Biosourced polymetallic catalysis: a surprising and efficient means to promote the Knoevenagel condensation

Pierre-Alexandre Deyris, Valérie Bert, Sébastien Diliberto, Clotilde Boulanger, Eddy Petit, Yves-Marie Legrand, Claude Grison

To cite this version:

Pierre-Alexandre Deyris, Valérie Bert, Sébastien Diliberto, Clotilde Boulanger, Eddy Petit, et al.. Biosourced polymetallic catalysis: a surprising and efficient means to promote the Knoevenagel condensation. Frontiers in Chemistry, Frontiers Media, 2018, 6, pp.art. 48. 10.3389/fchem.2018.00048. ineris-01863338

HAL Id: ineris-01863338
https://hal-ineris.archives-ouvertes.fr/ineris-01863338
Submitted on 28 Aug 2018

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.
Biosourced Polymetallic Catalysis: A Surprising and Efficient Means to Promote the Knoevenagel Condensation

Pierre-Alexandre Deyris 1, Valérie Bert 2, Sébastien Diliberto 3, Clotilde Boulanger 3, Eddy Petit 4, Yves-Marie Legrand 4 and Claude Grison 1*

1 Laboratoire de Chimie Bio-Inspirée et D’Innovations Ecologiques, UMR 5021 Centre National de la Recherche Scientifique – Université de Montpellier, Grabels, France, 2 INERIS, Clean and Sustainable Technologies and Processes Unit, DRC/RISK, Parc Technologique Alata, BP2, Verneuil-en-Halatte, France, 3 Institut Jean Lamour, UMR 7198, Université de Lorraine, Centre National de la Recherche Scientifique, Metz, France, 4 IEM, Université de Montpellier, Centre National de la Recherche Scientifique, ENSCM, Montpellier, France

Zn hyperaccumulator (Arabidopsis halleri) and Zn accumulator Salix “Tordis” (Salix schwerinii × Salix viminalis) have shown their interest in the phytoextraction of polluted brownfields. Herein, we explore a novel methodology based on the chemical valorization of Zn-rich biomass produced by these metallophyte plants. The approach is based on the use of polymetallic salts derived from plants as bio-based catalysts in organic chemistry. The formed ecocatalysts were characterized via ICP-MS, X-ray diffraction (XRD), Fourier transform infrared spectroscopy (FT-IR) in order to precise the chemical composition, structure, and behavior of the formed materials. The Doebner-Knoevenagel reaction was chosen as model reaction to study their synthetic potential. Significant differences to usual catalysts such as zinc (II) chloride are observed. They can principally be related to a mixture of unusual mineral species. DFT calculations were carried out on these salts in the context of the Gutmann theory. They allow the rationalization of experimental results. Finally, these new bio-based polymetallic catalysts illustrated the interest of this concept for green and sustainable catalysis.

Keywords: sustainable chemistry, ecocatalysis, bio-sourced catalyst, Knoevenagel condensation, phytoextraction

