bond formation).

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† Electronic supplementary information (ESI) available: Full experimental procedures, characterization data of all new compounds (including ¹H, and ¹³C-NMR spectra, and ADEQUATE, NOESY and dsel-HSQC-IPAP experiments). See DOI: 10.1039/d0sc05036a

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impact in intermolecular hydroamination reactions has also been substantiated.^{7g}

Encouraged by these results, we wondered about the possibility of developing new multi-cooperative catalytic approaches based on MLC. Another aim was to expand the scope of SCS Pd,Pt catalysts to more challenging transformations. In this work, we focused on Conia-ene cyclizations, *i.e.* the formation of carbo- and heterocycles by C–C coupling between an enolizable group, typically a β -dicarbonyl moiety, and an alkyne. As pointed out by Enders *et al.* in their recent review,¹² this is a very powerful transformation and much progress has been achieved recently from both methodological and synthetic viewpoints, but several limitations still need to be addressed. While investigating the ability of the MLC Pd indenediide complex to catalyze Conia-ene cyclizations, we discovered that it is extremely efficient and versatile when associated with a Lewis acid (LA). The combined use of TM catalysts and Lewis acids (often referred to as dual activation) has attracted increasing interest and led to impressive developments over the last decade,¹³ including in the Conia-ene reaction.^{12,14} Yet, synergistic effects between MLC and LA activation have not been realized so far. As reported hereafter, the association of $[(SCS)Pd]_2$ and $Mg(OTf)_2$ provides a first example of such a dual and multi-cooperative catalytic system. It is a very efficient and useful catalyst for Conia-ene cyclizations, displaying broad substrate scope and high regio as well as stereo-selectivity.

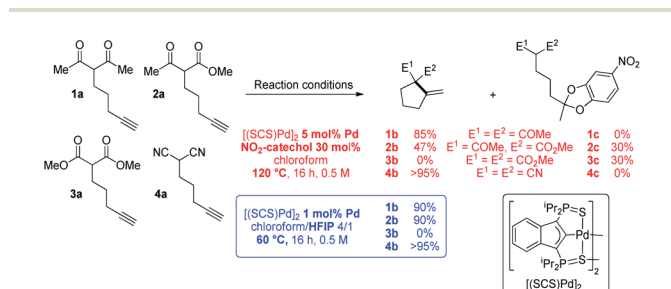
Results and discussion

To start with, the carbo-cyclization of the β -dicarbonyl compounds **1a–3a** and β -dinitrile **4a** bearing an alkyne moiety was studied applying the reaction conditions previously optimized for the cycloisomerization of alkynoic acids (Scheme 1).⁹ A series of hydrogen-bond donors (catechols..., 30 mol%, Table S1†)¹⁵ were probed as proton shuttling additives to the indenediide-derived Pd complex $[(SCS)Pd]_2$ (5 mol% Pd). The best results were obtained with 4-nitrocatechol. Heating at 120 °C for 16 h, full conversion could be achieved in all cases but **3a** (only 30% in this case). The expected cyclized products **1b** and **4b** were selectively obtained from the β -diketone and β -dinitrile substrates **1a** and **4a**, whereas **2a** led to the formation

of a secondary product **2c** (30% yield) resulting from double addition of the catechol to the C \equiv C triple bond,¹⁶ besides the targeted carbocycle **2b** (47% yield). As for the β -diester **3a**, no cyclized product **3b** was formed, only the secondary product **3c** was obtained.

The known ability of hexafluoroisopropanol (HFIP) to form hydrogen-bond networks and participate in proton shuttling¹⁷ prompted us to screen a series of fluorinated alcohols as additives. This resulted in noticeable improvement of activity and selectivity (Scheme 1 and Table S2,† entries 1–4).¹⁵ Using a 4/1 chloroform/HFIP solvent mixture, high conversions of the β -diketone **1a** and β -dinitrile **4a** were achieved in this case at 60 °C with only 1 mol% of Pd (entry 6). The fact that **4a** was readily cyclized is particularly noteworthy since β -dinitriles are usually significantly less reactive than β -dicarbonyl substrates in the Conia-ene reaction.¹² It is because the linear geometry of the cyano group is not suited for intramolecular hydrogen-bonding and thus tautomerization is more demanding. Under these milder conditions, the β -keto,ester **2a** could also be efficiently cyclized and **2b** was formed in high yield without a side product. In turn, the β -diester **3a** was unreactive. Due to their weaker acidity, β -diester substrates are notoriously less reactive and more challenging in the Conia-ene reaction.¹² The contribution of proton shuttling to the carbocyclization of **1a**, **2a** and **4a** in the presence of HFIP is supported by a strong solvent dependence. While toluene and dichloromethane gave results close to those obtained in chloroform, hydrogen-bond accepting solvents such as THF or DMSO completely inhibited the reaction.

