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# Association of Zn- and Mg-based Chain Transfer Agents in Coordinative Chain Transfer Polymerizations of Olefins for Enhanced Control and Activity

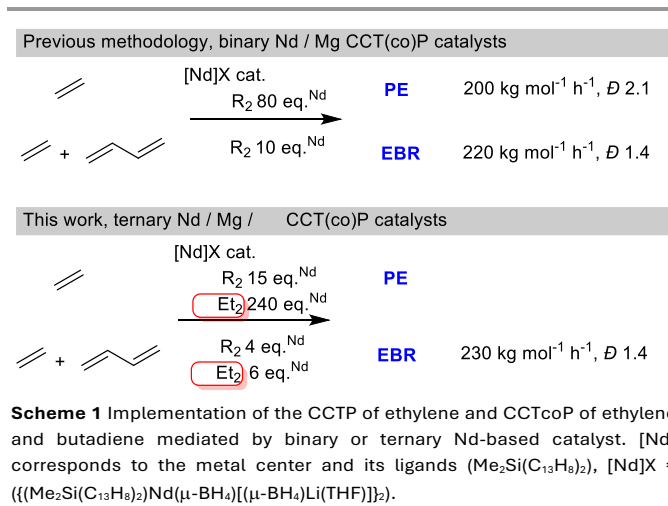
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Based on DFT-level computational studies, ZnEt<sub>2</sub> was implemented as a chain transfer agent (CTA) to promote the coordinative chain transfer (co)polymerization of ethylene for the first time with a metallocene complex of Nd. The use of ZnEt<sub>2</sub> in combination with Mg(<sup>n</sup>Bu)(<sup>n</sup>Oct) compared to the use of Mg(<sup>n</sup>Bu)(<sup>n</sup>Oct) alone, improves several key aspects of the polymerization process. When ethylene is polymerized, an increase of catalytic activities is observed and narrower molar mass distributions are obtained due to a reduced β-H transfer and faster reversible chain transfer reactions. When ethylene is copolymerized with butadiene, ZnEt<sub>2</sub> acts as an excellent CTA and has no impact on the polymerization kinetics and the microstructure of the final copolymer. Using ZnEt<sub>2</sub> in combination with MesMgBr rather than Mg(<sup>n</sup>Bu)(<sup>n</sup>Oct) enables selective chain transfer between neodymium and zinc and promotes a nearly quantitative chain-end functionalization with acyl chloride.

## Introduction

Polyolefins represent the most significant family of polymers in the industry. They are produced in large volumes and at low cost through radical polymerization or coordination catalysis and serve a broad spectrum of applications. A long-standing challenge in this field has been achieving the control of the chain growth to allow the formation of more complex architecture such as block copolymers. Both radical and coordination polymerizations can achieve this by utilizing systems leading to controlled and/or living polymerizations. For instance, recent advancements have applied reversible deactivation radical polymerizations and controlled coordination polymerizations to successfully control the polymerization of ethylene. Although not applied to the high-pressure and high-temperature gas-phase process used for producing low-density polyethylene (LDPE), recent studies have demonstrated that reversible deactivation radical polymerization of ethylene is achievable.<sup>1–4</sup> Besides, since the early 1990s, the development of coordinative chain transfer polymerization (CCTP)<sup>5–8</sup> has garnered increasing interest in both academia and industry. For example, Dow Chemical marketed olefin block copolymers (OBC) produced by this approach.<sup>9</sup> CCTP is a polymerization catalysis based on a degenerative transfer mediated by an organometallic compound such as AlR<sub>3</sub>, MgR<sub>2</sub>, ZnR<sub>2</sub> that acts as a reversible chain transfer agent (CTA).<sup>10</sup> Polymer chains grow on the catalyst and are reversibly transferred to the CTA, where they become intermittently dormant. In the absence of β-H elimination and with an excess of CTA relative to the catalyst, the polymerization process becomes controlled—meaning the number of polymer chains is fixed and directly proportional to the amount of CTA.

As a result, the degree of polymerization is predictable based on the ratio of monomer consumed to the initial amount of CTA used. Since the chains produced by CCTP remain connected to a metal center throughout the polymerization, this method not only allows for the production of precision polyolefins,<sup>11–13</sup> but also facilitates the introduction of functional groups at their chain ends<sup>14</sup> and gives access to diverse macromolecular architectures.<sup>15–20</sup> The CTA itself can be active under polymerization conditions. In the case of copolymerization of 1,3-dienes and styrene with lanthanide borohydride complexes, it is possible to adjust the microstructure of the copolymers by regulating the amount of MgR<sub>2</sub>.<sup>21–23</sup> In the same works, Zinck et al. have also shown the beneficial use of ternary systems based on the association of MgR<sub>2</sub> and AlR<sub>3</sub> to control the stereospecificity of 1,3-diene polymerization by CCTP. Neodymium-based metallocenes enable coordinative chain transfer (co)polymerization (CCT(co)P) of ethylene and butadiene using MgR<sub>2</sub> as the CTA such as Mg(<sup>n</sup>Bu)(<sup>n</sup>Oct) (BOMAG) (Scheme 1).<sup>24–26</sup> The unique properties of MgR<sub>2</sub> in this CCTP process has been elucidated through a theoretical mechanistic study.<sup>25</sup> ZnEt<sub>2</sub> is commonly used in association with a large panel of olefin polymerization catalysts resulting from the combination of Fe-, Hf- or Zr-based complexes with borate activators or methylaluminoxane (MAO).<sup>7,8</sup> In these systems, the underlying reaction mechanisms remain intricate. In addition, the group of Sita suggested that ZnEt<sub>2</sub> can mediate the chain transfer between the active catalyst and the second co-catalyst.<sup>13</sup>