INTRODUCTION

Over the last decade, there has been an awareness of the elemental sustainability (Hunt et al., 2015). The concerns over the depletion of the traditional sources of elements are actually rising. At the current rate, the known reserves of numerous element such as Zn will be consumed in <50 years (Hunt, 2013). However, elements are not lost but dispersed throughout our environment by human activities. For example, several wastes including municipal solid waste, e-waste, metallurgic, and mining wastes contain elevated levels of metals (Cui and Zhang, 2008; Jung and Osako, 2009). This dispersion is responsible for contamination and therefore degradation of soils. Moreover, this contamination can enter human bodies via inhalation, skin contact, or food chain and may have health impacts. Because of the elemental sustainability and the social issues, the recovery of metals from our waste can be considered as a hazard avoidance and an alternative to primary resources.
In this context, phytoremediation of industrial contaminated soils, a worldwide environmental issue, is a promising method for restoring soils and recovering metal. Certain plants can extract and accumulate metals from the soil into the biomass (van der Ent et al., 2013). This capacity called phytoextraction offers the opportunity to remediate soils (Losfeld et al., 2015). While the metal-enriched biomass might be considered as a hazardous waste, various valorization ways, exploiting recovered metal are currently investigated. Recently, we have studied the development of a series of novel approaches for the recycling and reuse of vital minerals for industrial chemistry, linked with the phytoremediation of contaminated soils on mining sites. Taking advantage of the remarkable ability of certain land plants to hyperaccumulate transition metals into shoots, we have addressed the use of metals derived from contaminated plant waste as Lewis acids, coupling catalysts, oxidizing, and reducing agents in organic synthesis of biomolecules. This was the first chemical recovery of new phytoextraction technologies. Recycling of waste from metal-enriched plants has led to the new concept in chemistry, namely “ecocatalysis” (Grison, 2015). The first results revealed that ecocatalysts (called Eco-M) can be more efficient and selective than traditional catalysts. For example, Eco-Zn were found to be very efficient in the Garcia Gonzalez reaction (Escande et al., 2014c) and showed unusual flexibility in catalyzed preparation of polyhydroxyalkyl furans. The polymetallic systems arising from the biomass produced by phytoextraction are unusual and lead to new associations of chemical species, such as CaMg₂Cl₆·12H₂O, Mg₃PO₄·11H₂O, or K₂ZnCl₄ (Escande et al., 2015a). They exhibit also a better chemo- and regio-selectivity than traditional catalysts in numerous reactions [multicomponent (Grison et al., 2013), domino (Escande et al., 2015d), cycloaddition (Escande et al., 2014b), catalytic reduction (Escande et al., 2015c, 2017b), and oxidation (Escande et al., 2015b, 2017a)]. They can be used as catalysts in synthetic transformations of biomass, leading to molecules of interest to industrial chemicals (peratures and cosmetics), oligomers of biological interest (Thillier et al., 2013), functionalized aromatic heterocyclic compounds (Clavé et al., 2016, 2017), and building blocks of industrial chemical processes.

In this paper, we wish to carry out a comparative study of new biomass, which derived from two metal-contaminated areas. Both sites are located in the Hauts-de-France region which was heavily impacted by metallurgical activities emissions. As a result, soils and canal sediments were contaminated by metals such as Zn, Cd, Pb, and Cu. For some metal polluted sites, phytomanagement seems to be a relevant option to handle the risks associated to the pollutants and valorize the produced biomass by ecocatalysis (Kidd et al., 2015). Herein, we present two different case studies: a Zn hyperaccumulator Arabidopsis halleri (Huguet et al., 2012) and a Zn accumulator Salix “Tordis” (Salix schwerinii × Salix viminalis). A. halleri was cultivated on a Zn-contaminated soil for phytoextraction purpose whereas S. “Tordis” was planted on a metal contaminated sediment landfill site for bioenergy production. Because both plant species accumulate Zn in a very important amount in their harvestable parts (i.e., leaves), this ability was suitable to ecocatalysis. According to the known optimal zinc accumulation capacity of each plant, the principal objective is to determine if the utilization of these plants is valuable in catalysis. To that end, the catalytic potential of each ecocatalyst was tested by the Knoevenagel condensation as model reaction.

The Knoevenagel condensation is widely used in organic synthesis especially in pharmaceutical industry. For example, it is present in one-step on the synthesis of atorvastatin, pioglitazone, pregabalin, rosuvastatin, or rosiglitazone (Harrington, 2011; Debarge et al., 2014; Vardanyan and Hruby, 2016). The catalysts commonly used are weak organic bases such as pyrrolidine or ammonium salts such as piperidine/acetic acid. Several Lewis acids such as ZnCl₂, CeCl₃·7H₂O·NaI, NbCl₅, and AlCl₃ have been reported (Shanthan Rao and Venkataraman, 1991; Bartoli et al., 2006; Yadav et al., 2009; Li et al., 2012).

MATERIALS AND METHODS
Phytomanagement on Metal-Contaminated Brownfield Area
Biomasses Cultivation and Collection
The willow cultivar “Tordis” was planted in 2012 in a very short rotation coppice (VSRC) on metal-polluted dredged sediments for the purpose of bioenergy production. The site is located at Fresnes-sur-Escuat (Hauts-de-France, France). The field site was devoted to metal immobilization by using a soil amendment and a soil grass cover and simultaneously to the production of woody biomass by growing willows. The 650 m² site which contains 350 willows produced ~1.4 tons of dried biomass in optimal conditions. As the willow “Tordis” leaves contained a large amount of Zn, the collection of leaves in November 2015 was undertaken to avoid metal pollution dispersion. At the same time, leaves of the Zn-hyperaccumulator A. halleri were collected in a site impacted by a Zn smelter (Auby, Hauts-de-France, France) in December 2015. On this site, numerous A. halleri plants are present, as a result of its spread on the entire site from the beginning of the nineteenth century, which allowed the collection of the desired amount of leaves to be processed. A. halleri from this site was extensively studied (Gomez-Baldaras et al., 2014). Like willows, the management of the Zn-enriched biomass produced on this site was undertaken by exploring the possibility to valorize the plant Zn content.