Thus, the indenediide pincer Pd complex $[(SCS)Pd]_2$ is a competent catalyst for the Conia-ene reaction of β -diketo, β -keto,ester and β -dinitrile substrates, when combined with HFIP as a co-solvent. However, it fails to cyclize the corresponding β -diester under the same conditions. Although the $[(SCS)Pt]_2$ complex proved to be superior for the cyclization of challenging alkynoic acids and amides,^{8c} no improvement was observed in the Conia-ene reaction upon changing Pd for Pt. The Pt complex proved to be in fact less active, with only 40% (*vs.* 90%) conversion of **2a** after 16 h (Table S2†).¹⁵ Looking for another way to boost the catalytic activity, we envisioned to use di or trivalent Lewis acids, which are known to activate β -dicarbonyl compounds and facilitate the formation of the corresponding enols. Combinations of hard and soft Lewis acids have been occasionally used in Conia-ene reactions. Such dual activation proved in particular successful to promote enantioselective cyclization of β -diketo and β -keto,ester derivatives.^{14a–cf} Given these precedents, we wondered about the possibility of implementing a new type of multi-cooperative catalytic approach by merging metal–ligand cooperation, as achieved by the (SCS) Pd complex, with the activation of the β -dicarbonyl moiety by a LA.¹⁸ Multi-cooperative catalytic systems are very attractive but challenging to achieve. They inherently raise compatibility issues and the different components must be carefully adjusted to act synergistically. In the case of the (SCS) Pd complex, the Lewis acid additive may interact with the electron-rich indenediide motif as well as the sulfur atoms and thereby interfere with the Pd–ligand cooperativity.¹⁹ A series of triflate salts were evaluated as LA additives to $[(SCS)Pd]_2$ in the cyclization of the β -keto,ester **2a** in 4/1 chloroform/HFIP as solvent. Initial



Scheme 1 Conia-ene cyclization of the β -dicarbonyl and β -dinitrile substrates **1a–4a** catalyzed by the $[(SCS)Pd]_2$ complex in the presence of 4-nitrocatechol or HFIP. Throughout the paper, the a/b letters in the compound numbers refer to the substrate/cyclized Conia-ene product, respectively.



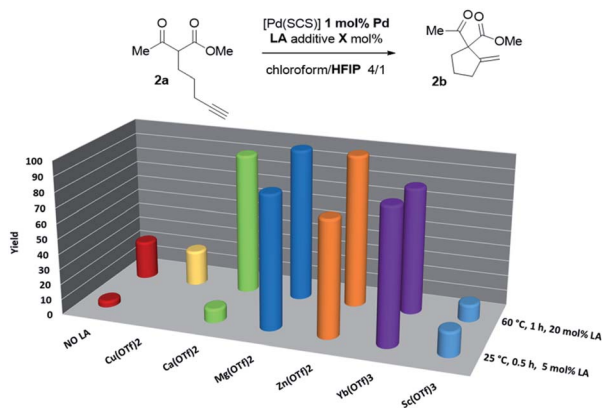


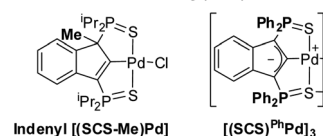
Fig. 2 Evaluation of the catalytic activity of the Pd complex $[(SCS)Pd]_2$ in the presence of LA additives.

screenings were performed at 60 °C with 1 mol% Pd and 20 mol% LA (Fig. 2 and Table S3†).¹⁵ While $Cu(OTf)_2$ and $Sc(OTf)_3$ had no or negative impact on catalysis, significant improvement of activity was observed with $Ca(II)$, $Mg(II)$, $Zn(II)$ and $Yb(III)$ salts.²⁰ High conversions were achieved in only one hour (>82% vs. only 25% without a LA), demonstrating the compatibility and synergy between MLC with the $[(SCS)Pd]_2$ complex and electrophilic activation of the β -dicarbonyl moiety with a LA. Taking advantage of the catalytic boost triggered by the LA additive, milder reaction conditions were then used. The temperature was decreased to 25 °C and the LA loading was reduced to 5 mol%. Under these mild conditions, $Mg(OTf)_2$ and $Yb(OTf)_3$ led to the best results (~90% conversion to **2b** within only 30 min, and >95% after 1.5 h). The β -diketo substrate **1a** was also readily cyclized under these conditions (>95% after 8 h). In contrast, similar results were obtained with and without Lewis acids in the case of the β -dinitrile **4a** (85% conv. is achieved in 16 h at 25 °C). Chelating coordination of the two cyano groups is not possible geometrically, and thus the LA cannot facilitate deprotonation, as in the case of the β -dicarbonyl substrates. Most remarkably, the combination of $[(SCS)Pd]_2$ with the LA allowed for the cyclization of the challenging β -diester **3a** as well. Here, $Mg(OTf)_2$ significantly outperformed $Yb(OTf)_3$ (83 vs. 13% conversion of **3a** within 30 min). Given the superior efficiency, abundance and biological tolerance of Mg over Yb, catalytic studies were continued with $Mg(OTf)_2$ as the LA.²¹

To ascertain the need and synergy of all the catalytic components, a series of control reactions were carried out for the cyclization of the β -keto,ester substrate **2a**. The results obtained under conditions deviating from the optimal ones [1 mol% Pd, 5 mol% $Mg(OTf)_2$, chloroform/HFIP 4 : 1, 25 °C] are disclosed in Table 1. As previously pointed out, the presence of HFIP and $Mg(OTf)_2$ is crucial to boost the catalytic activity and makes the Conia-ene reaction work at 25 °C with low Pd loadings. Conversely, in the absence of the Pd complex, no cyclized product (or only a very small amount) was formed with HFIP alone or with a mixture of $Mg(OTf)_2$ and HFIP. The electron-rich indenediide moiety of the SCS ligand also plays a major role, as

Table 1 Impact of the different components of the catalytic system (Pd complex, Lewis acid, and proton shuttle additive) on the carbocyclization of the keto,ester **2a**