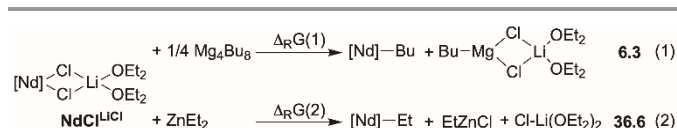
In light of these results, developing more efficient and versatile conditions for Nd-based CCTP is of significant interest, especially when compared to those achieved with MgR<sub>2</sub> as a lone CTA. However, a major challenge in using ZnEt<sub>2</sub> as a CTA with Nd-based pre-catalysts is its inability to activate the Nd precursor.<sup>27</sup>

In this communication, we circumvent this issue by combining ZnEt<sub>2</sub> and MgR<sub>2</sub> or RMgX with ((Me<sub>2</sub>Si(C<sub>13</sub>H<sub>8</sub>)<sub>2</sub>)Nd(μ-BH<sub>4</sub>)[(μ-BH<sub>4</sub>)Li(THF)]<sub>2</sub>) Nd-based complex **1** (Scheme S1). Building on our previous expertise with MgR<sub>2</sub> as an efficient CTA for the CCTP of ethylene and the CCT(co)P of ethylene and butadiene using a Nd-based catalyst,<sup>28–30</sup> and supported by preliminary DFT-level computational investigations, we evaluated the potential of ZnEt<sub>2</sub> as a CTA in MgR<sub>2</sub>-mediated CCTP systems. The resulting ternary Nd / Mg / Zn catalysts were then investigated for the polymerization of ethylene and its copolymerization with butadiene.

## Results and discussion

### Insights from molecular modelling

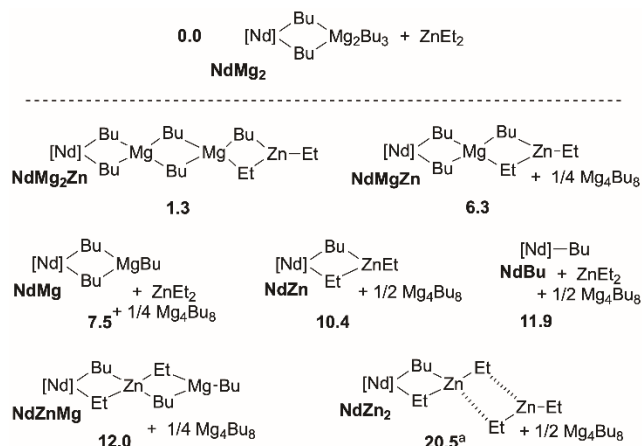
Before implementing the experimental design for the ternary catalytic system **1** / BOMAG / ZnEt<sub>2</sub>, we first obtained structural and mechanistic insights through DFT-level molecular modelling. This study presents several challenges, including the large size of the system (> 100 atoms), the presence of three different metals, and multiple polymer chains. To address these challenges, we relied on our previous theoretical modelling strategies for speciation of heteropolymetallic complexes and the exploration of chain exchange reactions.<sup>25,31</sup> To reduce computational costs, BOMAG was modelled using Mg(<sup>n</sup>Bu)<sub>2</sub> clusters, with the rhombohedral tetramer being selected as the most stable configuration.<sup>25</sup> Regarding ZnEt<sub>2</sub>, optimization of a dimer consistently leads to its dissociation into monomers. Since ZnEt<sub>2</sub> is purchased and used in a hydrocarbon solution, the most stable form present in this context is considered to be the monomer. The fully detailed computational method is described in the [Supporting Information](#).



**Scheme 2** Thermodynamics ( $\Delta_R G$ , kcal mol<sup>-1</sup>) of [Nd]-R species formation (R = <sup>n</sup>Bu or Et) as a function of the alkylating agent. [Nd] corresponds to the metal center and its Me<sub>2</sub>Si(C<sub>13</sub>H<sub>8</sub>)<sub>2</sub> ligand. Gaussian 09, B3PW91/D3BJ(SP), SMD(TolH), 298 K, 1 atm.

