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# From biodiversity to catalytic diversity: how to control the reaction mechanism by the nature of metallophytes

Vincent Escande · Tomasz K. Olszewski · Claude Grison

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**Abstract** Phytoextraction is widely used for the reclamation of degraded sites, particularly to remove trace metals from contaminated soils. Whereas this technique demonstrates several advantages, the biomass resulting from phytoextraction processes is highly enriched in metallic elements and constitutes therefore a problematic waste. We show here that this biomass can be used for the preparation of novel polymetallic extracts, with high potential as catalysts or reagents in organic synthesis. This new concept of ecocatalysis constitutes an innovative recycling of metallic elements whose current known reserves could be exhausted in the coming decades. The ecocatalysts Eco-Zn and Eco-Ni prepared respectively from Zn and Ni hyperaccumulating plants display two distinct chemical reactivities, starting from the same substrates. Eco-Zn led to the formation of esters of commercial interest for the fragrance industry, following a hydro-acyloxy-addition reaction pathway. In contrast, Eco-Ni afforded chlorinated products thanks to the hydrochlorination of alkenes. Both ecocatalysts allowed the synthesis of valuable products in high yields through methodologies in line with the spirit of sustainable chemistry.

**Keywords** Phytoextraction · Green chemistry · Ecocatalysis · Lewis acid catalysis · Metal-hyperaccumulating plants · Chemodiversity

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V. Escande · C. Grison (✉)  
FRE 3673 – Bioinspired chemistry and ecological innovation –  
CNRS, University of Montpellier 2, Stratoz – Cap Alpha, Avenue de  
l'Europe, 34830 Clapiers, France  
e-mail: [claude.grison@cnrs.fr](mailto:claude.grison@cnrs.fr)

T. K. Olszewski  
Faculty of Chemistry, Wrocław University of Technology, Wybrzeże  
Wspianskiego 27, 50-370 Wrocław, Poland

## Introduction

Phytoextraction of trace metals: environmental opportunities and innovative sustainable chemistry

The depletion of world reserves of metals has become a very serious problem, both from an economic and environmental point of view (Hunt et al. 2013). Due to an increasing demand for consumer goods and the continuous increase of the world population, there is a risk that the available reserves of many metals, widely used in industry, will be exhausted in less than 50 years (Hunt et al. 2013; Salazar 2013). Among those metals threatened with depletion, many of them are frequently used in synthetic industrial processes as catalysts. This is particularly the case of zinc, nickel, manganese, and platinum group elements (PGE) (Lloyd 2011; Yamamoto 2000). To cope with these risks of depletion, new recycling strategies are implemented (United Nations Environment Programme (UNEP) and United Nations University 2009), but today many metals are still recycled in less than 50 % (Hunt et al. 2013). Moreover, the need to maintain a constant supply of these metallic resources leads metal mining industry to develop new operations, which is a source of environmental and social impacts of high concern (Bridge 2004; Thornton 1996).

On the other hand, to remedy soil pollution caused by mining, phytoremediation programs have been developed in many countries (Chaney et al. 2000; Raskin and Ensley 2000; Singh et al. 2003). Among the techniques used in phytoremediation, the phytoextraction is often applied, a technique that relies on the use of so-called metal-hyperaccumulating plants or metallophytes, able to extract metallic trace elements (trace metals, TM) from the soil and concentrate them in their leaves (Anderson 2013). Phytoremediation processes are now among the most frequently used to restore the former mining sites, because of

their low cost of implementation, low environmental impact, and good public acceptance (Bhargava et al. 2012). However, numerous efforts are still necessary to exploit the ability of recovering and recycling metals from contaminated biomass produced during phytoextraction and fully develop the economic potential of this technique. Thus, innovative technologies are needed to receive the full benefit of the remediation process.

The existence of the aforementioned problems, namely pollution caused by the mining industry on the one hand and the depletion of metallic resources on the other hand, has led us to seek a solution to enhance the utility of the biomass issued from the phytoextraction processes and to improve the resulting economic opportunities. Therefore, we have developed a new range of catalysts for use in organic synthesis, called “ecocatalysts,” from biomass rich in metallic elements accumulated in the plant tissue issued after phytoextraction (Grison and Escarré 2011; Grison and Escande 2013a, b; 2014).

#### Use of the ecocatalysts in synthesis of valuable products for the fine chemical industry

Herein, we show the synthetic potential of ecocatalysts, underlying the fact that it is possible to control the reaction mechanism by choosing the right type of ecocatalyst, and the nature of metallophyte from which it derives. Thus, the presented here ecocatalysts Eco-Zn or Eco-Ni may promote different reaction pathways, based on the same substrates. These examples illustrate the importance of ecocatalysts in organic synthesis and the diversity of products possible to obtain according to ecocatalyst used. The catalysts used in this study are generated from biomass, issued from metal-hyperaccumulating plants, using the procedure discovered in our laboratory (Grison and Bès 2013; Saunier et al. 2013).

Eco-Zn was prepared from biomass derived from hyperaccumulating plants grown in France (Saint-Laurent-le-Minier, Gard) in order to restore old zinc mining sites. The catalyst Eco-Ni was prepared from nickel-hyperaccumulating plants grown in New Caledonia (South province), on sites degraded by the nickel mining industry. We sought to use these catalysts in chemical reactions of high economic interest and particularly for industrial production of fine chemicals.

At present, the chemical industry is changing to adapt its processes to new environmental requirements (Adams et al. 2013; Matus et al. 2010; Wilson and Schwarzman 2009). This transition towards new eco-compatible processes is particularly desirable in the field of fine chemicals, which produces large amounts of waste per kilogram of final product and use solvents, reagents, catalysts, and toxic pollutants (Federsel 2013; Watson 2012).

More specifically, the industrial sector of fragrances and cosmetics appears to be very concerned about the environmental compatibility of its processes, in particular for reasons of communication vis-à-vis its customers (Dobbs et al. 2000; Kraft et al. 2005; Leseurre et al. 2014).

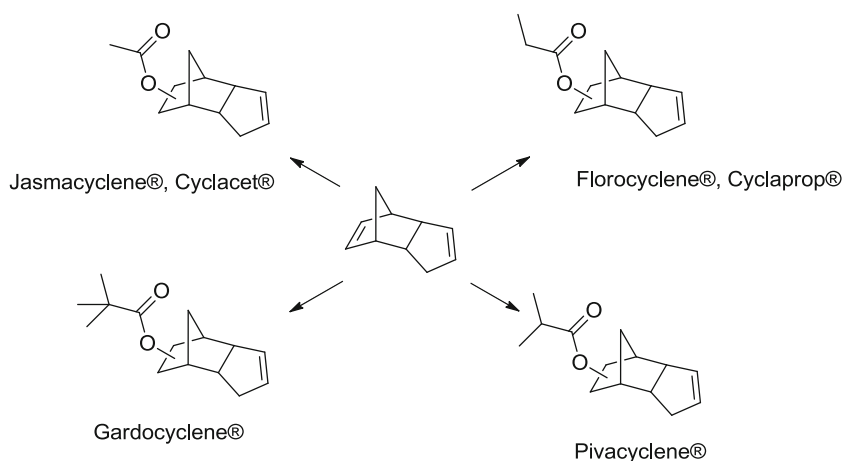
#### Catalysis of the hydro-acyloxy-addition reaction: context and advantages of the use of ecocatalysts

Due to the high potential of our ecocatalysts as Lewis acids, revealed in our previous works (Escande et al. 2014a, 2014b, 2014c; Grison et al. 2013), we became interested in the hydro-acyloxy-addition reaction known to produce valuable products under Lewis acid catalysis (Wunderly and Sowa 1937). This reaction is the source of many starting materials for the industry of perfumes and cosmetics, including verdyl acetate issued from dicyclopentadiene (Sell 2006), commercialized by Givaudan under the name Jasmacyclene® (Fig. 1).

