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Tuning the catalytic properties of rare earth borohydrides for the polymerisation of isoprene**

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Supporting information:
[†] CCDC 892289 and 892290. For crystallographic data in CIF or other electronic format see http://dx.doi.org/10.1039/C2DT31624B

Graphical abstract:
Abstract

Previous results obtained for the cis-polymerisation of isoprene with scandium half-sandwich complex \( \text{Cp}^* \text{Sc(BH}_4\text{)}_2(\text{THF}) \) \( 1\text{a} \) were extended to its neodymium analog. The X-ray structure of the already reported neodymium monomer compound \( \text{Cp}^* \text{Nd(BH}_4\text{)}_2(\text{THF})_2 \) \( 1\text{b} \) is presented.

\( \text{Cp}^* \text{Nd(BH}_4\text{)}_2(\text{THF})_2/\text{[CPh}_3\text{]}\text{[B(\text{C}_6\text{F}_5\text{)}_4]/Al(\text{tBu})}_3 \) catalytic system was found to be very active and stereoselective towards isoprene polymerisation, leading to highly 1,4-cis polyisoprene up to 92\%. The effect on isoprene polymerisation of the addition of a NHC molecule to \( \text{Cp}^*\text{Ln(BH}_4\text{)}_n(\text{THF})_n \) pre-catalyst (\( \text{Cp}^* = \eta^5\text{C}_5\text{Me}_5, \text{Ln} = \text{Sc}, n = 1, 1\text{a}; \text{Nd}, n = 2, 1\text{b}; \text{Sm}, n = 2, 1\text{c} \)) or to a trisborohydride \( \text{Ln(BH}_4\text{)}_3(\text{THF})_n \) (\( \text{Ln} = \text{Sc}, n = 1.5, 2\text{a}; \text{Nd}, n = 3, 2\text{b}; \text{Sm}, n = 3, 2\text{c} \)) was also studied. Several NHC ligands were assessed: the classical \( [1\text{-}\text{C}\{(\text{N}_t\text{BuCH})_2\} ] (L_1) \) and functional N-heterocyclic carbenes, two amino-tethered \( \text{HNBu}_2\text{CH}_2\text{CH}_2[1\text{-}\text{C}\{\text{N(CHCH)NR}\}] (HL_2-R) (R = \text{tBu, Mes (Mes = 2,4,6-Me}_3\text{-C}_6\text{H}_2)) \) and the hydroxyl-tethered \( \text{HOCMe}_2\text{CH}_2[1\text{-}\text{C}\{\text{N(CHCH)N}_i\text{Pr}\}] (HL_3) \). Neodymium-based complex \( (L_2\text{-tBu})\text{Nd(BH}_4\text{)}_2(\text{THF})_3 \) could be isolated and characterized. With some of the catalytic combinations tested, the introduction of the NHC ligand in the coordination sphere of the complex induces a switch of the stereoselectivity of the resulting polymer. Scandium complex \( 2\text{a} \), which produces mainly 1,4-cis polyisoprene when associated to a borate activator and aluminum alkyl, leads to 1,4-trans polymer up to 94\% regular when \( L_2\text{-tBu} \) carbene is added to the same reaction mixture. This result is the only example of highly trans-polyisoprene synthesized with a scandium based catalyst. Coordination of the carbene moiety to the rare earth centre is confirmed by NMR studies on paramagnetic neodymium pre-catalysts.

Introduction

Olefin elastomers are of primary industrial interest as they are used as high-performance materials in high-tech sectors such as tyres, sport accessories or adhesives.\(^1\) Among them, 1,4-cis- and 1,4-trans-polyisoprenes both exist as natural elastomers issued from Hevea Brasiliensis and Gutta Percha, respectively. They can alternatively be synthesized industrially at high scale by polymerisation of isoprene using stereoselective Ziegler–Natta type polymerisation catalysts.\(^2,3\) In this context, half-sandwich and monosubstituted derivatives of the rare earths have proved to behave as highly valuable catalysts towards the controlled (co)polymerisation of conjugated dienes, as highlighted by recent reports.\(^4-14\) Many very efficient catalytic systems suffer however from one main drawback that may hinder their development: the active species are generally based on organometallic derivatives containing sophisticated ligands, which require several upstream synthetic steps for their elaboration.\(^15\) In the search for alternative catalysts that would involve compounds requiring effortless syntheses, we recently demonstrated that the simple borohydrido half sandwich scandium complex \( \text{Cp}^*\text{Sc(BH}_4\text{)}_2(\text{THF}) \) \( 1\text{a} \) (\( \text{Cp}^* = \eta^5\text{C}_5\text{Me}_5, \text{THF} = \text{tetrahydrofuran} \)) is a precatalyst for very active catalytic mixtures towards isoprene cis-polymerisation.\(^16\) Actually, whereas the patents literature describes efficient catalysts based on paramagnetic neodymium developed by industry, many
catalysts reported in the academic literature are made of scandium, or lutetium, both diamagnetic, which is an advantage for organometallic syntheses and characterisations, but these latter metals are among the most expensive rare earths. It is thus desirable to identify analogs containing less costly rare earth elements, notably those which belong to the “early” lanthanides family, like neodymium, for use in the area of the controlled polymerisation of conjugated dienes. Then, the significant development of their catalytic performances has to be undertaken, and preferably in a simple manner.

In order to improve the efficiency of our systems with this plan in mind, we aimed to examine the influence of the direct addition of a neutral ligand towards the polymerisation of isoprene, regarding activity, selectivity, and control of molecular weights. Traditional strategies to switch the regio and the stereoselectivity in conjugated diene polymerisation involve adding alkylaluminum co-catalysts.\(^{17,18}\) An alternative approach, similar to that of Mountford \textit{et al.,}\(^{19}\) and Gade \textit{et al.,}\(^{20}\) and one that is more amenable to rapid catalyst screening, is the use of additional donor ligands. N-Heterocyclic carbenes (NHC’s) are well suited to the chemistry of rare earths\(^{21,22}\) and they may stabilize active and/or selective polymerisation catalysts from the larger rare-earth metals. To our knowledge, the effect of the addition of such ligands has not been assessed in rare earths mediated conjugated dienes polymerisation catalysis yet.