To validate the computational method, the thermodynamics of the formation of alkylated monometallic species [Nd]-R (R = <sup>n</sup>Bu or Et, [Nd] = Me<sub>2</sub>Si(C<sub>13</sub>H<sub>8</sub>)<sub>2</sub>Nd) from the Nd precursor Me<sub>2</sub>Si(C<sub>13</sub>H<sub>8</sub>)<sub>2</sub>Nd(μ-Cl)<sub>2</sub>Li(OEt)<sub>2</sub> (**NdCl<sup>LiCl</sup>**) was computed for both Zn and Mg alkylating agent (**Scheme 2**). This precursor has been taken as both as model and analog of **1**. While the activation reaction of **NdCl<sup>LiCl</sup>** is endergonic by 6 kcal mol<sup>-1</sup> in the case of Mg(<sup>n</sup>Bu)<sub>2</sub>, it is endergonic by 37 kcal mol<sup>-1</sup> in the case of ZnEt<sub>2</sub>. This is in line with experimental results for this specific system,<sup>27</sup> but also matches with the activation conditions reported for other polymerization catalytic systems.<sup>32,33</sup>

As suggested in the literature and mentioned above, ZnEt<sub>2</sub> may mediate the chain transfer between the catalyst and a second CTA, namely AlR<sub>3</sub>.<sup>13</sup> This mechanism hypothesis has been considered and will be compared to the chain transfer mechanism previously reported for the CCTP of ethylene mediated by BOMAG and [Nd]-<sup>n</sup>Bu.<sup>25</sup> Of note, the latter revealed that the chain exchange reaction proceeds via a two-step association / dissociation transmetalation in which the intermediate heterotrimetallic complex [Nd]-(μ-Bu)<sub>2</sub>-Mg<sub>2</sub>Bu<sub>3</sub> (**NdMg<sub>2</sub>**) is the resting state of the polymerization catalyst.<sup>25</sup> Considering now the addition of ZnEt<sub>2</sub> as a third component of the catalytic system, the speciation of thermodynamically accessible complexes has been performed.



**Scheme 3** Speciation of the alkylated species yielded from [Nd]-<sup>n</sup>Bu, Mg,<sup>n</sup>Bu<sub>8</sub> and ZnEt<sub>2</sub>. Only the most stable species are shown, the fully detailed speciation is provided in [SI](#).  $\Delta_R G$  in kcal mol<sup>-1</sup>. <sup>a</sup>In **NdZn<sub>2</sub>**, interaction between the ZnEt<sub>2</sub> fragments is weak, the endergonicity results from the entropy cost of the association of free and separated ZnEt<sub>2</sub>. Gaussian 09, B3PW91/D3BJ(SP), SMD(TolH), 298 K, 1 atm.

Relative to **NdMg<sub>2</sub>** and ZnEt<sub>2</sub>, a set of complexes can be formed with an endergonicity ranging from 1 to 10 kcal mol<sup>-1</sup> (**Scheme 3**). As a general trend, relative to separated **NdMg<sub>2</sub>** and ZnEt<sub>2</sub>,

the formation of complexes in which zinc is the  $\beta$  position to neodymium is endergonic by at least 10 kcal mol<sup>-1</sup>. Interestingly, the formation of the heterobimetallic dimer **NdZn** is almost as endergonic as the release of the **NdBu** complex from which monomer coordination-insertion mechanism takes place.<sup>25</sup>

From a kinetic point of view, the Gibbs energy barrier for ethylene insertion in the Nd–C bond of the **NdBu** complex is computed to 10.7 kcal mol<sup>-1</sup>. Concerning, the Nd / Mg binary system, in presence of Mg<sub>4</sub>Bu<sub>8</sub>, the overall Gibbs energy barrier is raised to 22.6 kcal mol<sup>-1</sup> relative to **NdMg<sub>2</sub>** and ethylene due to the formation of the **NdMg<sub>2</sub>** dormant complex.<sup>25</sup> In the Nd / Mg / Zn ternary system, the presence of zinc leads to the formation of the bimetallic species **NdZn** as an intermediate that can become the most abundant Nd-based complex along with **NdMg** if ZnEt<sub>2</sub> is added in large excess relative to BOMAG, the amount of **NdMg<sub>2</sub>** becoming small. **NdZn** and **NdMg** becoming the most abundant resting state of the catalyst, the overall Gibbs energy barrier for ethylene insertion is decreased from 22.6 kcal mol<sup>-1</sup> in a window ranging from 12.2 to 15.1 kcal mol<sup>-1</sup>. Thereof, adding an excess of ZnEt<sub>2</sub> to the catalytic mixture would lead to a significant boost in catalytic activity. Regarding the influence on the chain transfer, we were unable to adequately converge the geometry of the transition state(s) as it relies on association / dissociation processes, coupled with re-orientation of the chain end carbanion doublet by rotations.<sup>25,26</sup> However, an energy barrier of less than 5 kcal mol<sup>-1</sup> can be roughly estimated leading to an overall fast exchange between Nd and Zn, this barrier is thereof lower than the one computed between Nd and Mg by 8 kcal mol<sup>-1</sup>,<sup>25</sup> thus enabling a much faster chain shuttling. Finally, as ZnR<sub>2</sub> and presumably Zn(polymer)<sub>2</sub>, do not aggregate, a more straightforward chain transfer between Nd and Zn may be inferred.