A detailed study of this reaction has been published in various articles by Peterson and co-workers (Peterson 1960; Peterson and Allen 1962, 1963; Peterson and Tao 1964). Thus, the use of different catalysts for that transformation was reported thus far in the literature such as follows: boron trifluoride (Wunderly and Sowa 1937), Pd- and Ti(III)-based catalysts (Annby et al. 1993; Ferraz and Ribeiro 1992; Larock and Hightower 1993), sulfamic acid (Wang et al. 2004), triflic acid (Li et al. 2006; Rosenfeld et al. 2006), ferric triflate (Choi et al. 2008), copper(II) triflate (Taylor et al. 2005), indium(III) triflate (Chen and Lu 2007), cerium(IV) sulfate (Horiuchi et al. 2003), and ionic liquids (Gu et al. 2004). Although these systems effectively catalyze the hydro-acyloxy-addition reaction, they have certain drawbacks such as the use of metals whose reserves will be exhausted in the coming decades, the necessity of handling hazardous reagents and pollutants or the use of non-environmentally friendly solvents and nonreusable catalysts. Some improvements have been made by the use of heterogeneous catalysis supported on montmorillonite (Ballantine et al. 1981; Yang et al. 2006) and silica gel (Hinze et al. 2009). These applications, however, are substrate-dependent and provide products with moderate yields. The development of efficient and reusable catalytic systems based on abundant resources and the handling of which represents no risk for the environment is therefore highly desirable.

The characteristics of our ecocatalysts Eco-Zn and Eco-Ni perfectly meet these requirements. Additionally, this study highlights an unexpected reaction pathway, leading to different products starting from the same substrates, by simply changing the polymetallic composition of the ecocatalysts prepared from biomass from different metal-hyperaccumulating plants.

**Fig. 1** Starting materials for fragrance industry prepared via hydro-acyloxy-addition reaction from dicyclopentadiene



## Materials and methods

### Preparation of ecocatalysts

#### *Eco-Zn from Zn hyperaccumulating plants*

Leaves of *Noccaea caerulescens* (Brassicaceae) and *Anthyllis vulneraria* (Fabaceae) were harvested before flowering in Les Avinières, a former zinc mining site in Saint-Laurent-Le-Minier, southern France (same mass of the two plants), air-dried and crushed. The obtained solid (30 g) was heated to 400 °C for 5 h and the resulting powder (5 g) was added to 50 mL of 1 M HCl solution. The solution was heated at 60 °C and stirred for 2 h. The reaction mixture was filtered on a pad of celite. The resulting yellow solution, composed of different metal chlorides, was then concentrated under vacuum, yielding Eco-Zn (5.1 g). The catalyst was kept in a desiccator under vacuum.

#### *Eco-Ni from Ni-hyperaccumulating plants*

Leaves of *Geissois pruinosa* (Cunoniaceae) were harvested in the South province of New Caledonia, on Chemin de Prony, near Noumea. The leaves were air-dried and crushed, then the obtained solid (42 g) was heated to 400 °C for 5 h and the resulting powder (5 g) was added to 50 mL of 1 M HCl solution. The solution was heated at 60 °C and stirred for 2 h. The reaction mixture was filtered on a pad of celite. The resulting green solution, composed of different metal chlorides, was then concentrated under vacuum, yielding Eco-Ni (3.38 g). The catalyst was kept in a desiccator under vacuum.

### Characterization of ecocatalysts

Inductively coupled plasma mass spectrometry (ICP-MS) analyses were performed using the metal analysis of total dissolved solutes in water. The samples were acidified with nitric acid 2.5 % and stirred for 30 min. The digestates were

diluted to 0.005 g.L<sup>-1</sup>. Three blanks were recorded for each step of the digestion and dilution procedure on a HR-ICP-MS Thermo Scientific Element XR. Three analyses were performed for each sample in order to determine the standard deviation of the measurement.

X-ray diffraction (XRD) data measurements on the samples dried at 110 °C for 2 h were performed by using a BRUKER diffractometer (D8 advance, with a CuK $\alpha$  radiation  $\lambda=1.54086\text{\AA}$ ) equipped with a LynxEye detector.

FT-IR measurements were carried out using pyridine as probe molecule. A PerkinElmer Spectrum 100 FT-IR spectrometer was used for recording the spectra. Excess gaseous pyridine was adsorbed on the samples, then the samples were degassed for 15 min at 25 °C ( $10^{-3}$  Pa) and a first spectrum was recorded. The samples were then degassed for 15 min at 150 °C ( $10^{-3}$  Pa) in order to eliminate the physisorbed pyridine and a second spectrum was recorded.

The use of ecocatalysts Eco-Zn and Eco-Ni in organic synthesis

### Chemicals

Reagents and solvents were purchased from Sigma-Aldrich, and were used without further purification.

### *Purification and analysis of reaction products*

Flash column chromatography was performed using Chromagel Carlo Erba Reactifs SDS silica 60 A CC 35–70  $\mu\text{m}$ . Reactions were monitored using Merck Kieselgel 60 F254 aluminum. TLCs were visualized by UV fluorescence (254 nm) then by immersion in one of the following dyeing reagents: KMnO<sub>4</sub>, vanillin, ninhydrin, phosphomolybdic acid solution, and phosphotungstic acid solution.

NMR spectra were recorded on a Brüker Avance 300 spectrometer at room temperature, <sup>1</sup>H frequency is at 300 MHz and <sup>13</sup>C frequency is at 75 MHz. IR spectra were

recorded on a PerkinElmer Spectrum 100 FT-IR spectrometer, in ATR mode. GC-MS analyses were performed on a Shimadzu QP2010SE apparatus, equipped with a  $30\text{ m} \times 0.25\text{ mm} \times 0.25\text{ }\mu\text{m}$  ZB-5MSi Guardian column (Phenomenex®) with hydrogen as carrier gas.