We describe in this work our results in isoprene polymerisation using simple rare earth based borohydrido catalysts, according to two different strategies. The first part is an extension of our previous results obtained with \(\text{Cp}^*\text{Sc}(\text{BH}_4)_2(\text{THF})\) (1a)/borate/Al(n-Bu)\(_3\) combinations, to the neodymium analog \(\text{Cp}^*\text{Nd}(\text{BH}_4)_2(\text{THF})_2\) (1b).\(^{16}\) Since the X-ray structure of this latter compound was unprecedented, we describe it, as well as that of the hexamer isolated in nonpolar solvent, in a preliminary section. The second part shows the effect on isoprene polymerisation of the addition of a NHC molecule to bisborohydrides \(\text{Ln}(\text{BH}_4)_2(\text{THF})_n\) (Ln = Sc, \(n = 1\), 1a; Nd, \(n = 2\), 1b; Sm, \(n = 2\), 1c) and to trisborohydrides \(\text{Ln}(\text{BH}_4)_3(\text{THF})_n\) (Ln = Sc, \(n = 1.5\), 2a; Nd, \(n = 3\), 2b; Sm, \(n = 3\), 2c) as pre-catalysts. Complex (L\(^{2-}\) \(^{\text{dbu}}\))Nd(\(\text{BH}_4\)_2(\text{THF})_3 bearing \(\text{N}^{\text{Bu}}\text{CH}_2\text{CH}_2[1-\text{C}\{\text{N}(\text{CHCH})\text{N}^{\text{Bu}}\}]\) NHC ligand (L\(^{2-}\) \(^{\text{dbu}}\)) was isolated and characterized (Table 1).

\textbf{Table 1.} List of acronyms and abbreviations

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<td>HL(^3)</td>
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Results and discussion

Structural determination of Cp*Nd(BH$_4$)$_2$(THF)$_2$1b

Surprisingly, well-defined half-sandwich complexes of the early 4f-metals are rather scarce. Presumably due to their propensity to easily undergo disproportionation, only a small number among them have been structurally described.\textsuperscript{23,24} Cp$^5$LnX$_2$ (Cp$^R$ = cyclopentadienyl ligand) compounds can eventually be isolated if the X ligand is sufficiently bulky, for example in (C$_5$H$_5$)La(MBMP)(THF)$_3$ (MBMP = 2,2'-methylene-bis(6-tert-butyl-4-methyl-phenolate))\textsuperscript{25} or Cp*CeX$_2$ (X = 2,6-di-tert-butylphenolate, N(SiMe$_3$)$_2$, CH(SiMe$_3$)$_2$),\textsuperscript{26} Cp*LaCH(SiMe$_3$)$_2$,\textsuperscript{27} and more recently in Cp*Ln[AlMe$_4$]$_2$ (Ln = La, Nd).\textsuperscript{28} When they do not undergo disproportionation, half-sandwich complexes often adopt a cluster or oligomeric arrangement.\textsuperscript{29} This was often the case in the borohydride series.\textsuperscript{30} However, mono Cp borohydrido complexes could be isolated as discrete monomers with bulky Cp$_4$i (Cp$_4$i = η$^5$-C$_5$H$_i$Pr$_4$).\textsuperscript{31} In the Cp* series, the scandium compound was the only one to be structurally characterized,\textsuperscript{16} while with larger lanthanides, anionic “ate” [Cp$^8$Nd(BH$_4$)$_3$]$_2$[Mg(THF)$_6$] complexes were also prepared.\textsuperscript{32}

Complex Cp*Nd(BH$_4$)$_2$(THF)$_2$1b was first synthesized by Ephritikhine \textit{et al.} and its molecular structure deduced from $^1$H NMR, IR spectroscopy, and elemental analysis.\textsuperscript{33} We have been able to isolate X-ray quality crystals of this compound by recrystallization from THF–pentane mixtures and we resolved its molecular structure for the first time (Fig. 1). In addition and not surprisingly, we noticed that when this 1b compound is heated in a toluene or a benzene atmosphere, crystals of an unsolvated hexamer [Cp*Nd(BH$_4$)$_2$]$_6$1b’\textsuperscript{‡} are obtained. A similar behaviour had been previously observed in the C$_5$Me$_4$Pr series.\textsuperscript{30} Compound 1b crystallizes as a bis-THF adduct in the $P2_1/n$ space group with four molecules in the unit cell. The BH$_4$ ligands are terminal trihapto, whereas previously reported FTIR spectroscopic data indicated the presence of both tridentate and bidentate BH$_4$ ligands.\textsuperscript{33} It is likely that the compound isolated by Ephritikhine was a mix between the monomer and oligomeric forms of Cp*Nd(BH$_4$)$_2$(THF)$_x$, in which the [Cp*Nd] units are linked through borohydride bridges. Due to the larger ionic radius of neodymium vs. scandium, two THF molecules are necessary to complete the coordination sphere of the metal in the monomeric form, as expected from previous analyses. For comparison, the scandium analog 1a is a mono-THF adduct in the solid-state, with both borohydride groups acting as tridentate ligands, in accordance with short Sc–B bond lengths (2.345 Å).\textsuperscript{16} X-ray crystallographic analysis revealed that both borohydride groups in 1b act as tridentate ligands, in connection with short Nd–B bond distances in the range 2.648–2.663 Å. By comparison, such distances lay in the range 2.596–3.050 Å in [Cp*Nd(BH$_4$)$_3$]$_6$ hexamer 1b’ (see ESI†). The centroid (Cp*)-Nd distance is 2.43 Å as well as in the “ate” complex [Cp*Nd(BH$_4$)$_3$]$^{-32}$ and in the range 2.442–2.502 Å in [Cp*Nd(BH$_4$)$_2$]$_6$1b’ (see ESI†).
Figure 1. Displacement ellipsoid plot of Cp*Nd(BH₄)(THF)₂1b (30%). Only H-atoms on the borohydride ligand are represented. Selected bond lengths (Å) and angles (°): Nd1–B1 = 2.6486(18); Nd1–B2 = 2.6630(18); Nd1–H1C = 2.428; Nd1–H1D = 2.449; Nd1–H1E = 2.458; Nd1–O1 = 2.5460(11); Nd1-centroid = 2.438; B1–Nd1–B2 = 136.59(6).