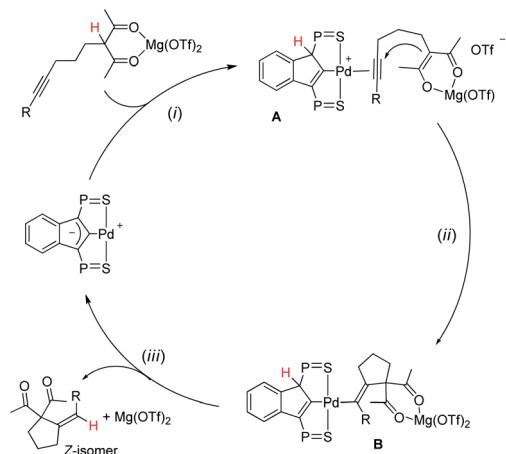
Reaction conditions, deviations from the optimal	Conv./reaction time
Optimal: $[(SCS)Pd]_2$ 1 mol% Pd, 5 mol% $Mg(OTf)_2$ chloroform/HFIP 4/1, 25 °C	>95%/1.5 h
$[(SCS)Pd]_2$ 5 mol% Pd, 20 mol% $Mg(OTf)_2$ chloroform, 25 °C, no HFIP	46%/60 h
No Mg salt, 60 °C	>95%/22 h
No Pd complex, no Mg salt, 60 °C	Traces/16 h
No Pd complex, 20 mol% $Mg(OTf)_2$	0%/2 h
Indenyl $[(SCS-Me)Pd]$ complex	>95%/8.5 h
Indenyl $[(SCS-Me)Pd]$ complex, 5 mol% Pd, no Mg salt, 60 °C	90%/15 days
$[PdCl(allyl)]_2$ 1 mol% Pd	6%/24 h
$[PdCl_2(PhCN)]_2$ 1 mol% Pd	7%/24 h
$[PdCl_2(PhCN)]_2$ 1 mol% Pd, dppe 1 mol%	35%/24 h
$[PdCl_2(PhCN)]_2$ 1 mol% Pd, dppe 1 mol%, iPr_2EtN 1 mol%	78%/24 h
$[(SCS)^{Ph}Pd]_3$ 5 mol% Pd, 20 mol% $Mg(OTf)_2$	>95%/20 min



substantiated by the noticeable decrease in activity observed when shifting to the corresponding 1-Me-2-indenyl Pd complex.

In addition, much lower conversions were observed with $[PdCl(allyl)]_2$ and $[PdCl_2(PhCN)]_2$ (6–7% after 24 h), as well as $[PdCl_2(dppe)]$ (35%). Even the combination of $[PdCl_2(dppe)]$ with a base (iPr_2EtN , 1 mol%) performed poorly (78% conversion after 24 h) compared to the $[(SCS)Pd]$ system (>95% in 1.5 h).

Seeking to evidence the non-innocent character of the ligand *via* the formation of indenyl $[(SCS-H)PdX]$ species, several catalytic reactions were *in situ* analyzed by ^{31}P NMR spectroscopy (see the ESI for a detailed description, Fig. S1–S4†). These monitorings were complicated by the fact that HFIP has an impact on the structure/association degree of the catalyst, and only broad signals could be observed (the dimer $[(SCS)Pd]_2$ was recovered intact after evaporation of the solvent at the end of the catalytic reaction). When the reaction was carried out in the absence of HFIP, only the dimer $[(SCS)Pd]_2$ was observed along the reaction, indicating that it is the resting state. This behaviour is reminiscent of that observed in the cycloisomerization of alkynoic acids,⁹ and is consistent with the tight association of the two (SCS)Pd fragments. We decided then to evaluate the behaviour of a pre-catalyst bearing Ph instead of iPr groups on the P atoms. The less electron-donating $Ph_2P=S$ sidearms led to weaker association of the $(SCS)^{Ph}Pd$ fragments.^{8a} Only broad signals were again observed in the presence of HFIP, but when the reaction was carried out in the absence of HFIP, minor signals corresponding to indenyl species were clearly detected by ^{31}P NMR spectroscopy. Although $[(SCS)^{Ph}Pd]_3$ remains the major species, the observation of indenyl species during the



Scheme 2 Catalytic cycle proposed to account for the Conia-ene reaction catalyzed by the (SCS)Pd complex and $\text{Mg}(\text{OTf})_2$ (the ^iPr groups at P are omitted for clarity).

catalytic transformation is consistent with the indenediide ligand backbone acting as a base.

A simplified 3-step mechanism for the Conia-ene reaction catalyzed by the association of $[(\text{SCS})\text{Pd}]_2$ and $\text{Mg}(\text{OTf})_2$ is depicted in Scheme 2: (i) *double activation of the substrate*: the LA chelates the β -dicarbonyl moiety, the acidic proton is transferred to the electron-rich indenediide backbone of the SCS ligand while the alkyne coordinates to palladium (intermediate **A**); (ii) *cyclization*: the ensuing enolate adds to the π -activated $\text{C}\equiv\text{C}$ triple bond to give the alkenyl Pd complex **B**; (iii) *product release*: back-transfer of the proton from the ligand backbone to the $\text{C}(\text{sp}^2)$ atom bonded to Pd results in proto-depalladation and regenerates the indenediide Pd catalyst. HFIP is presumed to help dissociate the dimeric pre-catalyst,^{17c} and to promote proton shuttling from the substrate to the indenediide

moiety in step (i) and from the ligand backbone to the $\text{C}(\text{sp}^2)$ carbon at Pd in step (iii).