### General procedure

All reactions were conducted in sealed vials (4 mL) equipped with magnetic stirring. Silica (1.0 g, Chromagel Carlo Erba Reactifs SDS silica 60 A CC 35–70  $\mu\text{m}$ ), carboxylic acid, alkene, and ecocatalyst (Eco-Zn or Eco-Ni, mass corresponding to amounts indicated in Tables 5, 6, 7) were mixed by grinding in a mortar. The resulting solid was introduced in a sealed vial and heated to 120 °C in sand bath for 2–24 h. Aliquots were taken for analysis in GC-MS and TLC, after dilution in dichloromethane. After completion of the reaction, the solid was washed with cyclohexane ( $3 \times 5\text{ mL}$ ) and filtered. The resulting solution was concentrated under reduced pressure and purified by flash column chromatography. The previous catalytic solid (mixture of ecocatalyst and silica) was recycled by washing with cyclohexane and drying in an oven (120 °C, 5 h), before being reused in the next run.

Synthesized products were identified by comparing their spectroscopic data (IR,  $^1\text{H}$  NMR,  $^{13}\text{C}$  NMR, MS) with those previously published (He and Goldsmith 2010; Magens et al. 2007; Schmidt et al. 1980; Taylor et al. 2005; Wang et al. 2004) or with those of authentic commercial samples. When necessary, the exact position of the introduced substituent was determined by key Heteronuclear Multiple Bond Correlation (HMBC) correlations or by 2D total correlation spectroscopy (TOCSY) experiments.

## Results and discussion

### Characterization of Eco-Zn and Eco-Ni ecocatalyst

After harvest, the Zn hyperaccumulating plants *N. caerulescens* (Brassicaceae) and *A. vulneraria* (Fabaceae) were transformed into Eco-Zn following a procedure designed to be simple to implement. After air drying and thermic treatment at 400 °C, to remove most of the organic matter, and addition of 1 M hydrochloric acid followed by concentration of the resulting solution, a mixture of a polymetallic salts was obtained. Analogous procedure was followed for biomass from the Ni-hyperaccumulating plant *G. pruinosa* (Cunoniaceae). The Eco-Zn and Eco-Ni were characterized by ICP-MS analysis and XRD techniques. Their Lewis acidity was studied by infrared monitoring of the adsorbed pyridine.

### ICP-MS analysis

*ICP-MS analysis of Eco-Zn* The mineral analysis of Eco-Zn was performed by ICP-MS after dilution of the solid in an aqueous solution of 2.5 % nitric acid (Table 1). The results of the analysis revealed that Zn is the main transition metal present in the analyzed polymetallic composition of Eco-Zn (6.57 wt%), and is accompanied mostly by Fe (1.80 wt%) and Al (1.74 wt%) (Table 1).

The reproducibility of the composition of Eco-Zn was studied in our previous article, which showed that the levels of transition elements are only slightly variable between samples of Eco-Zn prepared from batches of different plants (Escande et al. 2014c). Eco-Zn was prepared from most representative batches of the plants to homogenize the content of the transition elements and minimize possible variations in their composition.

*ICP-MS analysis of Eco-Ni* Analogous procedure using ICP-MS was employed to characterize the Eco-Ni catalyst (Table 2).

The analysis showed that Ni is the major transition metal present in the Eco-Ni catalyst (7.39 wt%). Small amounts of other transition metals were also observed, but at much lower levels than in the case of Eco-Zn: Fe (0.79 wt%), Mn (0.16 wt%), and Zn (1.25 wt%). The amount of Al present was in the range of 0.53 wt% whereas the content of Mg (7.63 wt%) was comparable to that observed for Eco-Zn. In turn, the amounts of Ca (11.98 wt%) were two to three times lower than those observed in the case of Eco-Zn.

### XRD analysis

*XRD analysis of Eco-Zn* The X-ray diffraction (XRD) analysis of Eco-Zn showed crystalline species present in the solid (Fig. 2). Various chlorides have been characterized, including the mixed chloride  $\text{CaMg}_2\text{Cl}_6(\text{H}_2\text{O})_{12}$  and  $\text{K}_2\text{ZnCl}_4$  and potassium chloride KCl (Table 3).  $\text{K}_2\text{ZnCl}_4$  is a masked form of  $\text{ZnCl}_2$  as shown in our previous work (Escande et al. 2014a). The presence of other species not detectable by XRD is very likely, since this technique does not reveal the presence of molecules containing, e.g., Fe or Al, detected by the elemental analysis. This is probably due to the non-crystalline nature of these species within the Eco-Zn catalyst, which did not allow their characterization by XRD.

*XRD analysis of Eco-Ni* The XRD analysis of Eco-Ni catalyst revealed the presence of several mixed chlorides:  $\text{CaMg}_2\text{Cl}_6(\text{H}_2\text{O})_{12}$  (also detected in the Eco-Zn catalyst),  $\text{KCaCl}_3$ , and  $\text{KNiCl}_3$  (Fig. 3 and Table 4).  $\text{KNiCl}_3$  is a masked form of  $\text{NiCl}_2$ , having interesting Lewis acidic properties and potential for use in organic synthesis. Several sulfates were also detected, including  $\text{NiSO}_4(\text{H}_2\text{O})_6$  and  $\text{Na}_2\text{SO}_4$ . This

**Table 1** ICP-MS analysis of Eco-Zn

Ecocatalyst	Latitude and longitude data of sampling site	Mg wt% ( $\pm$ SD)	Ca wt% ( $\pm$ SD)	Fe wt% ( $\pm$ SD)	Zn wt% ( $\pm$ SD)	Cd wt% ( $\pm$ SD)	Al wt% ( $\pm$ SD)	Pb wt% ( $\pm$ SD)
Eco-Zn	43.933031, 3.666407	3.76 ( $\pm$ 0.10)	35.81 ( $\pm$ 0.50)	1.80 ( $\pm$ 0.028)	6.57 ( $\pm$ 0.059)	0.35 ( $\pm$ 0.0040)	1.74 ( $\pm$ 0.049)	0.73 ( $\pm$ 0.0048)

observation is consistent with the high levels of sulfate ion observed in *G. pruinosa*, as described in our previous article (Grison et al. 2013). In turn, the presence of potassium chlorate is difficult to explain. Such an inorganic species have been observed previously in similar ecocatalysts (Escande et al. 2014a). A possible explanation can be that the presence of potassium chlorate is due to the decomposition of chlorides during the preparation of the Eco-Ni catalyst, with in situ formation of oxidative species, a fact described in the literature for several transition metal chlorides (Krishna et al. 2003; Mondelli et al. 2011). Thus, the present oxidizing species would then permit the formation of chlorate by oxidation of other salts (Patnaik 2003).

#### *Infrared characterization of the Lewis acidity of Eco-Zn and Eco-Ni by pyridine adsorption*

**Characterization of Eco-Zn** In order to determine the Lewis acidity of our ecocatalysts, we have used a method that implies, as a probe, pyridine adsorbed on the surface of the examined solid acid, and monitoring the bands in the range of 1,400–1,650  $\text{cm}^{-1}$  arising from its ring vibration modes (Parry 1963; Zaki et al. 2001). Infrared spectra of pyridine adsorbed on crude fraction of our ecocatalysts were recorded at 25 and 150  $^{\circ}\text{C}$  in order to distinguish frequencies of physisorbed pyridine from frequencies of pyridine coordinated to Lewis sites (Fig. 4).