Polymerisation of isoprene with 1b/borate/Al(iBu)₃

Factors that govern the selectivity in conjugated dienes polymerisation are closely connected to the coordination mode of the monomer to the catalyst (see Fig. 2). Whereas cis-polyisoprene is mainly formed via the coordination of both double bonds of the isoprene monomer to the metal center in η⁴ mode, trans-polyisoprene can result either from η² or η⁴ coordination modes. However, isomerization reaction can occur in between syn and anti allyl species formed after insertion of the monomer, leading to cis-units even in the case of a previous η² coordination of the monomer. A 3,4-polyisoprene results from single coordination of the monomer, which is generally accepted as in line with the steric hindrance around the catalyst metal. Thus, by subtle changes of the environment around the catalyst metal, one may possibly be able to tune the selectivity.
Figure 2. Coordination modes of the isoprene monomer and the stereochemistry of the resulting polyisoprene.

We investigated the polymerisation of isoprene using rare earths borohydride pre-catalysts of three types: half-sandwiches \( \text{Cp}^*\text{Ln} (\text{BH}_4)_2 (\text{THF})_n \), \textit{in situ} modified trisborohydrides \( \text{Ln} (\text{BH}_4)_3 (\text{THF})_n / \text{L} \), and \textit{in situ} modified half-sandwiches \( \text{Cp}^*\text{Ln} (\text{BH}_4)_2 (\text{THF})_n / \text{L} \), \( \text{L} \) being a \( \text{N} \)-heterocyclic carbene. The co-catalysts are of two types, Al trialkyl (\( \text{Al(iBu)}_3 = \text{triisobutylaluminum} \)) combined with a borate activator (\( \text{TB} = [\text{CPh}_3][\text{B}(\text{C}_6\text{F}_5)_4] \) or \( \text{HNB} = [\text{HNMe}_2\text{Ph}][\text{B}(\text{C}_6\text{F}_5)_4] \)), and Mg dialkyl (\( \text{BEM} = ^{10}\text{BuEtMg} ; \text{DBM} = ^{12}\text{Bu}_2\text{Mg} \)).

Selected results are collected in Table 2 (Scheme 1).

Scheme 1. Isoprene polymerisation with \( 1b-2a,b / \text{trityl borate/Al(iBu)}_3 \) ternary systems (Table 2).
Table 2. Isoprene polymerisation with half-sandwich based catalysts combined with borate/alkylaluminum compounds

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<td>30 min</td>
<td>100/68</td>
<td>2.5</td>
<td>34/800</td>
<td>90.0/4.3/5.7</td>
<td></td>
</tr>
<tr>
<td>1.22</td>
<td>2b</td>
<td>4000</td>
<td>2 TB</td>
<td>tol</td>
<td>50</td>
<td>20</td>
<td>8/1.1</td>
<td>nd</td>
<td>nd/—</td>
<td>nd/—</td>
<td></td>
</tr>
<tr>
<td>Ref. 15</td>
<td>2a</td>
<td>420</td>
<td>1 TB</td>
<td>tol</td>
<td>RT</td>
<td>10 min</td>
<td>97.0/166</td>
<td>1.9</td>
<td>27/700</td>
<td>55.2/40.8/4.0</td>
<td></td>
</tr>
<tr>
<td>1.23</td>
<td>2a</td>
<td>420</td>
<td>2 TB</td>
<td>tol</td>
<td>RT</td>
<td>1 min</td>
<td>100/174</td>
<td>3.0</td>
<td>28/600</td>
<td>69.7/27.9/2.4</td>
<td></td>
</tr>
<tr>
<td>1.24</td>
<td>—</td>
<td>1000</td>
<td>2 TB</td>
<td>tol</td>
<td>RT</td>
<td>1 min</td>
<td>44/—</td>
<td>ins</td>
<td>ins/ins</td>
<td>ins/ins</td>
<td></td>
</tr>
</tbody>
</table>

$^a$ Run typically conducted with ca. 5 μmol Ln precatelyst (Ln = Sc, Nd)/borate/10Al(Bu)$_3$, ins: insoluble; nd: not determined. $^b$ TB = [CPh$_3$][B(C$_6$F$_5$)$_3$], HNB = [HNMe$_2$Ph][B(C$_6$F$_5$)$_4$]. $^c$0.5 mL, tol = toluene, DCM = dichloromethane, CLB = chlorobenzene. $^d$ Average activity (kg Pi/mol Ln/h). $^e$ Determined by SEC against
The formation of the neodymium cationic species resulting from the reaction of
hours, but the
hence poorly soluble
observations account f
material. A similar trend was reported with pre
CH
In chlorinated
very low
(HNB = [HNMe
but to the detriment of the
High molecular weights (superior to 10
3.8 when the temperature is raised to 50 °C. At this
PDI (polydispersity index) values are in the range 1.7
below).
This c
could account for the results. Room temperature conditions allow a more controlled process,
In order to get catalyst efficiency at this temperature, the best combination is
1b/2[CPh3][B(C6F5)4]/10Al(′Bu)3, but to generate cis-selective polymerisations, ambient temperature
conditions are needed, which is detrimental to the overall activity. At room temperature, high loadings of
monomer (≥2000 equiv.) are necessary to produce polymer, otherwise the catalyst remains inactive (runs 1.8,
1.9). It has to be mentioned that experiments 1.8 and 1.9 were carried out with vacuum-dried samples of 1b
which had probably formed BH4-bridged clusters of 1b′ type, through loss of coordinated THF molecules (see
below).