The scope and functional group tolerance of the reaction were then explored (Fig. 3). Our aim was not only to probe the generality of the new MLC-LA dual catalytic approach, but also to improve the efficiency of the Conia-ene transformation and expand its synthetic interest. Substrates with a terminal alkyne and prone to 5-*exo* cyclization were investigated first. The (SCS) Pd-Mg association proved to be particularly efficient to cyclize β -diesters **5a–7a** featuring an adjacent heteroatom (N or O). The amide-tethered substrate **5a** was quantitatively cyclized in only 5 min at 25 °C. Shifting the amide moiety at the *exo* position (**6a**) slowed down the transformation somewhat, but it still proceeded rapidly and efficiently at 25 °C (>95% in 50 min). Cyclization of the ether-tethered substrate **7a** was more challenging, but heating to 75 °C enabled to achieve high conversion (90%) in only 4.25 h.

The length of the linker between the β -dicarbonyl moiety and the terminal alkyne was then modified with the aim to access larger rings (**8a–12a**). 6-*Exo* cyclization of β -keto, esters such as **8a** featuring a flexible butylene tether is tricky. It has only been scarcely achieved, using a Au(i) complex featuring a tris-alkynyl phosphine with bulky end caps (1 mol%, 25 °C, 1.5 h) or $\text{In}(\text{NTf}_2)_3$ (1 mol%, 80 °C, 4 h).^{22,23} Pleasingly, the methylene-cyclohexane product **8b** could also be obtained efficiently using the $[(\text{SCS})\text{Pd}]_2$ complex (5 mol% Pd) and $\text{Mg}(\text{OTf})_2$ (20 mol%) within 20 h at 75 °C (90%). The dual catalytic system proved to be particularly efficient when the enol position bears an oxygen or nitrogen atom. Such substrates have been shown by Hatakeyama *et al.* to provide straightforward entry to valuable heterocycles and very good results in terms of activity and selectivity were obtained using $\text{In}(\text{OTf})_3$ with DBU (5 mol% each) in refluxing toluene.²⁴ In our hands, the β -diester substrates **9a–11a** reacted smoothly and high yields (>85%) were achieved with low catalytic loadings (1 mol% Pd, 5 mol% Mg).

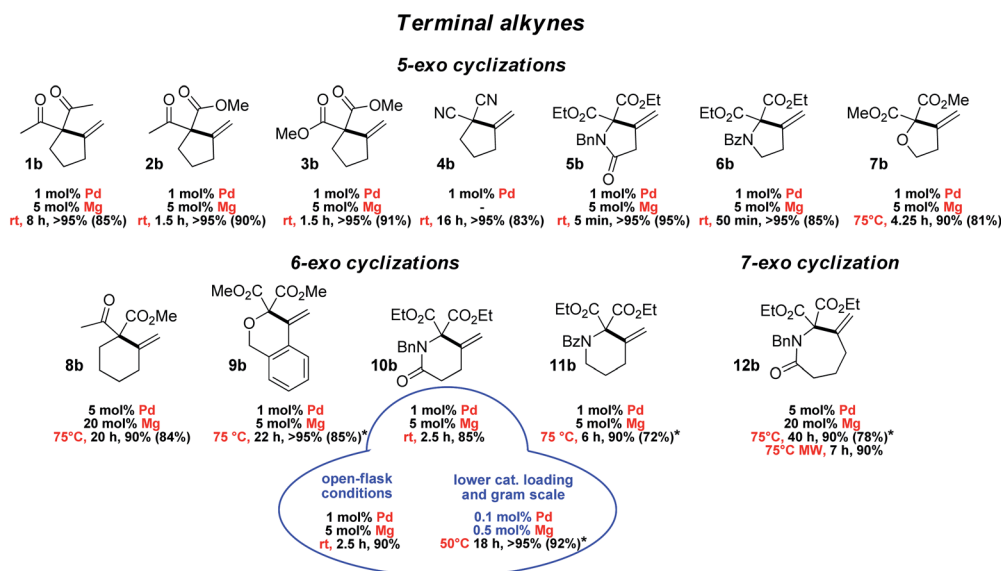


Fig. 3 Substrate scope for the Conia-ene reaction catalyzed by the (SCS)Pd complex and $\text{Mg}(\text{OTf})_2$: β -dicarbonyl substrates bearing a terminal alkyne (isolated yields are indicated in parentheses; * stands for 0.1 M substrate concentration instead of 0.5 M).



Particularly noteworthy is the formation of the δ -valerolactame **10b** which was accomplished in only 2.5 h at 25 °C. Even 7-*exo* cyclization was possible and highly yielding. Using the same conditions as for **8a**, the ϵ -caprolactame **12b** was obtained in 90% yield within 40 h, and the reaction time could be reduced to 7 h using microwave irradiation.²⁵ The preparation of such seven-membered rings was previously described using In(OTf)₃/DBU as the catalyst, but in modest yield (41%) and with high catalytic loading (15 mol%).²⁴

To assess the robustness of the Pd–Mg dual catalytic system, the cyclization of the amide-tethered substrate **10a** was carried out under the optimized conditions (25 °C, 1 mol% Pd, 5 mol% Mg, chloroform/HFIP 4/1) but under open-flask conditions and with technical grade solvents. The δ -valerolactame **10b** was obtained in similar yield to under inert conditions, demonstrating that the reaction does not require stringent precautions. Furthermore, when the catalytic loading was reduced by a factor of ten (0.1 mol% Pd and 0.5 mol% Mg), the product was still obtained in very high yield (>95% in 18 h at 50 °C). These reaction conditions were successfully applied to larger scale: 920 mg of pure product **10b** were prepared using only 1.4 mg of [(SCS)Pd]₂ and 4.5 mg of Mg(OTf)₂. Merging metal–ligand cooperation and Lewis acid activation, as realized with the [(SCS)Pd]₂ complex and Mg(OTf)₂, thus appears general, practical and of synthetic value. It promotes 5-, 6- and 7-*exo* cyclization of β -dicarbonyl substrates featuring terminal alkynes under mild conditions, and gives access to a variety of carbo- and heterocycles.