The presence of weakly bonded pyridine could be attributed to a band at 1,440  $\text{cm}^{-1}$  observed at 25  $^{\circ}\text{C}$  that disappeared after outgassing at 150  $^{\circ}\text{C}$  (Fig. 4) (Parry 1963). The first indication of the extract's Lewis acidity was the presence of a band at 1,450  $\text{cm}^{-1}$  at 150  $^{\circ}\text{C}$ , characteristic for pyridine still strongly bonded at this temperature, by coordination to Lewis acid sites (Zaki et al. 2001). The decrease of the band intensity at 1,486–1,487  $\text{cm}^{-1}$  is more delicate to explain. As the

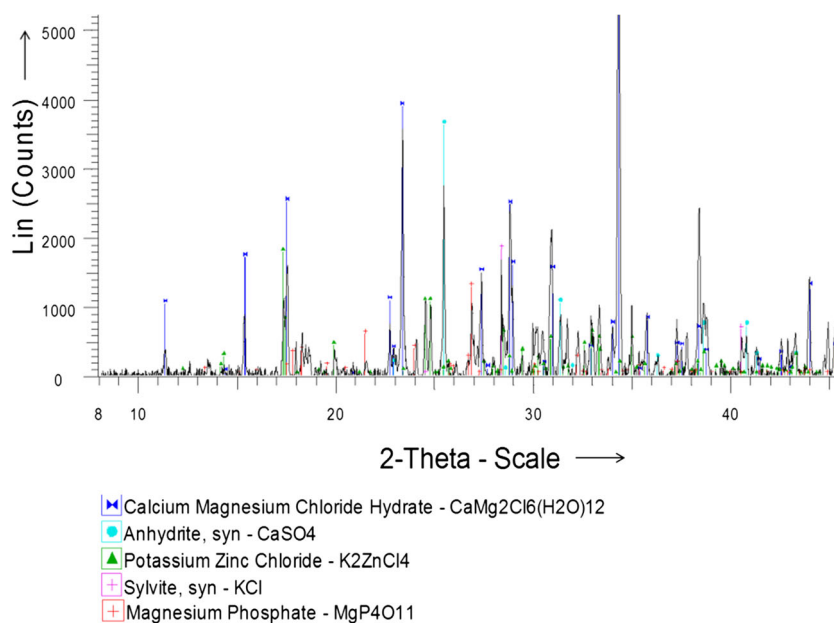
intensity was strongly reduced after heating, this band can be attributed to physisorbed pyridine. Looking at bands in the 1,590–1,640  $\text{cm}^{-1}$  region, the band at 1,599  $\text{cm}^{-1}$  observed at 25  $^{\circ}\text{C}$  disappearing at 150  $^{\circ}\text{C}$ , is characteristic for hydrogen-bonded pyridine (Parry 1963). Also, several bands were observed in the 1,600–1,640  $\text{cm}^{-1}$  range, with small variations in frequencies depending on outgassing temperature (1,608 and 1,631  $\text{cm}^{-1}$  at 25  $^{\circ}\text{C}$ ; 1,609, 1,628, and 1,639  $\text{cm}^{-1}$  at 150  $^{\circ}\text{C}$ ) (Fig. 4). The continued existence of bands in this region, in spite of heating at elevated temperature, is a proof of the presence of strongly bonded pyridine. Actually, these bands had been attributed to pyridine coordinated to Lewis acid sites, in previous studies. Additionally, their occurrence at different frequencies in this range may account for the involvement of different types of Lewis acid sites (Zaki et al. 2001) and this hypothesis was supported by the previous ICP-MS data (Tables 1 and 2), highlighting the presence of different potential Lewis acids in the extract.

**Characterization of Eco-Ni** The Eco-Ni catalyst was characterized using the same methodology (Fig. 5). The spectrum taken at 150  $^{\circ}\text{C}$  shows the persistence of an intense band at 1,447  $\text{cm}^{-1}$ , characteristic for the coordination of pyridine to Lewis acid sites. In addition, the Lewis acidity is confirmed by the presence of bands at 1,607 and 1,631  $\text{cm}^{-1}$  still intense after degassing at 150  $^{\circ}\text{C}$ . Knowing that the frequencies of the bands characteristic for the Lewis acidity increases with the strength of the Lewis acid (Parry 1963), allowed us to compare the Eco-Ni with Eco-Zn catalysts in terms of their Lewis acidity. The frequencies of the bands in the region 1,445–1,460  $\text{cm}^{-1}$  and 1,600–1,639  $\text{cm}^{-1}$ , observed in the case of the spectrum registered for Eco-Ni catalyst are always lower than those observed in the case of spectrum registered for Eco-Zn. The Lewis acidity of Eco-Ni is therefore lower than the Lewis acidity of Eco-Zn. This is consistent with the expected acidity

**Table 2** ICP-MS analysis of Eco-Ni

Ecocatalyst	Latitude and longitude data of sampling site	Mg wt% ( $\pm$ SD)	Ca wt% ( $\pm$ SD)	Fe wt% ( $\pm$ SD)	Ni wt% ( $\pm$ SD)	Mn wt% ( $\pm$ SD)	Al wt% ( $\pm$ SD)	Zn wt% ( $\pm$ SD)
Eco-Ni	-22.28417, 166.69556	7.63 ( $\pm$ 0.013)	11.98 ( $\pm$ 0.033)	0.79 ( $\pm$ 0.0050)	7.39 ( $\pm$ 0.042)	0.16 ( $\pm$ 0.00042)	0.53 ( $\pm$ 0.011)	1.25 ( $\pm$ 0.00027)

**Fig. 2** Characterization of Eco-Zn by XRD



of the main transition metal present in our ecocatalysts, since  $\text{Ni}^{2+}$  is a weaker Lewis acid than  $\text{Zn}^{2+}$  (Kobayashi et al. 2000; Satchell and Satchell 1969). However, unlike for the Eco-Zn catalyst, the infrared analysis of the adsorbed pyridine on the Eco-Ni catalyst shows the presence of a band at  $1,531\text{ cm}^{-1}$ , weak at  $25\text{ }^{\circ}\text{C}$ , but very intense after degassing at  $150\text{ }^{\circ}\text{C}$  (Fig. 5). This band has been described as characteristic for the frequency  $\nu_{19b}$  of the pyridinium ion formed by protonation of the pyridine in the presence of Brønsted acid sites (Zaki et al. 2001). The increase of its intensity after degassing at  $150\text{ }^{\circ}\text{C}$  can be connected to the influence of the heating on the kinetics of the protonation of pyridine.

The Eco-Ni catalyst comprises therefore a Lewis acid character, as well as Eco-Zn although slightly lower, and additionally a Brønsted acid character, unobserved in the case of Eco-Zn.

Catalysis of the hydro-acyloxy-addition reaction with Eco-Zn and Eco-Ni ecocatalysts

The hydro-acyloxy-addition reaction has been studied on the model example of condensation of acetic acid (**2a**) with

**Table 3** Crystalline species present in Eco-Zn catalyst and detected by XRD

Mineral	Formula (simplified)
Calcium magnesium chloride hydrate	$\text{CaMg}_2\text{Cl}_6(\text{H}_2\text{O})_{12}$
Anhydrite, syn	$\text{CaSO}_4$
Potassium zinc tetrachloride	$\text{K}_2\text{ZnCl}_4$
Sylvite, syn	$\text{KCl}$
Magnesium phosphate	$\text{MgP}_4\text{O}_{11}$

dicyclopentadiene (**1a**), in the presence of our ecocatalysts Eco-Ni or Eco-Zn (Scheme 1).