Regarding the half-sandwiches, we previously showed that Cp*Nd(BH4)2(THF)2 (1b) with 1 equiv. MgR2 co-
catalyst affords a catalyst for highly trans-selective isoprene polymerisation (>98%). Cp*Sc(BH4)2(THF)
(1a) is inactive under the same conditions, but combined with a borate as the mixture
Cp*Sc(BH4)2(THF)/[CPh3][B(C6F5)4]/Al(′Bu)3 it yields polyisoprene 97% cis-regular with excellent activity.16
Complex 1b also gives cis-rich polymer when associated to TB and triisobutyl aluminum. The best cis-rate
(91.7%, run 1.12) is obtained at high monomer loadings ([monomer]/[catalyst] = 6000) and with
2[CPh3][B(C6F5)4]/10Al(′Bu)3. It is noteworthy that with 1 equiv. trityl borate the resulting catalyst is poorly
active (19% yield in 7 h, run 1.1), at 50 °C, while less than 5 equiv. Al co-catalyst per borate activator affords
rather lower selectivity (run 1.2). In order to get catalyst efficiency at this temperature, the best combination is
1b′/[CPh3][B(C6F5)4]/10Al(′Bu)3 = 6000) and with
1b′/Al(′Bu)3), to result in insoluble
2.5 mL.

Complex 1b also gives cis-rich polymer when associated to TB and triisobutyl aluminum. The best cis-rate
(91.7%, run 1.12) is obtained at high monomer loadings ([monomer]/[catalyst] = 6000) and with
2[CPh3][B(C6F5)4]/10Al(′Bu)3. It is noteworthy that with 1 equiv. trityl borate the resulting catalyst is poorly
active (19% yield in 7 h, run 1.1), at 50 °C, while less than 5 equiv. Al co-catalyst per borate activator affords
rather lower selectivity (run 1.2). In order to get catalyst efficiency at this temperature, the best combination is
1b′/2[CPh3][B(C6F5)4]/10Al(′Bu)3, but to generate cis-selective polymerisations, ambient temperature
conditions are needed, which is detrimental to the overall activity. At room temperature, high loadings of
monomer (≥2000 equiv.) are necessary to produce polymer, otherwise the catalyst remains inactive (runs 1.8,
1.9). It has to be mentioned that experiments 1.8 and 1.9 were carried out with vacuum-dried samples of 1b
which had probably formed BH4-bridged clusters of 1b′ type, through loss of coordinated THF molecules (see
below).

High molecular weights (superior to 10^4 g mol^-1) can be obtained with this catalytic system (runs 1.6, 1.12).
When 1b is synthesized in situ,12 higher cis-rate (80.6 vs. 69.6%, runs 1.4 and 1.3, respectively) is obtained
but to the detriment of the polymer yield (66% vs. 100%). Replacement of the initiating trityl borate by HNB
(HNB = [HNMe2Ph][B(C6F5)4]) gives rise to the formation of a polymer that is 92% cis-selective but with
very low activity (run 1.18). However, this is notable since it is a rare example of cis-polymerisation at
elevated temperature with rare earth borohydrido based catalytic systems.15

In chlorinated solvents, two distinct behaviors are observed: polymerisation takes place within minutes in
CH2Cl2 (runs 1.13–1.14) very similarly as [CPh3][B(C6F5)4] does alone (run 1.24), to result in insoluble
material. A similar trend was reported with pre-catalyst 1a with similar catalytic combinations,16 such
observations account for a likely cationic process, which is known to favor cross-linking with polydienes and
hence poorly soluble polymer. In chlorobenzene (runs 1.15–1.17), good conversions are produced in a few
hours, but the cis-rate is inferior to that obtained in toluene.

The formation of the neodymium cationic species resulting from the reaction of 1b either with
[CPh3][B(C6F5)4] or [HNMe2Ph][B(C6F5)4]16,36 was established by^1H NMR experiments (see Experimental
When one equivalent of $[\text{CPh}_3][\text{B}(\text{C}_6\text{F}_5)_4]$ is added to 1b in THF-$d_8$, formation of a single product is observed with $\delta$(BH$_4$) = 100.3 ppm (initially at 67.9 ppm in 1b) and $\delta$(Cp*) = 8.96 ppm (initially at 7.46 ppm in 1b) likely corresponding to $[\text{Cp*Nd(BH}_4)(\text{THF})_x][\text{B}(\text{C}_6\text{F}_5)_4]$. When the reaction was conducted with $[\text{HNMe}_2\text{Ph}][\text{B}(\text{C}_6\text{F}_5)_4]$ instead of $[\text{CPh}_3][\text{B}(\text{C}_6\text{F}_5)_4]$, slightly different chemical shifts were observed ($\delta$(BH$_4$) = 100.4 ppm and $\delta$(Cp*) = 9.21 ppm) along with the formation of free amine PhNMMe$_2$ (Scheme 2). Attempts to crystallize this cationic species failed due to its high reactivity. When THF was used, the solution became viscous as the species polymerizes the solvent. Other attempts in solvents such as toluene or benzene also failed as the species forms insoluble oil in these solvents.  

Scheme 2. Probable formation of cationic $[\text{Cp*Nd(BH}_4)(\text{THF})_x][\text{B}(\text{C}_6\text{F}_5)_4]$ from 1b.