### Polymerization of ethylene and copolymerization of ethylene with butadiene

**Preliminary experimental observations.** Molecular modelling highlights that though ZnEt<sub>2</sub> is not capable of alkylating the complex **1**, its addition in excess relative to MgR<sub>2</sub> would enhance both the catalytic activity and the chain transfer leading to a fastest polymerization producing chains with more narrowly distributed molar masses. Thereof, we compared ethylene polymerization mediated by **1** / BOMAG catalyst with that mediated by **1** / BOMAG / ZnEt<sub>2</sub> ternary system, keeping the [CTA] / [Nd] ratio constant (Table 1). 4 g of PE was targeted corresponding to a theoretical  $M_n$  of 2000 g mol<sup>-1</sup>.

In a first blank experiment, ethylene was polymerized with the **1** / BOMAG catalyst ([Nd]:[Mg] = 1:80, run 1, Table 1). In agreement with our previous works,<sup>25</sup> the catalyst meets the requirement of CCTP but the chain exchange rate is low compared with the propagation rate as indicated by a dispersity value of 2.1. This has been attributed to a high-energy barrier for dissociation of the heterobimetallic species formed between the active species and BOMAG.<sup>25</sup> When the ternary system **1** / BOMAG / ZnEt<sub>2</sub> ([Nd]:[Mg]:[Zn] = 1:5:75) was used (run 2, Table 1), complex **1** was first activated by BOMAG in a separate vial and the mixture was transferred in a solution containing ZnEt<sub>2</sub> before starting the polymerization. A strong decrease of  $\bar{D}$

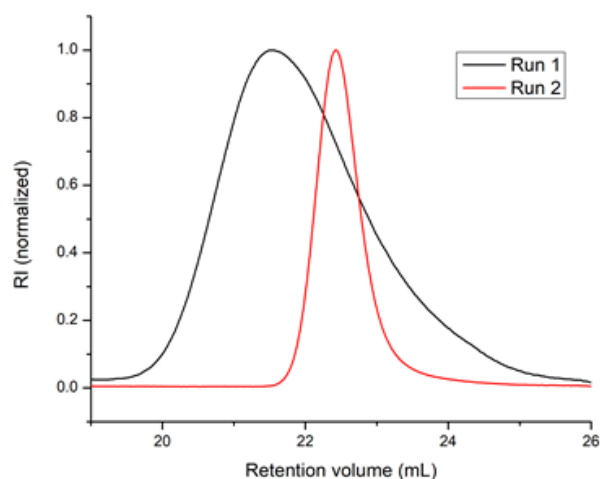
from 2.1 to 1.1 (Table 1 and Figure 1) is observed and the content of vinyl-terminated chains resulting from  $\beta$ -H elimination drops from 10 to 2%.

We also observe a good agreement between the theoretical number-average molar mass ( $M_n^{\text{theo}}$ ) calculated considering that ZnEt<sub>2</sub> is concomitantly acting as chain transfer agent with BOMAG and the experimental  $M_n$  confirming that ZnEt<sub>2</sub> definitively acts as a CTA in this CCTP.

**Table 1** Investigation of the influence of ZnEt<sub>2</sub> on **1** / BOMAG catalysts during ethylene polymerization.

run	[Nd]: [Mg]: [Zn]	yield g	activity kg mol <sup>-1</sup> h <sup>-1</sup>	$M_n^{\text{theo}}$ g mol <sup>-1</sup>	$M_n(\bar{D})^a$ g mol <sup>-1</sup>	PE <sup>vinyl</sup> % <sup>b</sup>
1	1:80:0	3.6	200	1800	2400(2.1)	10
2	1:5:75	4.1	3700	2050	2200(1.1)	2

80°C, toluene, [Nd] = 62.5  $\mu$ M,  $M_n^{\text{theo}}$  = yield/2\*(nMg + nZn); <sup>a</sup>determined by HT-SEC with PE standards; <sup>b</sup>determined by <sup>1</sup>H NMR.



**Figure 1** SEC traces of polyethylene obtained with **1** / BOMAG (run 1, Table 1) and **1**/BOMAG/ZnEt<sub>2</sub> (run 2, Table 1) catalysts.

