



Cite this: *Catal. Sci. Technol.*, 2015, 5, 1447

Received 12th January 2015,
Accepted 3rd February 2015

DOI: 10.1039/c5cy00058k

www.rsc.org/catalysis

The use of a custom-made palladium complex for the telomerisation of isoprene with alcohols under green conditions is described. Using a very low catalyst loading (0.1 mol%), this protocol allows for the reactions to be carried out without the need of high pressure equipment, at room temperature and under solventless conditions, affording high yields of telomerization products with high selectivity towards linear head-to-head and tail-to-head telomers.

Introduction

During the last fifty years, organic synthesis has reached a high level of sophistication with nearly no limits, allowing for the synthesis of highly complex molecules through protocols that provide specific chemo-, regio- and stereoselectivity. Nowadays, the main issue is the efficiency of a synthesis, which can be measured as the increase of complexity per transformation.¹ At the same time, there are economic and environmental factors that demand high atom efficiency, low E-factor (mass ratio of waste to desired product) and a decrease of waste generated, in accordance to the principles of green chemistry.² The development of synthetic protocols that can account for all these requirements is a challenge that in many cases can be only accomplished using catalysis, in any of its forms.³

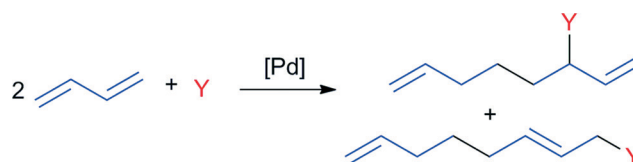
Telomerizations are 2 bond-forming, 100% atom-efficient reactions that involve the metal-catalyzed oligomerization of 1,3-dienes (also known as taxogens) followed by the addition of a nucleophile (or telogen) to achieve substituted longer chains (Scheme 1).⁴ The telomerization reaction has been a key route in the preparation of interesting industrial products

Room temperature, solventless telomerization of isoprene with alcohols using (N-heterocyclic carbene)–palladium catalysts†

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since it was discovered in 1967 by Smutny⁵ and Takahashi,⁶ independently. A vast range of products can be obtained due to the large variety of substrates and nucleophiles available. Butadiene and isoprene (2-methyl-1,3-butadiene) are by far the most popular dienes used due to the plethora of applications of the resulting products, although other taxogenes like myrcene have been used.⁷ The most common nucleophiles are water and alcohols, but other interesting products are obtained using other nucleophiles such as polyols,⁸ amines,⁹ carbon dioxide¹⁰ or acids.^{11,12} At a much smaller scale, telomerization reactions have also been applied in the synthesis of natural products.¹³

Isoprene is a common volatile diene naturally produced by many plants. Its skeleton is part of the terpenes family, which can be considered as chains consisting of multiple units of isoprene.¹⁴ The telomerization of isoprene is a more challenging reaction than the telomerization of butadiene, since isoprene is less reactive. In addition, not only isoprene is a non-symmetric molecule, meaning that the approach of the two units can be done in four different manners to give the head-to-head (H2H), head-to-tail (H2T), tail-to-head (T2H) and tail-to-tail (T2T) telomers, but also the attack of the nucleophile can occur in two different positions (Scheme 2). Therefore, eight different telomers are likely to form, as well as four dimers when no nucleophilic attack takes place, and trimers or oligomers if more than two isoprene units combine. Selectivity becomes an important goal to avoid separation of complex mixtures of products, whose compositions are strongly influenced by the catalyst employed and the reaction parameters, such as temperature and solvents.¹⁵