The catalytic system Nd(BH$_4$)$_3$(THF)$_3$/[CPh$_3$][B(C$_6$F$_5$)$_4$]/Al($'$Bu)$_3$ displays good cis-selectivity (90%) at room temperature, along with satisfactory activity (run 1.21). These data are not far from the ones noticed with the half-neodymocene 1b. We had already established that 2b can behave as an efficient catalyst for isoprene cis-polymerisation: the 2b/[HNMe$_2$Ph][B(C$_6$F$_5$)$_4$]/Al($'$Bu)$_3$ combination yielded more than 90% cis-polyisoprene, but the activity was significantly lower than with trityl borate as activator.  

The comparison of the results presented in Table 2 with those obtained with the related scandium catalyst (i.e. 1a/[CPh$_3$][B(C$_6$F$_5$)$_4$]/Al($'$Bu)$_3$) calls for the following comments: (i) under similar conditions, both activity and cis-selectivity remain inferior with neodymium; (ii) whereas the optimum precatalyst/activator/cocatalyst ratios were 1/2/20 with scandium, it seems that 1/2/10 ratio conditions are more preferable in the case of neodymium; (iii) activities of the same magnitude order (up to ca. 4000 kg PI mol$^{-1}$ catalyst h$^{-1}$) can then be reached with both catalysts, but the scandium one allows a better cis-selectivity. Moreover, it is noteworthy that 2a did not produce cis-PI when combined with [CPh$_3$][B(C$_6$F$_5$)$_4$]/Al($'$Bu)$_3$, whereas 2b does.

### Polymerisation of isoprene in the presence of NHC ligands

We then decided to examine the result of the addition of a neutral ligand to the borohydride derivatives studied previously, towards isoprene polymerisation efficiency. Several NHC ligands were assessed: the
classical \([1-\text{C\{(N'BuCH)\}_2}]\) (L\(^1\)) and functional N-heterocyclic carbenes, two amino-tethered HN'BuCH\(_2\)CH\(_2\)[1-C{N(CHCH)NR}] (HL\(^2\)-R) (R = \(^t\)Bu, Mes (Mes = 2,4,6-Me\(_3\)-C\(_6\)H\(_2\)))\(^{38}\) and the hydroxyl-tethered HOCMe\(_2\)CH\(_2\)[1-C{N(CHCH)NiPr}] (HL\(^1\), also represented under a zwitterionic form)\(^{39}\) (Scheme 3).

Such molecules were chosen as potentially capable to coordinate either to the rare centre or to the magnesium atom, due to the carbene moiety.\(^{21}\) In addition, HL\(^2\)-R and HL\(^3\) are prone to afford anionic ligation to the metal via deprotonation. Selected results are gathered in Table 3.

**Scheme 3.** NHC-ligands assessed as additional ligands in isoprene polymerisation.

Polymerisation experiments were conducted as described in the previous section, *i.e.* by combining the precatalyst with borate/Al\((t\)Bu\)_3 cocatalyst, but in the presence of the carbene ligand (Scheme 4). When the latter was bearing a tethered neutral protio-NHR function, the necessary amount of BEM \("\text{BuEtMg, or DBM = Mg}^{t}\text{Bu}_2\) for deprotonation was added, similarly as conducted in the case of the “in situ B/A route” (run 1.4, Table 2), since it was clear from \(^1\)H NMR spectroscopic analysis that 2b did not react with HL\(^2\)-tBu via protonolysis to form a putative (L\(^2\)-tBu)Nd(BH\(_4\)_2)(THF)\(_2\) (see Experimental section). From the results gathered in Table 3, one can observe that HL\(^2\)-R ligands behave differently depending on the R group, and on the metal considered. While 2b affords 100% yield of polymer in 30 min at room temperature (run 1.21, Table 2), the addition of a NHC ligand has a strong impact in terms of activity: the conversion needs 24 h to be completed in the presence of 1 equiv. L\(^1\), and 20 h with L\(^2\)-tBu (runs 2.1 and 2.2, respectively). This result is in line with a previous observation made by some of us: the carbene adduct of a samarium allyl initiator \(\text{C}_5\text{H}_4^{t}\text{Bu})_2\text{Sm(C}_3\text{H}_5)\text{[C(NPr)_2(CMe)\_2]}\) was poorly active for isoprene polymerisation.\(^{40}\) Interestingly however, the selectivity is switched with L\(^2\)-tBu from 90/4 to 52/46% cis/trans (run 2.2). When L\(^2\)-Mes is used, the reaction is even slower, but the cis-selectivity is preserved (run 2.3). The impact of a carbeneligand was less negative when added to pre-catalyst 2a. Whereas the system is not selective with 2a/TB/Al\((t\)Bu\)_3,\(^{16}\) the cis-selectivity was more pronounced with 2 equiv. trityl borate (run 1.23), and this was maintained in the presence of L\(^1\) carbene, though with lower activity (run 2.4). Higher cis-rate (79%) was received with 2.6 equiv. borate
activator (run 2.5). With 1 equiv. L\textsubscript{2-tBu}, no effect of the NHC was observed; both activity and selectivity being similar to the reference experiment (run 2.6, but 2 TB). On the other hand, a spectacular result was obtained when 1 equiv. HL\textsubscript{2-Mes} was added to the catalytic mixture: the selectivity switched to \textit{trans}-selective, along with a significant decrease of the reaction rate (run 2.7), which strongly differs with that taking place in the case of neodymium (run 2.3). This trend was confirmed with run 2.8: 2 equiv. L\textsubscript{2-tBu} were added to \(2a/\text{TB/Al(}^{t}\text{Bu)}\rangle\), which led to the formation of highly \textit{trans}-polyisoprene, accompanied with total conversion. This \textit{trans}-stereoregularity is tentatively ascribed to the steric hindrance around the scandium atom, due to the likely coordination of the bulky functionalized NHC ligands. An increase of the \textit{trans}-1,4 content in butadiene polymerisation with Nd catalysts in the presence of an excess of DIBAH was similarly ascribed to impediment of \(\eta^4\)-coordination of the monomer with benefit to \(\eta^2\)-coordination.\textsuperscript{41} To our knowledge, the result of run 2.8 is a rare example of \textit{trans}-polyisoprene obtained with a scandium cationic based catalyst, since steric congestion is generally known to favour \(\eta^2\) coordination of the monomer, and thus 3,4-polymerisation (Fig. 2).\textsuperscript{42} Regarding the half-sandwiches, a polymerisation test was conducted with 1b in the presence of L\textsubscript{1-carbene} (1 equiv.) at a monomer/precatalyst ratio of 500, which resulted in the production of polymer, with very high \textit{cis}-regularity (run 2.9). This result is in contrast with the one in the absence of L\textsubscript{1-carbene} which gave no polymer (run 1.8). One can advance here that the effect of L\textsubscript{1-carbene} is parallel to an increase of monomer/precatalyst ratio (run 1.10).\textsuperscript{43} Furthermore, one can also propose that the carbene moiety would help to dissociate the cluster structure – possibly of 1b′ type – of 1b after storage, since dry batches of precatalyst 1b are used. The addition of one equivalent of HL\textsubscript{2-Mes} to 1b switches the selectivity from highly 1,4-\textit{cis} in the presence of L\textsubscript{1-carbene} (1 equiv.) to 1,4-\textit{trans} at 69\% (run 2.10). This can be attributed to the bulk of the HL\textsubscript{2-Mes} ligand compared to L\textsubscript{1-carbene} resulting in \(\eta^2\) coordination mode of the monomer to the scandium atom. With scandium compound 1a, the behavior is different: in the presence of L\textsubscript{1-carbene} (1 equiv.) we observed, along with high activity (100\% conversion in 3 min, run 2.11), a switch in selectivity, this time oriented towards 3,4-polymerisation of isoprene. A stereoselective polymerisation of isoprene with NHC-tethered fluorenyl and indenyl supported catalysts was already reported to yield 3,4-polyisoprene, but the stereo-inductive effect of the NHC moiety was not established.\textsuperscript{44,45} Adding an additional equivalent L\textsubscript{1-carbene} led to a lower activity but similar 3,4-preference (run 2.12). When the half-sandwich 1a is associated to HL\textsubscript{2-Mes} the activity is even lower but the selectivity remains unchanged (run 2.13).