In addition, a sharp increase in activity from 200 to 3700 kg mol<sup>-1</sup> h<sup>-1</sup> is obtained, in good agreement with the modelling study. Indeed, the formation of a heterobimetallic **NdZn** (Scheme 3) formed between the active species Me<sub>2</sub>Si(C<sub>13</sub>H<sub>8</sub>)<sub>2</sub>NdR and ZnEt<sub>2</sub> is exergonic by only 1.5 kcal mol<sup>-1</sup>, whereas the heterobimetallic **NdMg<sub>2</sub>** (Scheme 3) species formed with BOMAG is exergonic by 11.9 kcal mol<sup>-1</sup>. Consequently, the overall energy barrier of the ethylene insertion is lowered in the presence of ZnEt<sub>2</sub> and chain exchange is much faster between Nd and zinc than between Nd and magnesium. However, at this stage of our study, chain exchange between magnesium and zinc is not demonstrated. To evaluate a potential chain transfer between magnesium and zinc, a model reaction was performed in an NMR tube. The <sup>1</sup>H NMR analysis of BOMAG and ZnEt<sub>2</sub> was carried out (Figure S1 and S2), followed by that of a BOMAG/ZnEt<sub>2</sub> mixture (Mg/Zn = 0.8, Figure S3 to S7). Three new signals are observed. Two triplets at 0.23 and 0.27 ppm and a quadruplet at 0.13 ppm (Figure S4). An assignment of the two triplets is better performed when the NMR analysis is carried with a spectrometer operating at 500

MHz (Figure S5). As the signals are as well defined as those of ZnEt<sub>2</sub>, it is reasonable to assume the formation of ZnEtR (R = <sup>n</sup>Oct and <sup>n</sup>Bu) and to confirm an exchange of alkyl groups between zinc and magnesium under these conditions.

**Investigation of the 1 / BOMAG / ZnEt<sub>2</sub> catalyst system for ethylene polymerization.** A series of experiments was further carried out by varying the Nd/Mg/Zn ratio. The overall concentration in Mg and Zn was kept constant, while [1] and [Mg]/[Zn] ratio were varied (Table 2). Given the low percentage

of PE-vinyl observed at 80°C when ZnEt<sub>2</sub> is used, the polymerization temperature was set at 90°C to improve polyethylene solubility and reach higher molar masses while keeping the control of the polymerization. As expected, a higher yield resulted in higher molar masses (runs 7 and 8 in Table 2). Besides, considering the strong increase in activity observed when ZnEt<sub>2</sub> is used, lower concentrations of 1 could be used (run 4, Table 2) emphasizing that neodymium metallocene catalysts offer performances close to that of the best molecular catalysts for ethylene polymerization.<sup>34</sup>

**Table 2** Investigation of the ternary catalyst 1 / BOMAG / ZnEt<sub>2</sub> in ethylene polymerization.

run	[Nd]:[Mg]:[Zn]	yield	time	activity	$M_n^{\text{theo}}$	$M_n^{\text{a}}$	$\bar{D}$	PE-vinyl
		g	min	kg mol <sup>-1</sup> h <sup>-1</sup>	g mol <sup>-1</sup>	g mol <sup>-1</sup>		% <sup>b</sup>
3	1:255:0	3.9	207	290	1950	2000	2.2	17
4	1:15:240	4.8	8	9200	2400	1900	1.2	5
5	1:60:20	4.1	48	410	2050	1400	1.6	29
6	1:40:40	4.9	14	1700	2450	2000	1.4	17
7	1:20:60	5.3	6	4600	2650	1800	1.4	5
8	1:20:60	9.7	12	3100	4850	4900	1.2	11
9	1:5:75	3.6	7	2500 <sup>c</sup>	1800	1400	1.2	5

[Nd] = 20 μM (runs 3 and 4), 62.5 μM (runs 5 to 9), 200 mL toluene, 4 bar, 90°C (run 8 at 80°C),  $M_n^{\text{theo}} = \text{yield}/2*(n\text{Mg} + n\text{Zn})$ ; <sup>a</sup>determined by HT-SEC with PE standards; <sup>b</sup>determined by <sup>1</sup>H NMR; <sup>c</sup>total deactivation of the catalytic system at the end of polymerization.

In the present case, ZnEt<sub>2</sub> increases the quantity of active neodymium as shown by the computational investigation. The good agreement between theoretical and experimental  $M_n$  indicates that polymerizations are controlled according to CCTP. The final PE show narrow molar mass distributions and the content of PE-vinyl chains decreases when the amount of ZnEt<sub>2</sub> increases. Decreasing the [Mg]/[Zn] ratio (runs 5-7, Table 2) leads to an increase in activity. When the lowest [Mg] is used (run 9, Table 2), a sudden deactivation of the catalyst is observed at the end of polymerization (Figure S8). The deactivation process appears to be linked to the increase in [Nd]-P active species in the presence of ZnEt<sub>2</sub>. This deactivation might occur from inter- or intra-molecular reactions between the active species, reactions that are not identified yet even by molecular modelling. In summary, the dialkylmagnesium compound not only acts as an alkylating agent for 1 but also brings stability to the catalytic system via a mechanism that still not understood at the molecular level. Building on these promising results on the PE chain transfer ability when additional ZnEt<sub>2</sub> is employed, the combination of 1 and mixed CTA based on Mg and Zn was further evaluated in the coordination chain transfer copolymerization (CCT(co)P) of ethylene and butadiene for the synthesis of ethylene butadiene rubber (EBR).