The good performance of the dual Pd–Mg catalytic system prompted us to explore then the cyclization of substrates bearing internal alkynes, which are significantly more challenging than terminal alkynes (Fig. 4). Being way less reactive, these substrates require harsher conditions, which often results in regio-selectivity issues with the formation of mixtures of *exo* and *endo* cyclization products.²⁶ Given the preference of 5-*endo* against 4-*exo-dig* cyclization, the β -keto,ester **13a** and β -diester **14a** were tested first. A longer reaction time was needed for **13a** than for **14a** (25 vs. 1.5 h) but in both cases, the reaction proceeded readily at 25 °C and afforded high yield (95%) of the 5-*endo* product. No trace of the less favored 4-*exo* product was detected by ¹H NMR spectroscopy. Our dual catalytic system does not compete with (Ph₃P)AuOTf for the formation of cyclopentene **13b** (83% yield was achieved in only 5 min at room temperature using 1 mol% Au),²⁷ but on the other hand, the gold catalyst works only with β -keto,esters and failed to cyclize β -diesters.²³ As far as the cyclization of **14a** is concerned, it had only been efficiently accomplished with Zn halides (10 mol%) but required high temperature and long reaction times (100 °C, 15 h).²⁸ We then investigated 5-*exo* Conia-ene cyclization with internal alkynes (β -diester substrates **15a–19a**). The terminal substituent was varied (R = Me, Et, Cy, Ph, ^tBu) to assess the impact of sterics and electronics. It should be noted that precedents of Conia-ene cyclization with internal alkynes mostly involve aryl and primary alkyl substituents, while examples with secondary alkyl groups are much rarer and tertiary ones are unprecedented.^{12,24,27,29} Due to the lower reactivity of these substrates, the catalytic loadings were fixed to 5 mol% Pd and 20 mol% Mg. For the less hindered substrates,

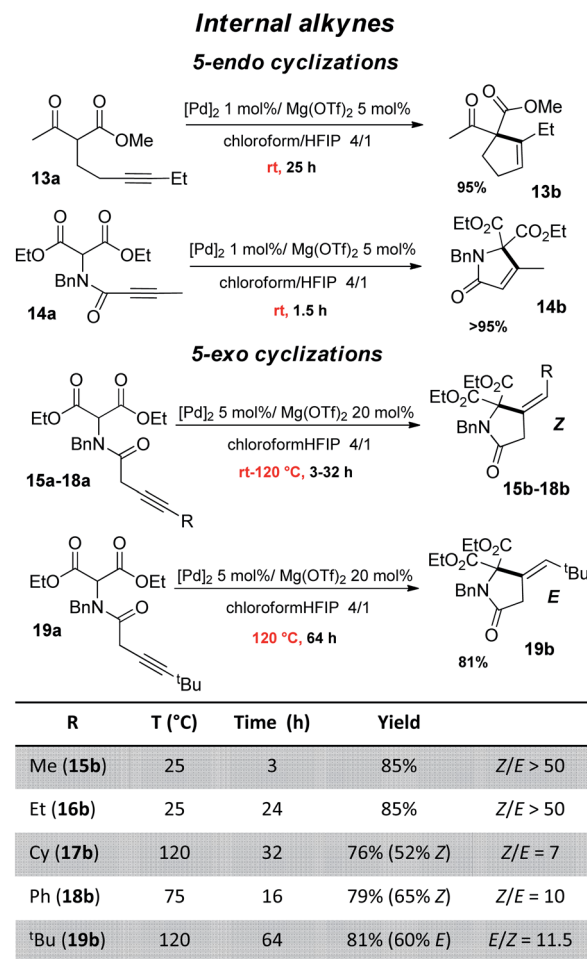


Fig. 4 Conia-ene reaction catalyzed by the (SCS)Pd complex and Mg(OTf)₂: β -dicarbonyl substrates bearing an internal alkyne (isolated yields in the Z isomer are indicated in parentheses).

with Me (**15a**) and Et (**16a**) substituents, cyclization occurred at 25 °C within 3 and 24 h, respectively, while those bearing Cy (**17a**), Ph (**18a**) and ^tBu (**19a**) groups required heating at 75–120 °C to achieve high conversions. For all substrates, including the ^tBu-substituted **19a**, the reaction was highly selective. The structures of the obtained products were unambiguously determined thanks to thorough ¹H and ¹³C NMR spectroscopic analyses including ADEQUATE, NOESY and dse1-HSQC-IPAP experiments.¹⁵ Complete selectivity in favor of 5-*exo* vs. 6-*endo* cyclization was observed, whatever the substituent at the alkyne (alkyl or phenyl). For **15a** and **16a**, a single 5-*exo* product was obtained, authenticated as the Z isomer. For the substrates **17a** and **18a** bearing bulkier groups, which required higher reaction temperature, two 5-*exo* products were formed. In all cases, the Z isomer was very major (Z/E ratio from 7 to 10). In the case of the ^tBu group, opposite stereo-selectivity was observed. Steric congestion favored here the E product (E/Z of 11.5). Reaction monitoring indicated that it is not due to isomerization of **19b** over prolonged heating, but the same E/Z ratio being observed from the beginning of the cyclization. From a mechanistic viewpoint, the Z-selectivity observed with **15a–18a** is consistent with the outer-sphere approach outlined in Scheme 2. The