\[\text{Scheme 4. Isoprene polymerisation with } 1a,b–2a,b/L_1 \text{ or HL}_2^2 \text{ (MgR}_2)_2/\text{borate/Al(}^{t}\text{Bu)}\rangle \text{ systems (Table 3).}\]
Table 3. Isoprene polymerisation with borohydride precatalysts combined with borate/alkylaluminum compounds in the presence of NHC ligands

<table>
<thead>
<tr>
<th>Run</th>
<th>Precatalyst/NHC</th>
<th>t (min)</th>
<th>Yield (%)</th>
<th>$M_n$ (g mol$^{-1}$)</th>
<th>PDI</th>
<th>$M_n$ (g mol$^{-1}$)</th>
<th>cis/trans/3,4 (%)</th>
</tr>
</thead>
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<td>1.21</td>
<td>2a–––</td>
<td>30</td>
<td>100</td>
<td>24 700</td>
<td>2.5</td>
<td>34 000</td>
<td>90.0/4.3/5.7</td>
</tr>
<tr>
<td>2.1</td>
<td>2b/L $^t$</td>
<td>1440</td>
<td>100</td>
<td>14 400$^t$</td>
<td>1.4</td>
<td>34 000</td>
<td>83.2/13.0/3.8</td>
</tr>
<tr>
<td>2.2</td>
<td>2b/0.5DBM/HL$^{2-3} $</td>
<td>1200</td>
<td>100</td>
<td>35 600$^t$</td>
<td>2.7</td>
<td>34 000</td>
<td>52.1/46.0/1.9</td>
</tr>
<tr>
<td>2.3</td>
<td>2b/0.5DBM/HL$^{3} $</td>
<td>960</td>
<td>15</td>
<td>–</td>
<td>–</td>
<td>5100</td>
<td>92.0/6.0/2.0</td>
</tr>
<tr>
<td>Ref. 16</td>
<td>2a</td>
<td>10</td>
<td>97.0</td>
<td>15 500</td>
<td>1.9</td>
<td>27 700</td>
<td>55.2/40.8/4.0</td>
</tr>
<tr>
<td>1.23$^f$</td>
<td>2a</td>
<td>1</td>
<td>100</td>
<td>42 500</td>
<td>3.0</td>
<td>28 600</td>
<td>69.7/27.9/2.4</td>
</tr>
<tr>
<td>2.4</td>
<td>2a/L $^t$</td>
<td>5</td>
<td>100</td>
<td>21 500</td>
<td>2.9</td>
<td>34 000</td>
<td>66.7/29.9/4.1</td>
</tr>
<tr>
<td>2.5</td>
<td>2a/L $^t$</td>
<td>5</td>
<td>100</td>
<td>24 300</td>
<td>4.0</td>
<td>34 000</td>
<td>79.7/16.2/3.4</td>
</tr>
<tr>
<td>2.6</td>
<td>2a/0.5DBM/HL$^{2-3} $</td>
<td>10</td>
<td>87.5</td>
<td>ins</td>
<td>ins</td>
<td>29 700</td>
<td>56.8/38.7/4.5</td>
</tr>
<tr>
<td>2.7</td>
<td>2a/0.5DBM/HL$^{3} $</td>
<td>960</td>
<td>48.3</td>
<td>16 200$^t$</td>
<td>2.0</td>
<td>16 400</td>
<td>9.4/88.6/2.0</td>
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<tr>
<td>2.8</td>
<td>2a/1DBM/2 HL$^{2-3} $</td>
<td>1440</td>
<td>100</td>
<td>6800</td>
<td>2.2</td>
<td>34 000</td>
<td>4.2/94.1/1.7</td>
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<td>2.9</td>
<td>1b/L $^t$</td>
<td>1440</td>
<td>41</td>
<td>6800</td>
<td>1.6</td>
<td>13 900</td>
<td>92.5/3.5/4.0</td>
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<tr>
<td>2.10</td>
<td>1b/0.5BEM/HL$^{3} $</td>
<td>3900</td>
<td>100</td>
<td>17 100</td>
<td>1.9</td>
<td>34 000</td>
<td>29.3/68.9/1.8</td>
</tr>
<tr>
<td>2.11</td>
<td>1a/L $^t$</td>
<td>3</td>
<td>100</td>
<td>13 900</td>
<td>5.1</td>
<td>34 000</td>
<td>41.1/3.1/55.8</td>
</tr>
<tr>
<td>2.12</td>
<td>1a/2L $^t$</td>
<td>1440</td>
<td>100</td>
<td>15 400</td>
<td>5.3</td>
<td>34 000</td>
<td>39.7/3.3/57.0</td>
</tr>
<tr>
<td>2.13</td>
<td>1a/0.5BEM/HL$^{3} $</td>
<td>3900</td>
<td>100</td>
<td>Multimodal</td>
<td>–</td>
<td>34 000</td>
<td>48(1,4)/52(3,4)</td>
</tr>
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</table>