Characterization of Biomasses
After homogenization of the collected leaves, a composite sample was formed and split in 10 and five samples for S. Tordis and A. halleri, respectively. Leaves were oven-dried at 40°C until constant weight. 0.5 g were ground and digested at 180°C for 20 min in 10 mL of nitric acid (67%) and 3 mL of ultra-pure water using a microwave digester (Mars Xpress, CEM). The digested samples were filtered to 0.45 µm (Millipore) and acidified to a pH < 2 for preservation. The analyses of metal concentrations were performed by ICP-AES (Agilent 720ES). Two standard references of plant material (white cabbage “BCR-679,” Community Bureau of Reference and tobacco leaves “CTA-VTL-2”) were included for analytical quality control.
Catalyst Preparation and Characterization
Preparation of the Ecocatalysts
Biomasses (usually 100 g) were first dried into an oven at 80°C and then thermally treated in an oven under air flow. The temperature program was a first gradient from 20 to 350°C in 1 h then 2 h at 350°C and a second one from 350 to 550°C in 1 h then 2 h at 550°C before a slow cooling overnight in the oven to obtain ashes. The resulting powder was subjected to a chemical treatment. In the typical procedure, the ashes (10 g) were added to 100 mL of 6 M HCl solution then heated and stirred at 60°C for 3 h. The reaction mixture was filtered through celite. The resulting yellow solution was refluxed during 5 h and then was evaporated at 80°C. The dry residues were crushed to fine powder in a mortar and placed in an oven at 90°C for 2 days. Purification steps are not mandatory in our process.

Preparation of the Double Salts
The preparation of the double salts was adapted from previously published work (Poddar et al., 2015). KCl and MgCl₂ or ZnCl₂ were dissolved separately in analytical grade water in the desired ratio. The resulting solutions were combined and then evaporated at 80°C. The dry residues were crushed to fine powder in a mortar and placed in an oven at 90°C for 2 days.

Characterization of the Catalysts
MP-AES analysis (microwave plasma-atomic emission spectrometer)
The samples were digested in 10 mL of reversed aqua regia [1:2 hydrochloric acid (37%):nitric acid (65%)] under a microwave-assisted digestion (Milestone ETHOS Tauch) with the following program: 20–90°C in 7 min, 90–170°C in 5 min, 170–210°C in 3 min, and then 20 min isothermal at 210°C. Samples were filtered and then diluted to 0.1 mg/L in nitric acid 2.5%.

X-ray diffraction analysis
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 Cu Ka radiation λ = 1.54086 Å) equipped with a Lynxeyes detector.

Pyridine-FT-IR (fourier transform infrared spectroscopy)
FT-IR measurements were carried out with pyridine probe. A PerkinElmer Spectrum 100 FT-IR spectrometer was used for recording the spectra. 10 ± 1 mg of catalyst were introduced in a 10 mL vial under argon. Dichloromethane (2 mL) was added followed by pyridine (100 μL) and the resulting mixture was stirred by a vortex during 1 min. Surplus pyridine was adsorbed, and the samples were degassed for 15 min at 25°C under an argon flux. A sample of the resulting powder (5 ± 1 mg) was picked up in order to record a first spectrum. The samples were then degassed for 25 min at 150°C (10⁻³ Pa) to eliminate the physisorbed pyridine. A second spectrum was then recorded with 5 ± 1 mg of the resulting powder.

General Procedure for Knoevenagel Condensations
Representative procedure
The catalyst was dried prior to use and was added to an argon flushed 5 mL round bottom flask followed by biphenyl, aromatic aldehyde (5 mmol), and methylene nucleophile (5 mmol). For control experiments, no catalyst was added. After stirring 1 h at 100°C under argon atmosphere, the reaction was cooled under running water and then diluted with ethyl acetate. The conversion of benzaldehyde was estimated by GC-MS analysis using biphenyl as an internal standard. To check the GC-MS calibration, some products were purified by flash-chromatography using ethyl acetate/cyclohexane as eluent.