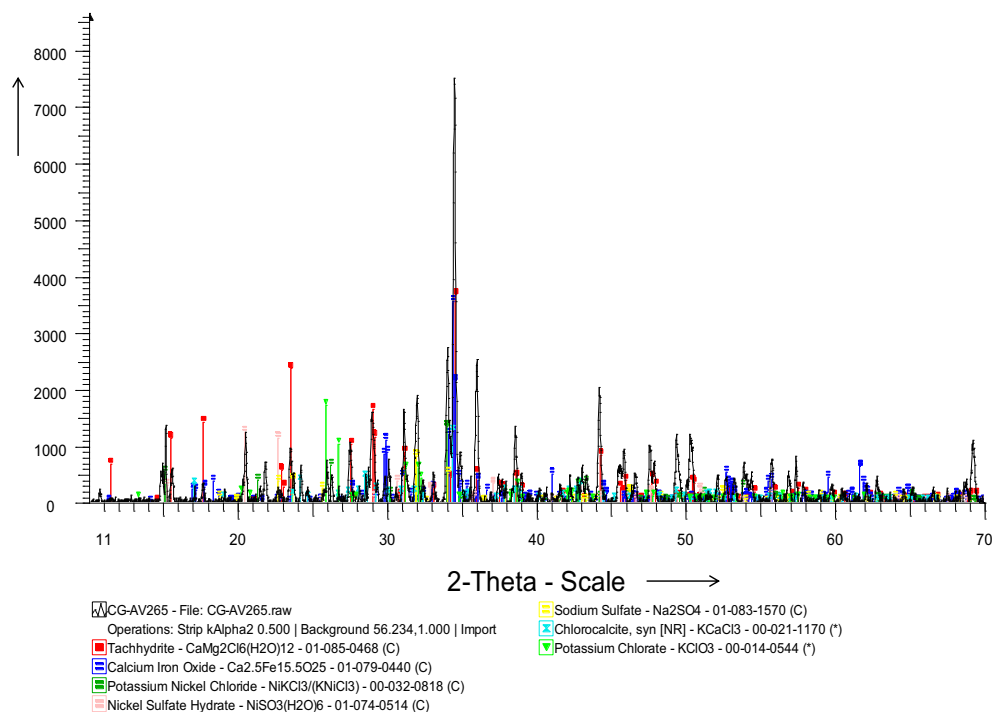
#### Reaction catalyzed by Eco-Zn catalyst

Optimization of reaction conditions was conducted by varying the amounts of acetic acid, Eco-Zn catalyst, as well as the temperature and reaction time (Table 5). At first, reaction with 10 mol% of Zn from Eco-Zn was performed during 4 h at  $120\text{ }^{\circ}\text{C}$  and gave 87 % of conversion (Table 5, entry 1). However, unexpectedly, verdyl acetate **3a** was not the major product, as it was formed only with 37 % yield. A second product was formed, with 50 % yield. After isolation by chromatography on silica gel and characterization by NMR, it was found that the major product of that reaction was the chlorinated compound **4a** (Scheme 2).

The presence of **4a** has not been reported in previous articles describing this reaction thus its formation must be specifically due to the nature of the Eco-Zn catalyst. By lowering the reaction temperature to  $60\text{ }^{\circ}\text{C}$ , the conversion reached only 6 % but **4a** became the major product (Table 5, entry 2). A comparison was made with 10 mol% of Zn using commercial  $\text{ZnCl}_2$  (Table 5, entry 3): the conversion was complete, and with a very high selectivity towards verdyl acetate **3a**, although traces of chlorinated product **4a** were also observed (around 2 % yield). This confirms that the formation of **4a** is specific to the use of Eco-Zn. Although a high yield of verdyl acetate **3a** could be obtained with the use of commercial  $\text{ZnCl}_2$ , it should be mentioned that this reagent, due to its very hygroscopic nature, is very difficult to handle, which is not observed for Eco-Zn.

Subsequently, in order to improve the conversion rate for the reaction catalyzed by Eco-Zn, heating was continued for 24 h

**Fig. 3** Characterization of Eco-Ni by XRD



(Table 5, entry 4). Under those conditions, the conversion was almost complete, and verdyl acetate **3a** was the major product isolated with 71 % yield. This example shows that the presence of **4a** in the reaction medium is only temporary since it is then obtained with only 28 % yield, whereas when reaction was heated for only 4 h, **4a** is isolated with 50 % yield (Table 5, entry 1).

Importantly, in order to avoid the use of solvent in this reaction, an excess of acetic acid can be employed (3 equivalents) (Table 5, entries 1–5). We sought to reduce the amount of acetic acid to 1 equivalent, while ensuring a good homogeneity of the reaction medium (Table 5, entries 6–10). We studied therefore the reaction in solid phase, using silica as dispersant. The reagents and Eco-Zn catalyst were co-ground in a mortar with silica gel, and the mixture thus obtained was stirred and heated (Table 5, entries 5–10). A first test reaction was performed under the same conditions as above, but with addition of silica gel to verify the influence of the heating of the solid

reaction medium (Table 5, entry 5). In this case, the complete conversion was observed and the yield of isolated verdyl acetate **3a** reached 89 %. Furthermore, the reaction time could be reduced from 24 to 12 h. This methodology proved to be effective in promoting the formation of verdyl acetate and allowed us to reduce the amount of acetic acid to 1 equivalent, while maintaining the high 84 % yield of the desired product **3a** (Table 5, entry 6). It has to mention that the reaction run in the presence of silica gel, without the Eco-Zn catalyst yielded verdyl acetate **3a** in only 17 % yield (Table 5, entry 7).