enolate adds to the π -activated alkyne *trans* to Pd, resulting in the *Z* product after proto-depalladation.³⁰ This *Z*-selectivity markedly contrasts with the *E*-selectivity induced by the In(OTf)₃/DBU catalyst (with R = Me, ⁿBu).²⁴ In the latter case, the reaction was proposed to proceed by *syn* carbometallation, with the In(III) center simultaneously activating the β -dicarbonyl and alkyne moieties.³¹ Formation of the *E* product **19b** most likely results from isomerization of the vinyl-Pd intermediate **B** prior to proto-depalladation, due to steric congestion.³²

Conclusions

When combined with Mg(OTf)₂ and HFIP, the indenediide-based SCS Pd pincer complex was found to be a very powerful catalyst for Conia-ene cyclizations. It works with a variety of substrates, β -ketoesters as well as β -dinitriles and β -diesters, and gives access with complete regio-selectivity to 5, 6 and 7-membered carbo- as well as heterocycles. It is robust and operates at low catalytic loadings (down to 0.1 mol% Pd and 0.5 mol% Mg). Most remarkably, it is also active and selective with substrates featuring internal alkynes. Whatever the substituent at the terminal position is (1°/2°/3° alkyl, aryl), 5-*exo* fully prevails over 6-*endo* cyclization. Moreover, except for the highly congested ^tBu product, the reaction occurs with high *Z*-selectivity. This behavior is in contrast with the *E*-selectivity observed with In-based catalysts, the other known catalysts for internal alkynes. This is due to the way the C–C bond formation and cyclization proceeds with the dual Pd/Mg catalyst; the Mg-chelated enolate attacks the π -coordinated C \equiv C triple bond *trans* to Pd.

These results expand the scope of MLC catalysis in terms of metal (the potential of Pd complexes in MLC catalysis is clearly under-exploited) and catalytic transformations (while hydrogenation/dehydrogenation processes have been extensively developed, examples of C–C bond forming reactions are very rare). Moreover, this work introduces a new multi-cooperative catalytic approach combining MLC and Lewis acid activation. The MLC/LA synergy parallels and complements that involving MLC and proton shuttling. It also represents a new type of dual activation, in which the TM catalyst displays MLC behavior (towards the enolizable C–H bond in this case) while the LA activates another reactive site of the substrate (the β -dicarbonyl motif). Future studies will seek to explore the generality of such MLC/LA synergy and to exploit it in other catalytic transformations.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Financial support from the Centre National de la Recherche Scientifique and the Université de Toulouse is gratefully acknowledged. E. M. thanks the European Commission for a MCIF (Au-MLC 841877). A. C. thanks the French government (MESRI, Ministère de l'Enseignement Supérieur, de la Recherche et de l'Innovation) for his PhD funding. The NMR service of

ICT is acknowledged for the ADEQUATE and NOESY NMR experiments (M. Vedrenne, P. Lavedan) and for the dsel-HSQMBC-IPAP experiments (Dr S. Massou).