Analysis of reaction products
NMR spectra were recorded on a Brüker Avance 300 spectrometer at room temperature. ¹H frequency is at 300 MHz, ¹³C 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 m x 0.25 mm x 0.25 μm ZB-5MSi Guardian column (Phenomenex®) with hydrogen as carrier gas.

DFT analysis
Density functional theory (DFT) calculations were performed using the Gaussian 09 computational package (Frisch et al., 2009). The B3LYP functionals was used as implemented in Gaussian. Geometry and electronic structure optimizations were carried out for the functional/basis set combinations using the default optimization algorithm in Gaussian 09. Energy and frequency calculations were performed on fully optimized geometries using the 6-311++G(2d,p) basis set. B3LYP functionals showed their relevance for studying metals (Lai et al., 2012) and it is widely used. Two different models have been applied to determine partial atomic charge: Mulliken (Mulliken, 1955), and Merz–Kollman (MK) (Besler et al., 1990).

RESULTS AND DISCUSSION
Phytoremediation
In this study, we used a Zn hyperaccumulator plant, A. halleri, and a Zn accumulator plant, Salix “Tordis.” Both are included in a large program of phytomanagement of metal-contaminated brownfields located in the north of France (Figure 1). After harvesting and drying the leaves of each plant, ICP-AES analyses were realized in order to measure element proportions in the biomass (Table 1). In addition to the normal of physiological species, both plants are able to accumulate Zn, and to a lesser extent Cd. As expected, A. halleri hyperaccumulates Zn, that represents more than 1% of its dry weight. The willow confirmed to have a Zn-accumulating behavior as its leaves contained Zn in concentration far above the baseline one. Based on Zn foliar concentrations A. halleri and willow leaves are suitable for the Zn-ecocatalyst production.

Preparation and Characterization of the Ecocatalysts
The ecocatalysts were prepared from an important quantity of biomass to homogenize their composition and minimize their variation. The biomass was calcined under air flow to remove most of organic matter (Garel et al., 2015). The mass loss for A. halleri and Salix was around 80–90%. The residues were treated with hydrochloric acid in order to form...
active metal chloride salts. According to MP-AES analysis, both
eccocatalysts contained zinc in good proportions, accompanied
by the physiological metals such as calcium, potassium, and
magnesium (Table 2). As expected, the quantity of zinc was
higher in the ecocatalyst made from A. halleri (Eco-A.h.), a
Zn hyperaccumulator (7.6%), than in those derived from Salix
“Tordis” (Eco-S.T.), a Zn accumulator (2.6%). In comparison
with our previous works related to Eco-Zn catalysts, which
derived from phytoextraction on mining sites (Grison et al.,
2015), the zinc concentration in Eco-A.h. is approximately
the same than the one found in the catalyst stemmed from Noccaea
cauulescens. However, Anthyllis vulneraria gave a catalyst with
the highest Zn content, better than the two brassicaceae, A. halleri
and N. cauulescens (Table 2). It should be noted that large
differences were observed for calcium: the Ca concentration is
significantly higher in Eco-S.T. than in Eco-A.h. (respectively,
17.1 and 9.7%).

The two biosourced catalysts were analyzed by X-ray
diffraction (XRD). The XRD characterization revealed the
presence of an unusual mixture of metal chlorides such as K₂ZnCl₄ (potassium tetrachlorozincate), KMgCl₃·6H₂O (carnallite), KCl (sylvite), and a remaining sulfate,
CaSO₄·(H₂O)₀.₅, in the crystalline fraction in both ecocatalysts
(Table 3). K₂ZnCl₄, a masked form of ZnCl₂, was observed in
Eco-A.h. but not in Eco-S.T. For the latter, zinc was not prese nt
in the crystalline phase or at a non-detectable level. Compar ed
to ecocatalysts derived from N. cauulescens and A. vulneraria,
the lack of CaMg₂Cl₆·12H₂O was as surprising as the presence of
carnallite (Escande et al., 2014a). Although XRD analyses only
testify the presence of crystalline species, eco-catalysts showed a
very original composition (see Supplementary Material).