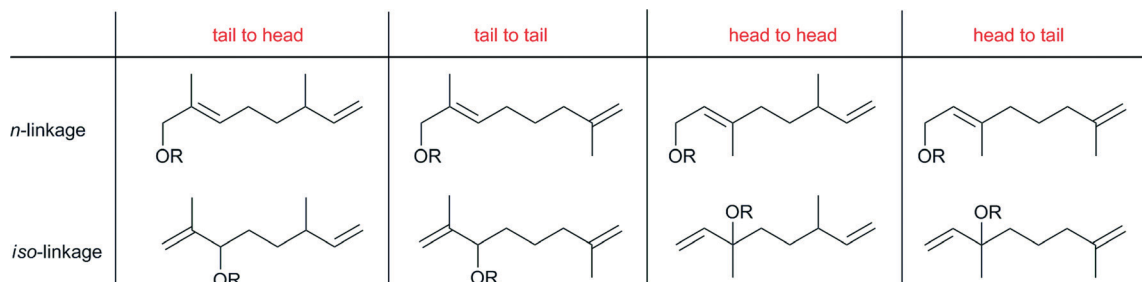
Scheme 1 Telomerization of 1,3-butadiene with a nucleophile Y.

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† Electronic supplementary information (ESI) available: ¹H and ¹³C NMR spectra of **1b**, **2b**, characterization of telomers. CCDC 1039375 (**1b**) and 912794 (**2b**). For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c5cy00058k



Scheme 2 Telomers from the telomerization of isoprene with ROH.

Palladium catalysts have been by far the most widely employed in this reaction. Protocols in which a palladium complex and a phosphorous ligand are combined *in situ* are the most common in the literature.¹⁶ The substitution of phosphines by NHCs (N-heterocyclic carbenes)¹⁷ has shown, in general, improved activity and selectivities.^{15,18} More recently, the use of well-defined (NHC)-bearing Pd complexes in the telomerization of dienes with alcohols has allowed for a much easier handling, better control of metal to ligand stoichiometry and significant improvements in activity and chemo- and regioselectivity.¹⁹ This is especially pronounced for complexes bearing NHCs with mesityl groups as N-substituents.^{15,19–21} Our group recently developed a new series of (NHC)-palladium complexes with the general formula (NHC)PdCl₂(TEA) (**1a**, **2a**, Fig. 1),²² which have proven to be very active pre-catalysts for cross-coupling reactions such as Suzuki–Miyaura and Buchwald–Hartwig²³ even under very mild reaction conditions. We decided to extend their application to telomerization reactions and therefore we synthesized two new complexes **1b** and **2b**, bearing mesityl N-substituents.

Results and discussion

The new complexes **1b** and **2b** were prepared in excellent yields following the standard procedure for the synthesis of complexes **1a** and **2a** (Scheme 3). Both complexes were fully characterized by means of elemental analysis and ¹H and ¹³C NMR spectroscopy. Crystals suitable for X-ray diffraction for each complex were obtained from dichloromethane/hexane solutions.²⁴ The solid-state structures are depicted in Fig. 2 and 3 showing, as expected, square planar conformations for both complexes with the NHC *trans* to TEA. As it is customary for (NHC)-Pd complexes, the Pd–C_{carbenic} distance is in the

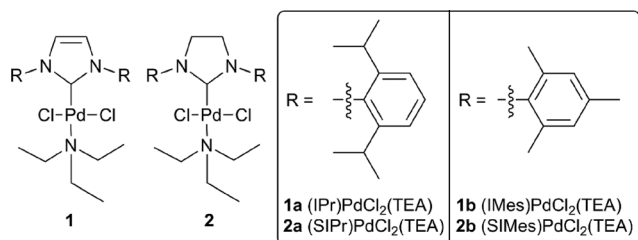
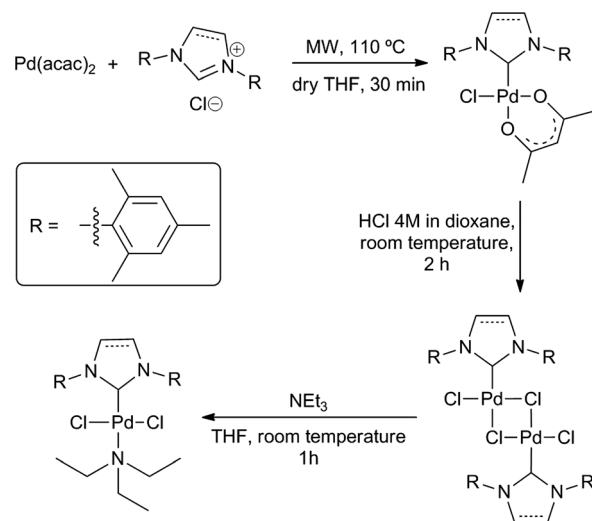


Fig. 1 Pre-catalysts tested in this study.