$^a$ [I]/[Ln] = 500; [TB]/[Ln] = 2; [Al]/[Ln] = 10, room temperature, V(toluene) = 0.5 mL; ins: insoluble; nd: not determined. $^b$ Determined by SEC against polystyrene standards. $^c$ $M_{n,th}$ = [I]/[Ln] × yield × 68. $^d$ From $^1$H and $^{13}$C NMR. $^e$ Bimodal: a minor contribution is observed at high $M_n$ values. $^f$ Partly soluble. $^g$ [I]/[Ln] = 420; [TB]/[Ln] = 1. $^h$ [I]/[Ln] = 420; [TB]/[Ln] = 2. $^i$ [TB]/[Ln] = 2.6.

Polymerisation experiments were further conducted with pre-catalysts 1b, Cp$^*$Sm(BH$_4$)$_2$(THF)$_2$ (1c), 2b and Sm(BH$_4$)$_3$(THF)$_3$ (2c), in the presence of NHC ligands, and in combination with BEM as co-catalyst (Scheme 5). Selected results are given in Table 4. As before, with amino-tethered $N$-heterocyclic carbenes HL$^{2-R}$, an additional equimolar amount of alkyl reagent was added to ensure the deprotonation of the ligand (see Experimental). This was not however necessary when the carbene ligand was bearing a tethered protio–OH group (HL$^3$), due to higher acidity of the latter.

Scheme 5. Isoprene polymerisation with 1b,c–2b,c/L$^1$, HL$^2$ or HL$^3$/MgR$_2$ systems (Table 4).
Table 4. Isoprene polymerisation with borohydride precatalysts combined with BEM in the presence of NHC ligands

<table>
<thead>
<tr>
<th>Run</th>
<th>Pre-catalyst/NHC</th>
<th>[Mg]/[Nd]</th>
<th>t (h)</th>
<th>Yield (%)</th>
<th>$M_{n,exp}$ (g mol$^{-1}$)</th>
<th>PDI$^c$</th>
<th>$M_{n,th}$ (g mol$^{-1}$)</th>
<th>cis/trans/3,4$^e$ (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ref. 42</td>
<td>2b/---</td>
<td>1</td>
<td>2</td>
<td>87.0</td>
<td>52 100</td>
<td>1.3</td>
<td>58 100</td>
<td>---/95.5/---</td>
</tr>
<tr>
<td>3.1</td>
<td>2b/1 L$^1$</td>
<td>1</td>
<td>2</td>
<td>30.9</td>
<td>27 000</td>
<td>1.2</td>
<td>21 000</td>
<td>1.5/96.5/2.0</td>
</tr>
<tr>
<td>3.2</td>
<td>2b/1 HL$^2$</td>
<td>1</td>
<td>2</td>
<td>26.5</td>
<td>34 600</td>
<td>1.3</td>
<td>18 000</td>
<td>1.6/96.1/2.3</td>
</tr>
<tr>
<td>3.3</td>
<td>2b/1 HL$^3$</td>
<td>1</td>
<td>18</td>
<td>80.2</td>
<td>39 800</td>
<td>1.2</td>
<td>54 500</td>
<td>1.0/96.6/2.4</td>
</tr>
<tr>
<td>3.4</td>
<td>2b/1 (H$_2$L$^7$)$^7b$</td>
<td>1</td>
<td>16</td>
<td>25.0</td>
<td>6 200</td>
<td>1.7</td>
<td>8500</td>
<td>2.0/94.0/3.0</td>
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<td>2b/2 HL$^{2b}$</td>
<td>1</td>
<td>16</td>
<td>50.2</td>
<td>9 800</td>
<td>1.6</td>
<td>17 100</td>
<td>1.9/96.1/2.0</td>
</tr>
<tr>
<td>3.6</td>
<td>2c/1 HL$^3$</td>
<td>1</td>
<td>96</td>
<td>18.6</td>
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<td>2</td>
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<td>27 000</td>
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<td>50 100/17 100</td>
<td>1.1/1.2</td>
<td>35 100</td>
<td>0.4/97.3/2.3</td>
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\(a\) \(T = 50^\circ\mathrm{C}\); \(V(\text{toluene}) = V(\text{isoprene})\); [I]/[Nd] = 1000. \(b\) [I]/[Nd] = 500. \(c\) Determined by SEC against polystyrene standards and corrected by a factor of 0.5 in case of \(\text{trans-PI}^\text{13}d\) \(M_{n,th} = [\text{I}]/[\text{Nd}] \times \text{yield} \times 68.\) \(e\) From \(^1\text{H}\) and \(^13\text{C}\) NMR.