The Lewis acidity of our ecocatalysts has been studied with
an empirical method that implies the coordination of adsorbed
pyridine on the surface of acidic solids. By studying the bands

---

**TABLE 1 | ICP-MS Analysis of leaves derived from Arabidopsis halleri and Salix “Tordis” (ppm).**

| Plant           | As   | Ca    | Cd    | Cr    | K     | Mg    | Mn    | Ni    | P     | Pb    | S     | Zn    |
|-----------------|------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| Arabidopsis halleri | 3.05 | 18,060| 141   | 1.92  | 21,180| 3,624 | 61    | 7.64  | 3,150 | 118   | 7,570 | 12,960|
|                 | (1.28)| (1,266)| (40)  | (1.05)| (3,235)| (380) | (20)  | (2.66)| (443) | (98)  | (698) | (2,449)|
| Salix “Tordis”  | 0.28 | 25,840| 9.57  | 4.33  | 9,140 | 1,780 | 63    | 1.83  | 1,739 | 3.03  | 5,690 | 3,159 |
|                 | (0.05)| (1,857)| (0.84)| (1.06)| (659) | (94)  | (5)   | (0.14)| (99)  | (0.74)| (407) | (271) |

Values are means ± SE (in brackets).
in the range of 1,640–1,400 cm\(^{-1}\), Lewis and Bronsted acidities can be evaluated for each catalyst and compared with classical metal chlorides (Parry, 1963; Zaki et al., 2001). Infrared spectra of pyridine adsorbed on crude fraction of ecocatalysts were recorded at 25°C and after heating at 150°C in order to distinguish the frequencies of physisorbed pyridine from those of pyridine coordinated to Lewis sites (Figure 2). After outgassing at 25°C, IR studies did not show a huge difference between both of the ecocatalysts. Bands at 1,445 cm\(^{-1}\) and at 1,443 cm\(^{-1}\) for Eco-A.h. and Eco-S.T, respectively, denoted the presence of weakly bonded pyridine and thus Lewis acid sites as well as bands at 1,607 and 1,608 cm\(^{-1}\), which are characteristic of pyridine strongly bonded to Lewis acid sites, observed after outgassing at 25°C. The intensity of this band was therefore persistent after outgassing at 150°C. No Bronsted acidic character was observed. This observation was consistent with this salt’s aspect, which is not deliquescent after contact with air compared to ZnCl\(_2\) and makes it easier to handle. K\(_2\)ZnCl\(_4\) was characterized with the same methodology. Figure 2 shows that the band at 1,400 cm\(^{-1}\) observed at 25°C disappeared after outgassing at 150°C. The presence of intense bands at 1,440–1,480 and 1,617 cm\(^{-1}\) at 150°C indicated the presence of strongly bonded pyridine. Frequencies of these bands are higher than those observed for K\(_2\)ZnCl\(_4\) that makes K\(_2\)MgCl\(_2\)-6H\(_2\)O a better Lewis acidity than K\(_2\)ZnCl\(_4\). The catalytic potential of K\(_2\)MgCl\(_2\)-6H\(_2\)O is little known. To the best of our knowledge, the catalytic activity of K\(_2\)Cl–6H\(_2\)O is not described in the literature but its potential should be a source of interest. It seemed interesting to compare these results with MgCl\(_2\), which is most widely used in organic synthesis. At 150°C, MgCl\(_2\) was characterized by the frequencies of bands at 1,448, 1,608, and 1,635 cm\(^{-1}\). The Lewis acidity of MgCl\(_2\) was therefore lower than the Lewis acidity of K\(_2\)MgCl\(_2\)-6H\(_2\)O. However, unlike K\(_2\)ZnCl\(_4\) and K\(_2\)MgCl\(_2\)-6H\(_2\)O, the spectrum of MgCl\(_2\) showed the presence of a very intense band at 1,537 cm\(^{-1}\) after degassing at 150°C. This band is characteristic of the presence of Bronsted acid sites, unobserved with K\(_2\)MgCl\(_2\)-6H\(_2\)O. These results highlighted the presence of different acidities in these ecocatalysts.