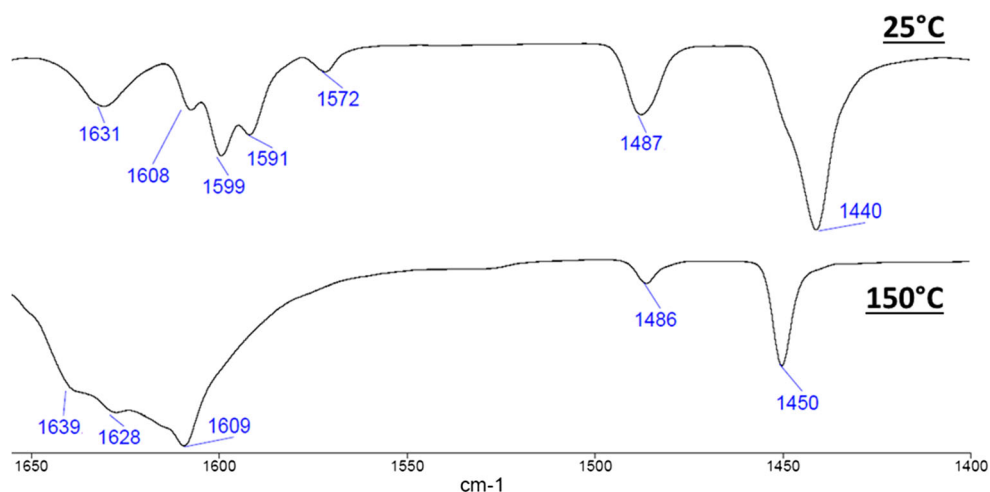
We then sought to reduce the amount of Eco-Zn catalyst, since 10 mol% can be considered as high Zn content. The reaction carried out with 1 mol% of Zn from Eco-Zn catalyst provided, on solid phase, 99 % conversion and 87 % yield of isolated verdyl acetate **3a** in just 12 h (Table 5, entry 8). This shows the effectiveness of Eco-Zn catalyst even at low loading, as the conversion increased from 17 % with the use of just silica gel (Table 5, entry 7) to 99 % with 1 mol% of Zn from Eco-Zn catalyst (Table 5, entry 8). In addition, recycling tests of the catalytic medium silica gel/Eco-Zn catalyst were performed (Table 5, entries 9 and 10). Washing with cyclohexane, filtration and drying in an oven (120 °C, 5 h) of the solid phase at the end of reaction provide the recycled catalytic system. Other solvents in which the metal chlorides present in the Eco-Zn catalyst are not soluble can also be used instead of cyclohexane. For example, ethyl acetate proved equally effective to wash the solid phase. After recycling the solid phase containing Eco-Zn, we used it again in the reaction under the same conditions as describe above, and only slight decrease in the conversion was detected (Table 5, entries 9 and 10).

**Table 4** Crystalline species present in Eco-Ni catalyst and detected by XRD

Mineral	Formula (simplified)
Calcium magnesium chloride hydrate	$\text{CaMg}_2\text{Cl}_6(\text{H}_2\text{O})_{12}$
Calcium iron oxide	$\text{Ca}_{2.5}\text{Fe}_{15.5}\text{O}_{25}$
Potassium nickel chloride	$\text{KNiCl}_3$
Nickel sulfate hydrate	$\text{NiSO}_4(\text{H}_2\text{O})_6$
Sodium sulfate	$\text{Na}_2\text{SO}_4$
Chlorocalcite, syn	$\text{KCaCl}_3$
Potassium chlorate	$\text{KClO}_3$



**Fig. 4** IR spectra of pyridine adsorbed on Eco-Zn, after brief outgassing at the indicated temperatures



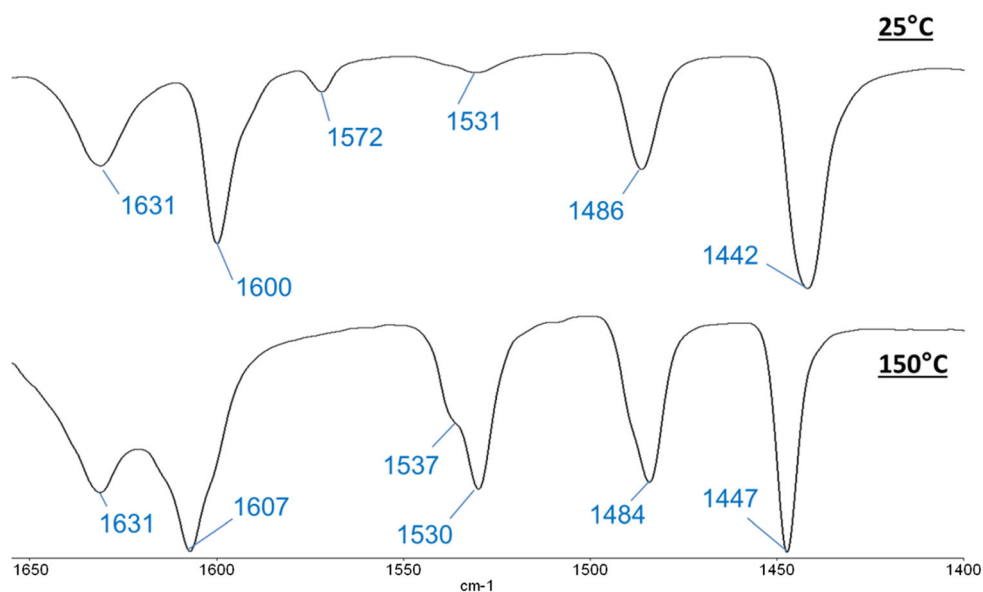
#### Reaction catalyzed by Eco-Ni catalyst

The IR studies described earlier revealed a significant difference between the Eco-Zn and Eco-Ni catalytic systems. Eco-Ni catalyst shows a lower Lewis acidity than Eco-Zn; however, it demonstrates a strong Brønsted acid character. It seemed therefore interesting to compare the activity of both catalytic systems in the hydro-acyloxy-addition reaction of acetic acid (**2a**) with dicyclopentadiene (**1a**).

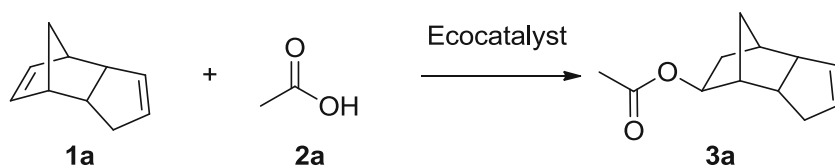
The Eco-Ni catalyst was first tested in the reaction under similar conditions as it was for Eco-Zn: loading 10 mol%, 4 h reaction time at 120 °C. Under these conditions, the conversion observed was 58 % (Table 6, entry 1), which is less than the conversion observed previously with Eco-Zn (87 %, Table 5, entry 1). This result can be explained by the lower Lewis acidity of Eco-Ni compared to Eco-Zn catalyst. It is

therefore in agreement with the IR study of the two cocatalysts. However, the proportion of chlorinated product **4a** versus verdyl acetate **3a** was higher with the use of Eco-Ni (**4a/3a**: 44/14) than with the Eco-Zn (**4a/3a**: 50/37). This result can also be connected with the data obtained during infrared characterization of the Lewis acidity by pyridine adsorption (“Infrared characterization of the Lewis acidity of Eco-Zn and Eco-Ni by pyridine adsorption” section). Unlike Eco-Zn, the Eco-Ni catalyst showed a significant Brønsted acid character, probably due to the presence of HCl linked to this catalyst. A clear evidence supporting this assumption is the tendency of Eco-Ni to promote hydrochlorination of alkenes. Thus with Eco-Ni, having a lower Lewis acidity than Eco-Zn catalyst, the formation of verdyl acetate **3a** remains a secondary reaction. The structure of the Eco-Ni is therefore ideal to promote the formation of the chlorinated product **4a**.