Notes and references

- (a) Y. J. Park, J. W. Park and C. H. Jun, *Chem. Res.*, 2008, **41**, 222–234; (b) A. E. Allen and D. W. C. MacMillan, *Chem. Sci.*, 2012, **3**, 633–658; (c) Z. Du and Z. Shao, *Chem. Soc. Rev.*, 2013, **42**, 1337–1378; (d) *Cooperative Catalysis*, ed. R. Peters, Wiley-VCH, Weinheim, 2015; (e) S. Afewerki and A. Córdova, *Chem. Rev.*, 2016, **116**, 13512–13570; (f) D. S. Kim, W. J. Park and C. H. Jun, *Chem. Rev.*, 2017, **117**, 8977–9015.
- (a) J. R. Khusnutdinova and D. Milstein, *Angew. Chem., Int. Ed.*, 2015, **54**, 12236–12273; (b) T. Higashi, S. Kusumoto and K. Nozaki, *Chem. Rev.*, 2019, **119**, 10393–10402.
- Pincer Compounds, Chemistry and Applications*, ed. D. Morales-Morales, Elsevier, 1st edn, 2018.
- (a) C. Gunanathan and D. Milstein, *Science*, 2013, **341**, 1229712; (b) C. Gunanathan and D. Milstein, *Chem. Rev.*, 2014, **114**, 12024–12087; (c) R. H. Crabtree, *Chem. Rev.*, 2017, **117**, 9228–9246; (d) L. Alig, M. Fritz and S. Schneider, *Chem. Rev.*, 2019, **119**, 2681–2751.
- Selected examples: (a) S. Kuwata and T. Ikariya, *Chem.–Eur. J.*, 2011, **17**, 3542–3556; (b) S. Tobisch, *Chem.–Eur. J.*, 2012, **18**, 7248–7262; (c) A. Nerush, M. Vogt, U. Gellrich, G. Leitus, Y. Ben-David and D. Milstein, *J. Am. Chem. Soc.*, 2016, **138**, 6985–6997; (d) U. K. Das, S. Chakraborty, Y. Diskin-Posner and D. Milstein, *Angew. Chem., Int. Ed.*, 2018, **57**, 13444–13448; (e) S. Chakraborty, P. Daw, Y. Ben David and D. Milstein, *ACS Catal.*, 2018, **8**, 10300–10305; (f) J. Bruffaerts, N. von Wolff, Y. Diskin-Posner, Y. Ben-David and D. Milstein, *J. Am. Chem. Soc.*, 2019, **141**, 16486–16493.
- P. Oulié, N. Nebra, N. Saffon, L. Maron, B. Martin-Vaca and D. Bourissou, *J. Am. Chem. Soc.*, 2009, **131**, 3493–3498.
- (a) A. Scharf, I. Goldberg and A. Vigalok, *J. Am. Chem. Soc.*, 2013, **135**, 967–970; (b) A. Scharf, I. Goldberg and A. Vigalok, *Inorg. Chem.*, 2013, **53**, 12–14; (c) J. Takaya and N. Iwasawa, *Chem.–Eur. J.*, 2014, **20**, 11812–11819; (d) A. Scharf, I. Goldberg, A. Vigalok and A. Vedernikov, *Eur. J. Inorg. Chem.*, 2015, **28**, 4761–4768; (e) V. Salamanca, A. Toledo and A. C. Albéniz, *J. Am. Chem. Soc.*, 2018, **140**, 17851–17856; (f) H. Kameo, J. Yamamoto, A. Asada, H. Nakazawa, H. Matsuzaka and D. Bourissou, *Angew. Chem., Int. Ed.*, 2019, **58**, 18783–18787; (g) M. Virant, M. Mihelač, M. Gazvoda, A. E. Cotman, A. Frantar, B. Pinter and J. Košmrlj, *Org. Lett.*, 2020, **22**, 2157–2161.
- (a) N. Nebra, J. Monot, R. Shaw, B. Martin-Vaca and D. Bourissou, *ACS Catal.*, 2013, **3**, 2930–2934; (b) N. Á. Espinosa-Jalapa, D. Ke, N. Nebra, L. Le Goanvic, S. Mallet-Ladeira, J. Monot, B. Martin-Vaca and D. Bourissou, *ACS Catal.*, 2014, **4**, 3605–3611; (c) D. Ke, N. Á. Espinosa, S. Mallet-Ladeira, J. Monot, B. Martin-Vaca and D. Bourissou, *Adv. Synth. Catal.*, 2016, **358**, 2324–2331.
- J. Monot, P. Brunel, C. E. Kefalidis, N. Á. Espinosa-Jalapa, L. Maron, B. Martin-Vaca and D. Bourissou, *Chem. Sci.*, 2016, **7**, 2179–2187.