### TABLE 2 | MP-AES Analysis of ecocatalysts derived from Arabidopsis halleri (Eco-A.h.) and Salix “Tordis” (Eco-S.T.) (wt %).

| Ecocatalyst       | K            | Mg           | Ca            | Zn            |
|-------------------|--------------|--------------|---------------|---------------|
| Eco-A. h.         | 11.8 (± 0.65%) | 1.9 (± 4.5%) | 12.5 (± 0.51%) | 7.6 (± 0.18%) |
| Eco-S. T.         | 10.0 (± 0.42%) | 1.5 (± 0.78%) | 19.8 (± 0.44%) | 2.6 (± 1.36%) |
| Eco-Zn (Noccaea caerulescens) | –           | 1.5 (±0.17%)  | –             | 7.3 (±0.4%)   |
| Eco-Zn (Anthyllis vulneraria) | –           | 3.6 (±0.18%)  | –             | 15.4 (±0.7%)  |

### TABLE 3 | Crystalline species in ecocatalysts (√, presence; ✗, lack).

| Catalyst       | Carnallite KMgCl\(_2\)-6H\(_2\)O | Calcium sulfate CaSO\(_4\)·(H\(_2\)O)\(_{0.5}\) | Sylvite KCl | Potassium tetrachlorozincate K\(_2\)ZnCl\(_4\) | Tachhydrite CaMgCl\(_2\)·12H\(_2\)O |
|----------------|----------------------------------|----------------------------------------------|------------|-----------------------------------------------|-----------------------------------|
| Eco-A.h.       | √                                | √                                            | ✗          | ✗                                            | ✗                                  |
| Eco-S.T.       | √                                | ✗                                            | √          | ✗                                            | ✗                                  |
Ecocatalyzed Knoevenagel Condensations

The catalytic activity of each ecocatalyst was tested using the Knoevenagel condensation, a widely used reaction in organic synthesis. Initial reactions were carried out with benzaldehyde and different nucleophiles using the conditions described for zinc dichloride (Shanthan Rao and Venkataratnam, 1991). Reactions were performed neat, at 100°C, during 1 h and using an amount of ecocatalyst which corresponds to 5 mol% of zinc with respect to aldehyde. Control reactions without catalyst exhibited no conversion of benzaldehyde. Results were compared with commercial ZnCl$_2$ and K$_2$ZnCl$_4$ under same conditions (Table 4).

Considering these results, the first unexpected trend is that ecocatalysts are more efficient or comparable than pure synthetic salts in most of cases. In the case of methyl cyanoacetate, Eco-A.h. and Eco-S.T. gave 53 and 84% of conversion, respectively, compared to ZnCl$_2$ and K$_2$ZnCl$_4$ which converted benzaldehyde only at 28 and 44% (Table 4, entry 2). Both ecocatalysts have a better ability to convert acetylacetone and dimethylmalonate while pure salts barely reach 5% (Table 4, entries 3 and 5). Interestingly, malononitrile gave a better result by reacting with ZnCl$_2$ compared to Eco-A.h. and K$_2$ZnCl$_4$, but Eco-S.T. provided an excellent conversion in spite of its weak Zn concentration (Table 4, entry 1). The second unexpected trend is that the catalytic behavior of K$_2$ZnCl$_4$ is disappointing. Except for the special case of malononitrile, K$_2$ZnCl$_4$ provided lower conversions than Eco-A.h.. According to these results, the presence of this salt in Eco-A.h. cannot explain the catalytic efficiency of this ecocatalyst. The third observation is closely linked to the second one: Eco-S.T. which is less concentrated in Zn than Eco-A.h. gave better conversions in four cases (Table 4, entries 1–4) and was comparable in the last case (Table 4, entry 5). Eco-S.T. did not exhibit K$_2$ZnCl$_4$ in XRD analyses but this ecocatalyst provided better results. There is every reason to believe that the active species is not K$_2$ZnCl$_4$. In terms of selectivity, reactions only gave Knoevenagel product excepted for entries 3 and 4. These formed products possess a α,β-unsaturated carbonyl group which can undergo a Michael reaction with another equivalent of nucleophile when the reaction time is extended. To check the range of applicability of these ecocatalysts and the electronic effect of different substituents, the reaction was extended to other benzaldehyde derivatives with acetylacetone as nucleophile (Table 5). The results obtained in Table 5 were coherent with the electronic effects of the substituent on the reaction mechanism. As expected, donor groups lower the conversion (Table 5, entries 1 and 2) whereas attractor groups increased it (Table 5, entries 3 and 4). Additionally, Eco-S.T. led to better results than Eco-A.h. in all cases. This suggested that ZnCl$_2$ and K$_2$ZnCl$_4$, and therefore zinc salts, were not the active species during the catalytic process. Infrared studies in Figure 2 revealed a significant Lewis acidity for KMgCl$_3$·6H$_2$O which was detected in both ecocatalysts by XRD analysis. This salt was thus synthesized in order to be tested. The amount of potassium magnesium trichloride was calculated by considering the magnesium content of Eco-S.T.. Results are summarized in Table 6. The three catalytic systems are active in the Knoevenagel condensations of benzaldehyde with various nucleophiles. In most of cases, KMgCl$_3$·6H$_2$O gave results which are closer to those obtained with Eco-S.T. than MgCl$_2$ or zinc salts (Table 6, entries 1–3). According to these encouraging results, it is clear that KMgCl$_3$·6H$_2$O and its strong Lewis acidic character play a significant role into the catalytic activity applied to the Knoevenagel condensation. It is also important to consider the great interest of using KMgCl$_3$·6H$_2$O in organic synthesis. Indeed, carnallite is a natural ore which the source of magnesium and potassium for industrial needs. This potentiality of the novel Lewis acid in green catalysis required complementary study.