**Fig. 5** IR spectra of pyridine adsorbed on Eco-Ni, after brief outgassing at the indicated temperatures



**Scheme 1** Hydro-acyloxy-addition reaction catalyzed by the ecocatalysts



Due to the selectivity observed with Eco-Ni, we looked at whether it is possible to obtain more selectively the chlorinated product **4a**. Unlike the reaction where verdyl acetate **3a** is formed as the major product, the synthesis of chlorinated compound **4a** requires the use of Eco-Ni not only as the catalyst but also as a reagent, a source of Cl. Therefore, we increased the amount of Eco-Ni, using 1 equivalent of Ni from Eco-Ni. We also reduced the reaction time to 2 h, since it had been observed previously that the product **4a** is converted to **3a** with prolonged reaction time (Table 5, entries 1 and 4). Under these conditions (1.0 equiv Eco-Ni, 120 °C, 2 h), the conversion reached 100 %, but the selectivity towards the chlorinated product **4a** was very low (Table 6, entry 2). In turn, using 0.5 equiv of Ni from Eco-Ni, allowed us to conserve the high conversion and to improve the selectivity towards the desired chlorinated product **4a** (64 % yield, Table 6, entry 3). Thus, it can be concluded that the poor selectivity observed when using 1.0 equivalent of Ni originated from the large amount of Ni present, which led to the formation of undesired verdyl acetate **3a** at the cost of **4a**.

Further optimization of the reaction with 0.5 equiv of Ni led to the isolation of desired chlorinated product **4a** with 68 % yield when the reaction time was 1 h (Table 6, entries 3–5). So far the reactions were conducted with the use of 3 equiv of acetic acid, in order to maintain comparable conditions to those applied for Eco-Zn. However, since the application of Eco-Ni focused on the formation of chlorinated product **4a**, acetic acid could probably be superfluous. We therefore carried out an experiment without the acetic acid and under optimal conditions for the formation of product **4a** (Ni 0.5 equiv, 120 °C, 2 h). Surprisingly, the desired product **4a** was isolated only with 29 % yield (Table 6, entry 6). It therefore appears that the presence of acetic acid is required in the reaction with Eco-Ni if chlorinated product **4a** is to be obtained with good yield. It is possible that acetic acid plays a role in the process of releasing of HCl from Eco-Ni, e.g., by stabilizing the metal ions resulting from departure of Cl, to which chlorine was attached in the Eco-Ni before reaction. However, the amount of acetic acid could be reduced to 1 equivalent maintaining the good yield of the desired chlorinated product **4a** (Table 6,

**Table 5** Hydro-acyloxy-addition reaction catalyzed by Eco-Zn catalyst: optimization of the reaction conditions

Entry <sup>a</sup>	Equiv. of AcOH	Solid medium	T (°C)	mol% of Zn from Eco-Zn	Time (h)	Conversion (%) <sup>b</sup>	Yield (%) <sup>b</sup>		
1	3	–	120	10	4	87	37	<b>3a</b>	<b>4a</b>
2	3	–	60	10	4	6	1		
3	3	–	120	10 <sup>c</sup>	4	100	98		
4	3	–	120	10	24	99	71		
5	3	SiO <sub>2</sub> <sup>d</sup>	120	10	12	100	89		
6	1	SiO <sub>2</sub> <sup>d</sup>	120	10	12	100	84		
7	1	SiO <sub>2</sub> <sup>d</sup>	120	0	12	17	17		
8	1	SiO <sub>2</sub> <sup>d</sup>	120	1	12	99	87		
9	1	SiO <sub>2</sub> <sup>d</sup>	120	1 <sup>e</sup>	12	85	72		
10	1	SiO <sub>2</sub> <sup>d</sup>	120	1 <sup>f</sup>	12	73	62		

<sup>a</sup> Reaction performed on 1.0 mmol of dicyclopentadiene

<sup>b</sup> Conversion and yield were determined by GC-MS with biphenyl as internal standard

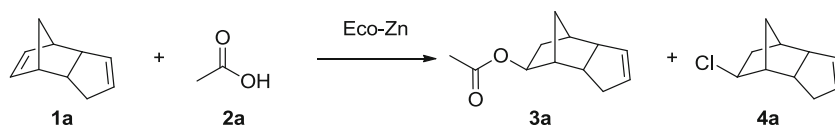
<sup>c</sup> Commercial ZnCl<sub>2</sub> was used as the source of Zn

<sup>d</sup> A 1.0 g of SiO<sub>2</sub> was used as solid medium

<sup>e</sup> Catalyst recycled one time

<sup>f</sup> Catalyst recycled two times

**Scheme 2** Products obtained with the Eco-Zn catalyst



entry 7) and even better ratio of products **4a/3a** (**4a/3a**: 70/10, Table 6, entry 7) than that obtained with the use of 3 equiv of acetic acid (**4a/3a**: 68/23, Table 5, entry 4) (which is a consequence of the use of an excess of acetic acid, favoring the formation of the acetate **3a**).

Additionally, we have tested this reaction under solvent-free conditions with addition of silica gel as a dispersing agent (in order to provide a better homogenization of the reaction medium). Under those conditions, the chlorinated product **4a** was obtained with 69 % yield (Table 5, entry 8), which is comparable to the results obtained without the addition of silica gel (Table 5, entry 7). Ratio of products **4a/3a** however was slightly lower (**4a/3a**: 69/29). Extending the reaction time to 5 h did not improve those results (Table 5, entry 9). Therefore, the use of a solid dispersant is not as advantageous in the case of the formation of chlorinated product **4a** with Eco-Ni, as it was in the case of formation of verdyl acetate **3a** with Eco-Zn (see Table 5).

Finally, a reaction carried out under the optimized conditions (Ni 0.5 equiv, 120 °C, 1 h), but with a commercial nickel chloride (Table 6, entry 10) led to a lower yield of **4a** than that observed with the use of Eco-Ni (45 % yield with the use of NiCl<sub>2</sub>·6H<sub>2</sub>O versus 69 % yield with the use of Eco-Ni; Table 6, entry 8). These observations show that the results

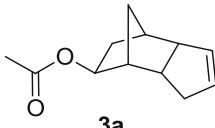
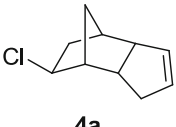
obtained with Eco-Ni depend directly on the nature of the plant-based extract, whose constituents have unique and specific properties in organic synthesis. The Eco-Ni catalyst thus provides a synthetic pathway for green and efficient synthesis of compound **4a**, which may serve as a precursor in organic synthesis, especially in coupling reactions that frequently involve the use of halogenated compounds.

#### Comparison of the catalytic activity of Eco-Zn with Eco-Ni

Having in our hand the optimized conditions for the hydro-acyloxy-addition reaction catalyzed by Eco-Zn and Eco-Ni, we have applied this protocol to different substrates, in order to examine the reaction scope and limitations (Table 7). As depicted on Fig. 6 and in Table 7, reactions with Eco-Zn allowed selective preparation of esters **3a–h** whereas the use of Eco-Ni yielded selectively the chlorinated products **4a–c**.