**Investigation of 1 / BOMAG / ZnEt<sub>2</sub> catalyst system for the copolymerization of ethylene (E) and butadiene (B).**-A first preliminary test was performed with ZnEt<sub>2</sub> alone combined with 1 ([Zn]/[Nd] = 5) and a [B]/[E] monomer feeding ratio of 80/20, at 70°C in toluene but as expected no polymerization was observed. Nevertheless, the addition of 4 equivalents of Mg(<sup>n</sup>Bu)(<sup>n</sup>Oct) (BOMAG) in the polymerization medium activates the metallocene and an EBR was isolated. The

determination of the  $M_n$  highlighted that both BOMAG and ZnEt<sub>2</sub> acted again as CTA.

Copolymerizations of ethylene and butadiene were then carried out with a ([Mg]+[Zn])/[Nd] ratio fixed to 10 and targeting 15 g of polymer, corresponding under these conditions to a theoretical  $M_n^{\text{theo}}$  of 15000 g mol<sup>-1</sup> (Table 3). Very similar molar masses (between 12500 and 14200 g mol<sup>-1</sup>) consistent with the targeted value were obtained for various Mg/Zn ratios showing again here that both BOMAG and ZnEt<sub>2</sub> are acting as CTAs.

**Table 3** Copolymerization of ethylene with butadiene using 1 / BOMAG / ZnEt<sub>2</sub> ternary catalytic system.

run	[Nd]:	yield	time	activity	$M_n^{\text{a}}$	$\bar{D}$
	[Mg]:	g	min	kg mol <sup>-1</sup> h <sup>-1</sup>	g mol <sup>-1</sup>	
	[Zn]					
10	1:10:0	14.9	81	220	13800	1.4
11	1:8:2	15.4	87	210	14200	1.5
12	1:5:5	14.8	94	190	13800	1.5
13	1:4:6	14.8	76	230	12500	1.4
14 <sup>b</sup>	1:2:8	1.6	35	60 <sup>b</sup>	-	-

[Nd] = 250 μM, 4 bar (ethylene/butadiene = 80/20), 90°C, 200 mL toluene; <sup>a</sup>determined by SEC in THF using an universal calibration; <sup>b</sup>low activity then total deactivation of the catalytic system after 35 minutes of polymerization.

For CCTcoP of ethylene and butadiene, the number of polymer chains is fixed by the number of magnesium-alkyl and zinc-alkyl bonds. For both binary and ternary catalytic systems, similar molar mass distributions are obtained (Figure S9), but contrary to ethylene homopolymerization, the use of ZnEt<sub>2</sub> as co-CTA in CCTcoP of ethylene and butadiene does not raise the catalytic activity nor narrow the molar mass distribution that respectively remain around 200 kg mol<sup>-1</sup> h<sup>-1</sup> and 1.4 (Table 3). The microstructure of the EBR formed in the presence of ZnEt<sub>2</sub> is similar to the one of the EBR obtained with BOMAG alone

(Table 4). This suggests that the same active species are formed in both systems. Indeed, whereas the overall energy barrier for ethylene insertion includes the dissociation of dormant heterobimetallic species into the active species, the kinetics of the copolymerization of ethylene and butadiene is controlled by monomer insertions into the Nd-allyl and the Nd-(vinylcyclohexyl)methylene active chain ends, as demonstrated by joint experimental and computational mechanism investigation.<sup>26,31</sup> The nature of the CTA has thus very little impact on the kinetics.

As in the case of ethylene polymerization (run 9, Table 2), the catalytic system deactivates at low Mg/Nd ratios (run 14, Table 3). In CCTcoP case, since the initial activity for this run is low (60 kg mol<sup>-1</sup> h<sup>-1</sup>, Table 3), the deactivation of the catalytic system may result from the inefficient activation when using only 2 equivalents of magnesium relative to neodymium.

**Table 4.** Microstructure of EBR obtained with **1** / RMgX / ZnEt<sub>2</sub> ternary catalysts (X = Bu or Mes) determined by <sup>1</sup>H NMR.

Run	Nd/Mg/Zn	E/B	Vinyl %	Trans-1,4 %	1,2-cyclohexyl %	Ethylene <sup>a</sup> %
10	1/10/0	80.0/20.0	7.7	6.0	10.4	75.9
11	1/8/2	81.2/18.8	6.3	5.9	10.7	77.2
12	1/5/5	80.7/19.3	6.6	6.2	10.7	76.6
13	1/4/6	81.0/19.0	6.4	6.0	10.7	76.9
17	1/16/10	79.7/20.3	8.6	5.8	9.9	75.8

<sup>a</sup> Except those included in cyclic units.