DFT Analysis of Catalytic Salts and Discussion

In view of the results obtained with KMgCl$_3$ in the Knoevenagel condensations, it seemed interesting to compare the reactivity of MgCl$_2$ and KMgCl$_3$. For this, we relied on the concept of Gutmann (Staemmler, 1979; Jensen, 1980), which has been supplemented by Denmark (Denmark and Wynn, 2001). According to Gutmann, the formation of an acid-base adduct
induces an overall increase in the electron density in the acceptor fragment of the adduct. Similarly, Denmark, then Massa (Massa et al., 2010) demonstrated that a low Lewis acid, SiCl₄, can be activated by a Lewis base to promote allylation and propargylation of aldehydes. Similarly, Marcantoni described a surprising catalytic activity for the CeCl₃-NaI system (Bartoli et al., 2006). According to the concept pushed forward by Gutmann, Denmark and the observations of Marcantoni, we can anticipate an interesting reactivity of KMgCl₃ driven by an increasing positive charge of Mg. Indeed, the binding of an additional chloride could enhance the electrophilicity character of the Mg. In this hypothesis, KMgCl₃ should be a better Lewis acid than MgCl₂, which is supported by IR data. We performed DFT calculations to evaluate the electronic redistribution resulting of an additional chloride in KMgCl₃ in comparison with MgCl₂. As shown by the other experiments, the potassium (K⁺) does not seem to interfere in the reactivity. The calculations have therefore been performed without the K⁺ counterion. Furthermore, as it is presumed that the metal is interacting with the benzaldehyde carbonyl during the catalytic process, another set of experiments assesses the electron density of the metal in presence of benzaldehyde (BA). All the experiments were performed in the gas phase (see Experimental section for details). The results are shown in Table 7. The Mulliken population analysis is often considered as fair standard to evaluate the atomic charge, but it suffers a lot from basis set dependency. On the contrary the Merz-Singh-Kollman (MK) charges are known as being rather basis independent. In both cases, there is a clear tendency that can be drawn: the Mg atom is more electropositive when holding an extra Cl. This nicely converges with the Gutmann analysis and the experimental data showing that the MgCl₂⁻ is more reactive than MgCl₂.

**GENERAL CONCLUSION**

The aim of this work was basically to determine if the utilization of the Zn-accumulating plant *Salix Tordis* and the hyperaccumulating plant *A. halleri* was valuable in catalysis. To answer this question, two ecocatalysts were synthesized from biomass used in phytomanagement, in order to be tested in the Knoevenagel condensation reaction. Both ecocatalysts were able to convert aldehydes but interestingly, the conversion promoted...
With these experimental results, we were able to show the relevance of both Zn-accumulating plants. Finally, these new bio-based polymetallic catalysts illustrated the interest of this concept for green and sustainable catalysis.

AUTHOR CONTRIBUTIONS

CG is the principal investigator of the work. P-AD carried out the organic synthesis. VB is responsible for the phytomanagement part. Y-ML and EP are responsible for the DFT calculation. CG and P-AD are responsible for the writing. CB is responsible for the XRD analyses. SD realized the XRD analyses.