Scheme 3 General procedure for the synthesis of (NHC)PdCl₂(TEA) complexes.

range of a single bond, with very similar values for **1b** and **2b**. Interestingly, both Pd–N1 distances are also very similar, implying a similar extent of electron donation from the NHC to the Pd centre.

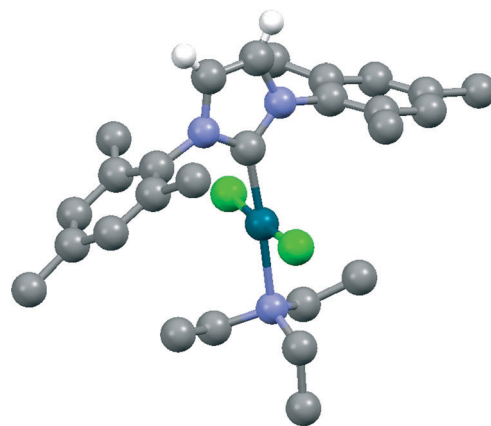


Fig. 2 Crystal structure of (IMes)PdCl₂(TEA) (**1b**). Hydrogen atoms are omitted for clarity except those on the backbone of the NHC. Selected distances (Å): Pd–C_{carbenic}, 1.968(2); Pd–Cl1, 2.2931(6); Pd–Cl2, 2.3250(6); Pd–N1, 2.206(2).



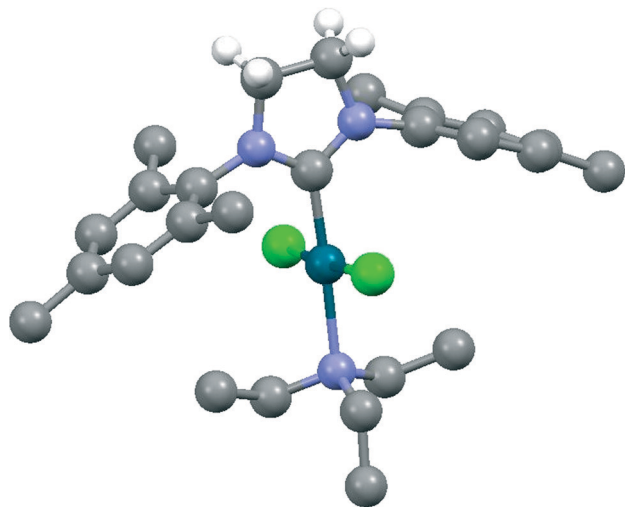


Fig. 3 Crystal structure of (SIMes)PdCl₂(TEA) (**2b**). Hydrogen atoms are omitted for clarity except those on the backbone of the NHC. Selected distances (Å): Pd–C_{carbenic}, 1.974(4); Pd–Cl1, 2.3088(10); Pd–Cl2, 2.3013(10); Pd–N1, 2.198(3).

A review of the literature revealed that Pd-catalyzed telomerization reactions are performed under an inert atmosphere, usually in a pressure vessel or an autoclave, with temperatures ranging in most cases from 40 to 90 °C.^{16–21} Our previous experience with this type of complexes in catalytic cross-coupling reactions prompted us to attempt our telomerizations at 25 °C. Also, since different solvents, like THF or *n*-hexane, have proven to have some influence on the selectivity, we decided to carry out our reactions without solvent or excess of any of the coupling partners. In addition and to make our catalytic system as user-friendly as possible, we performed our reactions in regular glass pressure vials fitted with a screw-cap with a septum, using a low catalyst loading of 0.1 mol%.²⁵ All the reagents except isoprene were loaded in open air. It is important to remark that all the reagents were used as received without further purification or drying. After closing it, the air in the vial was flushed out with N₂, followed by the injection of the corresponding amount of isoprene through the septum. The reactions were carried out at room temperature and stopped after 72 h, when they were analyzed by gas-chromatography/mass-spectrometry using pentadecane as internal standard. An