**Functionalization of PE and EBR chains obtained using optimized ternary catalyst systems.** In addition to the improved performances provided using a combination of the two chain transfer agents, the advantage of using zinc is the selectivity of organozinc compounds towards certain chemical reactions that can be advantageously used for chain-end

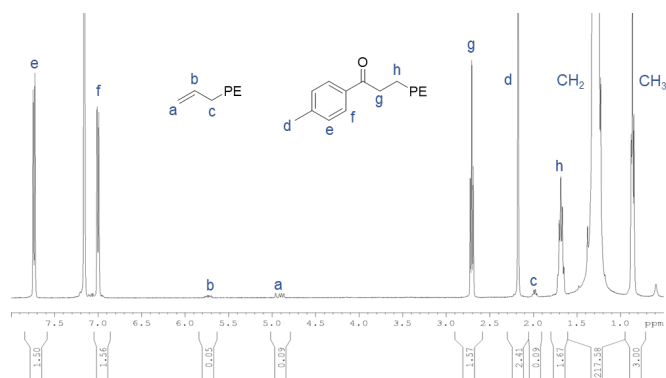
functionalization. For example, whereas reactions between organomagnesium compounds and acyl chlorides are poorly selective towards ketone formation<sup>35-37</sup> unless specific additives are used,<sup>38</sup> organozinc compounds shows a good selectivity.<sup>39</sup> In the perspective of chain-end functionalization, a potential limitation of the ternary systems depicted here above is the storage of polymer chains on both zinc and magnesium atoms. Therefore, identifying a combination of organozinc and organomagnesium compounds that can efficiently alkylate compound **1** with the organomagnesium and facilitate reversible chain transfer on Zn centres will ultimately enable the selective end-functionalization of polymer chains produced by CCT(co)P. Of interest, we have recently shown that MesMgBr is not an activator for the complex **1** but nevertheless reacts with **1**, as shown by the solubilization of the resulting species in toluene.<sup>19,40</sup> As a consequence, our strategy was to use MesMgBr exclusively as an arylating agent for **1** and ZnEt<sub>2</sub> as the CTA. The catalytic ternary system **1** / MesMgBr / ZnEt<sub>2</sub> was first assessed in CCTP of ethylene (Table 5).

Here again, 4 g of PE was targeted, corresponding to a theoretical  $M_n$  of 2000 g mol<sup>-1</sup> considering that ZnEt<sub>2</sub> governs the number of chains formed, *i.e.* all chains were initiated by an ethyl group (Table 5). Whatever the reaction conditions used, polymerization takes place. Considering that ZnEt<sub>2</sub> does not alkylate **1** and that the Nd-Mes was previously shown not to initiate ethylene polymerization,<sup>40</sup> initiation of the polymerization presumably proceeds via an arylation of the Nd center by MesMgBr followed by a transmetalation between Nd-Mes and Zn-Et bonds. This was indeed confirmed by a good agreement between experimental and theoretical  $M_n$ . The reversibility of the chain transfer between Nd and Zn centers is also confirmed by lower the dispersity values obtained ( $\mathcal{D} < 1.3$ ) compared to the one obtained when the experiment was conducted with BOMAG alone ( $\mathcal{D} = 2.1$ , run 1, Table 5).

**Table 5** Optimization of **1** / MesMgBr / ZnEt<sub>2</sub> catalyst for ethylene polymerization or ethylene and butadiene copolymerization and subsequent chain end functionalization by reaction with an acyl chloride.

run	Nd:Mg:Zn	yield	time	activity	$M_n^{\text{theo}}$	$M_n^c$	$\mathcal{D}$	PE-vinyl	PE(EBR)-C(O)C <sub>6</sub> H <sub>4</sub> X
		g	min	kg mol <sup>-1</sup> h <sup>-1</sup>	g mol <sup>-1</sup>	g mol <sup>-1</sup>		% <sup>d</sup>	%
1 <sup>a</sup>	1:80:0	3.6	82	200	1800	2400	2.1	10	-
15	1:20:80	4.2	31	690	2010	2000	1.1	3	79 <sup>d</sup>
16	1:8:80	3.5	35	500 <sup>e</sup>	1750	1700	1.1	4	86 <sup>d</sup>
17 <sup>b</sup>	1:16:10	10.3	174	70	10000	16400	1.3	-	34 <sup>f</sup>

[Nd] = 62.5 μM, 4 bar, 80°C, 200 mL toluene; <sup>a</sup>reference test with BOMAG; <sup>b</sup>[Nd] = 250 μM, 4 bar (ethylene/butadiene feed ratio: 80/20), 80°C, 200 mL toluene; <sup>c</sup>determined by HT-SEC with PE standards for PE samples and by SEC-THF with universal calibration for the EBR sample; <sup>d</sup>determined by <sup>1</sup>H NMR; <sup>e</sup>partial discoloration of the solution, loss of exotherm and decrease of ethylene consumption; <sup>f</sup>functionalization yield =  $M_n^{\text{SEC}}/M_n^{\text{RMN}}$ .