ACKNOWLEDGMENTS

Authors would like to thank Franck Pelissier (ChimEco/CNRS-UM UMR 5021) for the MP-AES analysis and ANR (PHYTOCHEM) and ANR (DYNMULTIFEC) programs for financial supports.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2018.00048/full#supplementary-material

REFERENCES

Bartoli, G., Beleggia, R., Giuliani, S., Giuliani, A., Marcantoni, E., Massaccesi, M., et al. (2006). The CeCl3·H2O–NaI system as promoter in the synthesis of functionalized trisubstituted alkenes via Knoevenagel condensation. Tetrahedron Lett. 47, 6501–6504. doi: 10.1016/j.tetlet.2006.07.031

Bisler, B. H., Merz, K. M., and Kollman, P. A. (1990). Atomic charges. Softness a MK charge on Mg

TABLE 6 | GC-MS results of Knoevenagel condensations of benzaldehyde with different nucleophiles catalyzed by magnesium salts.

| Entry | R1  | R2  | KMgCl2 | MgCl2  | Eco-S.T.d |
|-------|-----|-----|--------|--------|----------|
|       | Conv. (%) | Select. (%) | Conv. (%) | Select. (%) | Conv. (%) | Select. (%) |
| 1     | 70   | >99  | 61     | >99    | 93       | 98       |
| 2     | 92   | >99  | 81     | >99    | 84       | 99       |
| 3     | 59   | 65   | 55     | 82     | 33       | 60       |
| 4     | 22   | 96   | 41     | 98     | 10       | –        |

aConversion determined by GC-MS with biphenyl as internal standard; b112 mg of KMgCl2; c60 mg of MgCl2; d614 mg of Eco-S.T.; eKnoevenagel product isolated yield.

| TABLE 7 | Summary of Mulliken, MK charges, and softness on Mg assessed by DFT calculations performed using Gaussian09.

|          | MgCl2 | MgCl2 | BA–MgCl2 | BA–MgCl2 |
|----------|-------|-------|----------|----------|
| Mulliken charge on Mg³⁺ | 0.6505 | 0.996948 | 0.8083 | 1.1180 |
| MK charge on Mg³⁺ | 0.8357 | 0.8382 | 0.7978 | 0.8809 |
| Softness³ | 4.155 | 4.351 | 6.942 | 8.946 |

³Calculated at the B3LYP/6-31G(d,p) level of theory. ²Used for the HSAB concept (Hard Soft Acids and Bases), also known as the Pearson acid base concept (Pearson, 1968), the hardness (nH) and softness (nS) can be estimated using the following formulas (Pearson, 2005): nh = E_LUMO – E_HOMO and nS = 1 2 where E_LUMO = lowest unoccupied molecular orbital energy and E_HOMO = highest occupied molecular orbital energy.

by Eco-S.T. was better than with Eco-A.h. in spite of its lower zinc concentration. While our hypotheses were focused on zinc salt to act as the active specie during the catalytic process, the surprising activity of the magnesium salt KMgCl2 was out of our expectations. The results obtained in catalysis with this salt confirmed its strong Lewis acid character which was first observed in infrared studies and also its remarkable role to convert benzaldehyde compared to K2ZnCl4 and MgCl2. These observations were supported by DFT calculations which showed that magnesium is a better Lewis acid when it possesses three chlorine atoms in its first coordination sphere instead of two.

REFERENCES

Clavé, G., Pelissier, F., and Campidelli, S. (2017). Ecocatalyzed Suzuki cross coupling of heteroaryl compounds. Green Chem. 19, 4093–4103. doi: 10.1039/C7GC01672G

Cui, J., and Zhang, L. (2008). Metallurgical recovery of metals from electronic waste: a review. J. Hazard. Mater. 158, 228–256. doi: 10.1016/j.jhazmat.2008.02.001

Debarge, S., McDaid, P., O’Neill, P., Frahill, J., Wong, J. W., Carr, D., et al. (2014). Evaluation of several routes to advanced pregabalin intermediates: synthesis and enantioselective enzymatic reduction using ene-reductases. Org. Process Res. Dev. 18, 109–121. doi: 10.1021/op4002774

Denmark, S. E., and Wynn, T. (2001). Lewis base activation of lewis acids: catalytic hydroformylation over silica-supported Lewis acids. J. Am. Chem. Soc. 123, 6199–6200. doi: 10.1021/ja016017e

Deyris et al. 2018.00048/full#supplementary-material