initial comparison of the performance of complexes **1a–2b** in the telomerization reaction of isoprene with MeOH was performed (Table 1). In all the cases, only *n*-linkage products were obtained. The conversion of isoprene to products was very high regardless of the complex utilized, but only those bearing unsaturated carbenes led to telomerization products (Table 1, entries 3, 4). (IMes)PdCl₂(TEA) showed to be the most selective towards the formation of a given telomer, affording 83% of the T2H and H2H product (88% yield of all combined telomers). On the other hand, SIMes- and SIPr-bearing complexes resulted to be efficient catalysts for the synthesis of oligomers and polymers that are formed by condensation of at least three isoprene units, posing as potential catalysts for isoprene polymerization protocols. These results are consistent with previous findings in the literature, since bulkier carbenes such as SIPr have been reported to promote trimerization of isoprene producing sesquiterpenes.¹⁵

The promising results accomplished with complex **1b** encouraged us to test the scope of the reaction of isoprene with different alcohols. The same procedure used for the telomerisation of isoprene with methanol was applied for ethanol, 1- and 2-propanol and butanol (Table 2). The mixtures of telomers were isolated by distillation, analyzed by GC/MS, ¹H and ¹³C NMR and identified by comparison with previously reported data (see ESI†). In all cases, with the exception of MeOH, this protocol allows for the selective formation of only two of the telomers, T2H and H2H. In general, the results showed that when the length of the alcohol chain increases, the conversion decreases, probably as a consequence of the increased steric hindrance of bulkier nucleophiles, as mechanistic studies suggest.¹⁵ The selectivity towards telomerization *vs.* dimerization and polymerization also decreased from methanol to butanol. Interestingly, while in most cases the H2H telomer is obtained as the major telomerization product, the selectivity switches towards the T2H when ¹PrOH is the nucleophile, which could also be a consequence of the bulk of the alcohol and the interactions with the corresponding dimers. To the best of our knowledge, these are the highest values of TON for telomerizations of isoprene with alcohols carried out at room temperature.

The reaction of isoprene with methanol using complex **1b** was also attempted without an inert atmosphere, under the same conditions. Although the conversion to products

Table 1 Activity comparison in the solventless telomerization of isoprene with MeOH

$\text{Isoprene (2 equiv)} + \text{MeOH (1 equiv)} \xrightarrow[\text{NaOMe, 0.2 equiv, room temperature, 72 h}]{\text{complex, 0.1 mol\%}^a} \text{products}$									
Entry	Complex	Conversion ^b (%)	H2T ^c (%)	T2T ^c (%)	T2H ^c (%)	H2H ^c (%)	Dimers ^c (%)	Trimers ^c (%)	Polymers ^c (%)
1	2a	99	—	—	—	—	10	13	77
2	2b	89	—	—	—	—	1	1	98
3	1b	99	—	5	7	76	4	6	2
4	1a	99	—	15	2	29	32	8	15

^aCatalyst loading calculated as (mmol catalyst × 100)/(mmol alcohol).^b Conversion of isoprene to products. ^c Percentage of component in the product mixture.



Entry	R	Conversion ^c (%)	TON ^c	H2T ^d (%)	T2T ^d (%)	T2H ^d (%)	H2H ^d (%)	Dimers ^d (%)	Oligomers ^d (%)
1	Me	99	871	—	5	7	76	4	8
2	Et	91	637	—	—	27	43	24	6
3	Pr	80	616	—	—	30	47	18	5
4	iPr	64	358	—	—	53	3	22	22
5	Bu	75	443	—	—	14	45	20	21

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Acknowledgements

We thank the EPSRC UK National Crystallography Service at the University of Southampton for the collection of the crystallographic data for **1b**. O.N. thanks the National Science Foundation (CHE-0924324) for funding initial stages of this work.

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