**Figure 2**  $^1\text{H}$  NMR spectrum of the *p*-methylphenone-PE obtained with the ternary catalytic system **1** / MesMgBr / ZnEt<sub>2</sub> after deactivation with *p*-toluoyl chloride (run 16, Table 5).

Concerning chain-end functionalization, preliminary results gathered on the reactivity of MgPE<sub>2</sub> prepared with (C<sub>5</sub>Me<sub>5</sub>)<sub>2</sub>NdCl<sub>2</sub>Li(OEt)<sub>2</sub>/BOMAG binary catalyst reveals that functionalization upon the reaction with two equivalents of *p*-toluoyl chloride does not provide the expected ketone functionalized PE chains. Instead, a mixture of secondary and tertiary alcohols (PE-CH(OH)-(C<sub>6</sub>H<sub>4</sub>Me): 25 %, PE<sub>2</sub>-C(OH)-(C<sub>6</sub>H<sub>4</sub>Me): 27 %, PE-vinyl (30 %) et PE-CH<sub>3</sub> (16 %) (Scheme S2 and Figure S10) is obtained. Using **1** / MesMgBr / ZnEt<sub>2</sub> ternary system, chain end functionalization with two equivalents of *p*-toluoyl chloride relative to Zn was performed (runs 15 and 16, Table 1). A representative  $^1\text{H}$  NMR spectrum of the final PE obtained is shown on Figure 2 for run 16 (Table 5). The assignment of the signals performed on Figure 2 strongly supports the selective formation of the targeted *p*-methylphenone-PE and shows the absence of any other signals including those related to chains initiated by a mesityl moiety. High functionalization yields from 79 to 86% were obtained using only one equivalent of acyl chloride per Zn-alkyl bond. This selectivity outcome reinforces the role played by Zn in mostly supporting PE chains and eludes an efficient chain transfer reactions between Nd or Zn with Mg. Eventually, run 16 (Table 5), with Mg / Nd = 8, shows a slowdown in polymerization rate over time. As already mentioned in the case of BOMAG (Table 2), an excess of magnesium compound compared to neodymium is required to achieve stable activity.

**1** / MesMgBr / ZnEt<sub>2</sub> catalyst system was further tested for the copolymerization of ethylene and butadiene (run 17, Table 5). An EBR (20.3 % of inserted butadiene) with a similar microstructure (Table 4) to that obtained when BOMAG is used (run 10, Table 3) was isolated however with lower activity. The polymerization medium was deactivated by addition of 4-dimethylaminobenzoyl chloride. The  $^1\text{H}$  NMR spectrum shows

the characteristic signal of methyl groups from -NMe<sub>2</sub> group in agreement with the formation of *p*-dimethylaminophenone chain end (Figure S11). A functionalization rate of 34% was measured. This relatively low functionalization efficiency in comparison to functionalization reaction conducted on PE chains can be ascribed to the different nature of the chain ends in the two systems<sup>26</sup> and their respective reactivity but also to the higher molar mass obtained for EBR.<sup>41</sup>

## Conclusions

In this work, we have demonstrated that ZnEt<sub>2</sub> is an effective CTA when used in combination with **1** for CCTP of ethylene or CCTcoP of ethylene and butadiene to produce EBR. However, being unable to alkylate **1** when used alone, the effectiveness ZnEt<sub>2</sub> as CTA depends on the additional presence of an organomagnesium compound that additionally stabilize the resulting catalyst. Compared to magnesium, the storage of the polymer chains on zinc modifies the reactivity of the chain ends, enabling a selective terminal ketone formation through deactivation reactions of the polymerization medium with acyl chlorides. Despite this, the functionalization rate remains low for EBR, making the synthesis of highly functionalized EBRs with high molar masses a future challenge. Additionally, combining ZnR<sub>2</sub> and MgR<sub>2</sub> CTA could open new avenues for integrating Nd metallocenes in original catalyst combinations for the production of original OBC. This approach holds the potential to combine the properties of polyolefins exclusively accessible with Nd metallocene with other polyolefins.

## Author Contributions

The original ideas were conceived by the corresponding authors FdA, MEP, and CB in collaboration with industrial partners FJBdD and RN. Experimental work was carried out by AD, VL and NB, while the computational studies were conducted by LV. The manuscript was written collectively by all the authors.

## Conflicts of interest

There are no conflicts to declare.

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ZnEt<sub>2</sub> is used for the first time as a reversible transfer agent in olefin polymerization with a neodymium metallocene catalyst. Its presence has improved the catalyst's performance for the coordinative chain transfer polymerization of ethylene.