Reaction between cyclohexane **1b** and acetic acid **2a** is a good illustration of the tendency observed during the preliminary optimization of the reaction conditions with the use of dicyclopentadiene as substrate. When optimized conditions favoring the formation of acetate with Eco-Zn were used (1 mol%, 120 °C, 24 h), the desired acetate **3b** was obtained with 71 % yield (Table 7, entry 1). In turn, application of the

**Table 6** Hydro-acyloxy-addition reaction catalyzed by Eco-Ni catalyst: optimization of the reaction conditions

Entry <sup>a</sup>	Equiv. of AcOH	Solid medium	T (°C)	Equiv of Ni from Eco-Ni	Time (h)	Conversion (%) <sup>b</sup>	Yield (%) <sup>b</sup>		
									
1	3	–	120	0.1	4	58	14	<b>3a</b>	<b>4a</b>
2	3	–	120	1.0	2	100	52		
3	3	–	120	0.5	2	99	35		
4	3	–	120	0.5	1	91	23		
5	3	–	120	0.5	0.5	76	16		
6	0	–	120	0.5	2	29	0		
7	1	–	120	0.5	1	80	10		
8	1	SiO <sub>2</sub> <sup>c</sup>	120	0.5	1	98	29		
9	1	SiO <sub>2</sub> <sup>c</sup>	120	0.5	5	100	36		
10	1	SiO <sub>2</sub> <sup>c</sup>	120	0.5 <sup>d</sup>	1	73	28		

<sup>a</sup> Reaction performed on 1.0 mmol of dicyclopentadiene

<sup>b</sup> Conversion and yield were determined by GC-MS with biphenyl as internal standard

<sup>c</sup> A 1.0 g of SiO<sub>2</sub> was used as solid medium

<sup>d</sup> Commercial NiCl<sub>2</sub>·6H<sub>2</sub>O was used as source of Ni

**Table 7** Eco-Zn- and Eco-Ni-catalyzed hydro-acyloxy-addition reaction with the use of various substrates

Entry <sup>a</sup>	Alkene	Carboxylic acid	Eco-M	Time (h)	Conversion (%) <sup>b</sup>	Products	Yield (%) <sup>b</sup>	Yield (%) <sup>b</sup>	
1			Eco-Zn	24	100		71		29
2			Eco-Ni	1	100		12		88
3			Eco-Zn	24	100		95		5
4			Eco-Ni	1	100		20		80
5			Eco-Zn	24	53		53		0
6			Eco-Zn	24	100		74		26
7			Eco-Zn	24	89		58		31
8			Eco-Zn	24	100		93		7
9			Eco-Zn	24	100		92		8

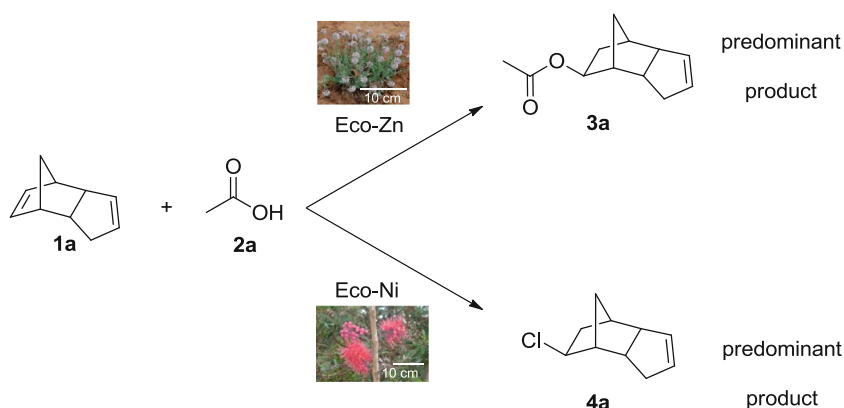
<sup>a</sup> Reaction conditions : alkene (1.0 mmol), carboxylic acid (1.0 mmol), Eco-Zn (10 mg, amount corresponding to 0.01 mmol of Zn, according to ICP-MS analysis) or Eco-Ni (entries 2 and 4) (385 mg, amount corresponding to 0.5 mmol of Ni, according to ICP-MS analysis), and silica gel (1.0 g), 120 °C

<sup>b</sup> Conversion and yield were determined by GC-MS with biphenyl as internal standard

optimized conditions for the formation of chlorinated product with Eco-Ni (0.5 equiv, 120 °C, 1 h) produced the desired product **4b** with 88 % yield (Table 7, entry 2). Similarly, reaction with the use of norbornene **1c**, as alkene, furnished selectively the acetate **3c** (95 % yield, Table 7, entry 3) with the use of Eco-Zn and the chlorinated product **4c** (80 % yield,

Table 7, entry 4) with the use of Eco-Ni catalyst. Subsequently, various carboxylic acids were tested in order to check the possibility to produce different esters. Reaction with formic acid **2b** and dicyclopentadiene **1a** produced the desired ester **3b** with 53 % yield. Formation of the corresponding chlorinated product **4a** was not observed (Table 7, entry

**Fig. 6** Selective formation of product **3a** or **4a** depending on the nature of the ecocatalysts used



5). The use of propanoic acid **2c** and butanoic acid **2d** and dicyclopentadiene **1a** allowed the preparation of corresponding esters **3e** (74 % yield, Table 7, entry 6) and **3f** (58 % yield, Table 7, entry 7). The ester **3e** is a compound of commercial interest for the perfume industry, and is sold under the commercial name Florocyclene<sup>®</sup> and Cyclaprop<sup>®</sup> (Fig. 1). It should be noted that the yield of ester decreases when increasing the length of the carbon chain in the carboxylic acid used, from acetic to butanoic acid. In turn, very good yields of the reaction of propanoic acid with norbornene (93 %, Table 7, entry 8) and butanoic acid with norbornene (92 %, Table 7, entry 9) were obtained in the presence of Eco-Zn catalyst. In terms of selectivity, it should be noted that the formation of esters and chlorinated products from norbornene and dicyclopentadiene proceeded with >95 % exo selectivity, as determined by <sup>1</sup>H NMR.

## Conclusion

In conclusion, we showed that two different biosourced polymetallic ecocatalysts Eco-Zn and Eco-Ni prepared from biomass derived from metal-hyperaccumulating plants *N. caerulescens* (Brassicaceae), *A. vulneraria* (Fabaceae), and *G. pruinosa* (Cunoniaceae) can efficiently and selectively mediate two different addition reactions on alkenes, leading to commercially valuable fine chemicals. The developed synthetic protocols based on the use of Eco-Zn and Eco-Ni are characterized by several advantages: low catalyst loading and high yields of reaction products, short reaction times, reaction without solvent or in solid state, broad substrate scope, and the possibility of the catalyst recovery and reuse. Eco-Zn thus provides an efficient way of access to various esters following the hydro-acyloxy-addition reaction of carboxylic acids with alkenes (53–95 % yield). In contrast, starting from the same substrates, Eco-Ni shows a different reactivity and affords chlorinated products by hydrochlorination of the starting alkenes (70–88 % yield). These results can be connected to the

original compositions and properties of the ecocatalysts, whose reactivity thus correlates with the botanical origin of the metal-hyperaccumulating plant. Phytoextraction proves therefore to be useful not only for the environment but also for organic chemistry. This constitutes an encouragement to develop phytoextraction for the recovery of degraded sites, by giving a new scientific and economic value to the resulting biomass.

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