- 10 (a) J. Li, Y. Shiota and K. Yoshizawa, *J. Am. Chem. Soc.*, 2009, **131**, 13584–13585; (b) S. Qu, Y. Dang, C. Song, M. Wen, K. W. Huang and Z. X. Wang, *J. Am. Chem. Soc.*, 2014, **136**, 4974–4991; (c) X. Yang, *ACS Catal.*, 2014, **4**, 1129–1133; (d) S. Chakraborty, P. O. Lagaditis, M. Forster, E. A. Bielinski, N. Hazari, M. C. Holthausen, W. D. Jones and S. Schneider, *ACS Catal.*, 2014, **4**, 3994–4003; (e) U. Gellrich, J. R. Khusnutdinova, G. M. Leitun and D. Milstein, *J. Am. Chem. Soc.*, 2015, **137**, 4851–4859.
- 11 N. E. Smith, W. H. Bernskoetter and N. Hazari, *J. Am. Chem. Soc.*, 2019, **141**, 17350–17360.
- 12 D. Hack, M. Blumel, P. Chauhan, A. R. Philipps and D. Enders, *Chem. Soc. Rev.*, 2015, **44**, 6059–6093.
- 13 See recent reviews: (a) G. Bouhadir and D. Bourissou, *Chem. Soc. Rev.*, 2016, **45**, 1065–1079; (b) W. Guan, G. Zeng, H. Kameo, Y. Nakao and S. Sakaki, *Chem. Rec.*, 2016, **16**, 2405–2425; (c) D. You and F. P. Gabbaï, *Trends Chem.*, 2019, **1**, 485–496; (d) J. Becica and G. E. Dobereiner, *Org. Biomol. Chem.*, 2019, **17**, 2055–2069.
- 14 (a) B. K. Corkey and F. D. Toste, *J. Am. Chem. Soc.*, 2005, **127**, 17168–17169; (b) A. Matsuzawa, T. Mashiko, N. Kumagai and M. Shibasaki, *Angew. Chem., Int. Ed.*, 2011, **50**, 7616–7619; (c) S. Suzuki, E. Tokunaga, D. S. Reddy, T. Matsumoto, M. Shiro and N. Shibata, *Angew. Chem., Int. Ed.*, 2012, **51**, 4131–4135; (d) P. Barrio, M. Kumar, Z. Lu, J. Han, B. Xu and G. B. Hammond, *Chem.–Eur. J.*, 2016, **22**, 16410–16414; (e) M. Cao, A. Yesilcimen and M. Wasa, *J. Am. Chem. Soc.*, 2019, **141**, 4199–4203; (f) T. Horibe, M. Sakakibara, R. Hiramatsu, K. Takeda and K. Ishihara, *Angew. Chem., Int. Ed.*, 2020, **38**, 16470–16474.
- 15 See the ESI.†
- 16 For related transformations, see: (a) M. Li and R. Hua, *J. Org. Chem.*, 2008, **73**, 8658–8660; (b) Y. Oonishi, A. Gómez-Suárez, A. R. Martin and S. P. Nolan, *Angew. Chem., Int. Ed.*, 2013, **52**, 9767–9771.
- 17 (a) I. Colomer, A. E. R. Chamberlain, M. B. Haughey and T. J. Donohoe, *Nat. Rev. Chem.*, 2017, **1**, 0088; (b) C. Yu, J. Sanjosé-Orduna, F. W. Patureau and M. H. Pérez-Temprano, *Chem. Soc. Rev.*, 2020, **49**, 1643–1652; (c) V. Pozhydaev, M. Power, V. Gandon, J. Moran and D. Leboeuf, *Chem. Commun.*, 2020, **56**, 11548–11564.
- 18 Milstein reported very recently a tandem catalytic approach for the oxidation of alkenes by water. In(OTf)₃ promotes the hydration of the alkene and a PNP Ru pincer complex and then promotes dehydrogenation: S. Tang, Y. Ben-David and D. Milstein, *J. Am. Chem. Soc.*, 2020, **142**, 5980–5984.
- 19 For reactions of organic and metal-based electrophiles on the indenediide ligand backbone, see: (a) P. Oulié, N. Nebra, S. Ladeira, B. Martin-Vaca and D. Bourissou, *Organometallics*, 2011, **30**, 6416–6422; (b) N. Nebra, N. Saffon, L. Maron, B. Martin-Vaca and D. Bourissou, *Inorg. Chem.*, 2011, **50**, 6378–6383; (c) N. Nebra, S. Ladeira, L. Maron, B. Martin-Vaca and D. Bourissou, *Chem.–Eur. J.*, 2012, **18**, 8474–8481.
- 20 ³¹P NMR monitoring showed a rapid reaction between [(SCS)Pd]₂ and Cu(OTf)₂, while no change was observed for the other salts.
- 21 Enabling direct reaction monitoring by NMR spectroscopy; Mg(OTf)₂ has also a practical advantage over Yb(OTf)₃.
- 22 A. Ochida, H. Ito and M. Sawamura, *J. Am. Chem. Soc.*, 2006, **128**, 16486–16487.
- 23 6-Exo cyclization of a β-keto,ester derived from cyclopentanone was also reported *via* gold catalysis: J. J. Kennedy-Smith, S. T. Staben and F. D. Toste, *J. Am. Chem. Soc.*, 2004, **126**, 4526–4527.
- 24 K. Takahashi, M. Midori, K. Kawano, J. Ishihara and S. Hatakeyama, *Angew. Chem., Int. Ed.*, 2008, **47**, 6244–6246.
- 25 The formation of medium to large-sized carbocycles from ω-alkynyl-β-keto,esters has been reported using In(NTf₂)₃ as the catalyst: (a) H. Tsuji, K. I. Yamagata, Y. Itoh, K. Endo, M. Nakamura and E. Nakamura, *Angew. Chem., Int. Ed.*, 2007, **46**, 8060–8062; (b) Y. Itoh, H. Tsuji, K. I. Yamagata, K. Endo, I. Tanaka, M. Nakamura and E. Nakamura, *J. Am. Chem. Soc.*, 2008, **130**, 17161–17167.
- 26 (a) H. Ito, Y. Makida, A. Ochida, H. Ohmiya and M. Sawamura, *Org. Lett.*, 2008, **10**, 5051–5054; (b) W. Chaladaj, A. Kolodziejczyk and S. Domanski, *ChemCatChem*, 2017, **9**, 4334–4339.
- 27 S. T. Staben, J. J. Kennedy-Smith and F. D. Toste, *Angew. Chem., Int. Ed.*, 2004, **43**, 5350–5352.
- 28 W. Hess and J. W. Burton, *Adv. Synth. Catal.*, 2011, **353**, 2966–2970.
- 29 Upon Conia-ene cyclization of β-ketoesters featuring internal alkynes (R = Me, ⁿBu, ⁱPr, Ph) with a Au(i) complex, *anti* addition and *Z*-selectivity were also observed, but the regio-selectivity was poor (mixtures of 5-*exo*/6-*endo* were formed). See ref. 26a.
- 30 For a review in proto-depalladation, see: M. L. O'Duill and K. M. Engle, *Synthesis*, 2018, **50**, 4699–4714.
- 31 Carbometalation-type mechanisms have also been proposed to account for the *syn* addition observed with Zn (see ref. 28) and Cu catalysts: S. Montel, D. Bouyssi and G. Balme, *Adv. Synth. Catal.*, 2010, **352**, 2315–2320.
- 32 For examples of vinyl-Pd *Z/E* isomerization, see: (a) C. Amatore, S. Bensalem, S. Ghalem and A. Jutand, *J. Organomet. Chem.*, 2004, **689**, 4642–4646; (b) N. Panda and R. Mothkuri, *J. Org. Chem.*, 2012, **77**, 9407–9412; (c) J. Aziz, G. Frison, P. Le Menez, J. D. Brion, A. Hamze and M. Alami, *Adv. Synth. Catal.*, 2013, **355**, 3425–3436.