At the addition of NHC ligands, one can observe that the \(\text{trans}\)-character of the reference catalysts Ln(BH$_4$)$_3$(THF)/BEM (95.5%)\(^{46}\) is preserved, and in general little improved. On the other hand, the activity is severely lowered (runs 3.1, 3.2, 3.3). It was established by a series of NMR experiments that the functionalized NHC ligands act as chelates, the carbene moiety being coordinated to the Nd atom in solution (see Experimental). The result of the reaction of HL$^3$ with 2b affords a set of paramagnetic resonances in the NMR spectra that can be assigned to an L$^1$-supported Nd compound (Scheme 6, top). Despite our efforts, it was not possible to isolate this compound in bulk conditions. The reaction of 2b with the protio imidazolium precursor (H$_2$L$^7$)$^{7b}$, \(\text{I}\) gave (with intense bubbling) a compound in which the \(^1\text{H}\) NMR paramagnetic spectrum was very similar, thus confirming the previous reaction. Run 3.4, which was carried out with the protio ligand, gives as expected a highly \(\text{trans}\)-regular polymer, though in little quantity. The same tendency was noticed in the presence of 2 equiv. HL$^3$, \(\text{i.e.}\) \(\text{trans}\)-selectivity and lesser activity (run 3.5). One can note however that in the case of samarium, the addition of 1 equiv. HL$^3$ yields some polymer, with fair \(\text{trans}\)-selectivity (run 3.6), whereas 2c/BEM was inactive in the absence of carbene ligand.\(^{46}\) With amino-tethered \(N\)-heterocyclic carbenes HL$^{2R}$, whatever the substituent \(\text{tert}\)-butyl or mesityl, the activity is low, and the selectivity remains \(1,4\)-\(\text{trans}\) (runs 3.7–3.9). The reaction of HL$^{2R}$ with 2b after deprotonation with DBM and the coordination of the carbene moiety to the Nd atom are clearly established by \(^1\text{H}\) NMR (Scheme 6, bottom, and Experimental part). The scale up synthesis of the reaction of HL$^{2b}$/BEM with 2b in the presence of BEM led to the isolation of \(\ldots\)
the bis THF adduct (L^{2-\text{Bu}})_2Nd(BH_4)_2(THF)_23 as a crude product (43% yield). After work-up and prolonged drying, a non-solvated compound (L^{2-\text{Bu}})Nd(BH_4)_23', according to elemental analysis, was finally obtained as a blue-red solid (see Experimental section). Unfortunately no suitable crystals were obtained for X-ray analysis. This compound which initially bears two THF molecules per neodymium when freshly prepared, has a tendency to desolvate as the unsolvated form is obtained when the complex is further dried.\textsuperscript{30}

Scheme 6. Reaction of 2b with HL^3 (top) and with HL^{2-\text{Bu}} in the presence of dialkylmagnesium (bottom).

In contrast with homoleptic trisborohydrides, half-sandwich precatalysts remain quite active when NHC molecules are present (conversion >60% in 2 h, runs 3.10 and 3.11). Whatever L\textsuperscript{1} or HL\textsuperscript{3}, the selectivity remains highly trans-, in the range observed with the reference 1b/BEM catalyst.\textsuperscript{14} The same is also noticed with samarium, although the process is less controlled, with bimodal SEC curves (runs 3.12, 3.13). These results are somewhat different from those reported by Anwander et al.: the presence of donor nitrogen functionality in quinolyl-substituted cyclopentadienyl yttrium and lanthanum catalysts was shown to slightly disfavor the trans-selectivity towards isoprene polymerisation.\textsuperscript{9}

Finally, \textsuperscript{1}H and \textsuperscript{11}B NMR monitoring experiments were carried out with simple L\textsuperscript{1} carbene in order to check just the coordination to Nd (1b) and to Sc (2a) (see Fig. 3 and Experimental part). The signals of the initial compound in both cases disappear, in favour of new sets of signals. With paramagnetic Nd, one new Cp* and
one new BH₄(¹H and ¹¹B) signals can be observed, along with some diamagnetic THF, and one may clearly
distinguish between paramagnetic and diamagnetic peaks for the L¹ ligand. Thus, some L¹ carbene is probably
coordinated under a single new species. With scandium, the initial compound has reacted as well, but several
¹¹B resonances can be observed, and many peaks are assignable to coordinated L¹ ligand in the ¹H NMR
spectrum. This would possibly indicate a constrained molecular geometry which could be tentatively
rationalized as in line with the surprising switch of selectivity from cis- to trans- with pre-catalyst 2a (runs
2.6–2.8, 2.10, Table 3). To summarize, the coordination of the L¹ carbene is established unambiguously, but
the result may probably be interpreted in terms of dynamic exchange, which unfortunately does not allow
isolation of well-defined adducts in this borohydride series.

Figure 3. ¹H (left) and ¹¹B (right) NMR signals of borohydride groups (benzene-d₆). Grey curves: 1b alone;
black curves: 1b + 1 equivalent of L¹ NHC ligand.

The situation is obviously more complicated in polymerisation experiments since several metallic species –
the pre-catalyst and the co-catalyst – are present in the mixture. The carbene moiety may coordinate either to
the Nd or the Mg atom (Table 4), Nd or Al, Sc or Al (Table 3), and even Nd/Sc, Mg or Al, when all three
metals are present (Table 3). This depends on the type of polymerisation, and thorough investigations which
require the use of NHC ligands that are specific to a metal, are necessary to go further in the understanding of
these phenomena. This will be the subject of our further studies in this area.

Conclusion

We reported in a previous work the efficiency of the ternary scandium-based catalytic system
Cp*Sc(BH₄)₂(THF)/[CPh₃][B(C₆F₅)₄]/Al(iBu)₃ towards the polymerisation of isoprene, leading to highly cis-
polyisoprene along with a strong activity. These results were extended to neodymium which is less costly than
scandium metal and more utilized as a catalyst in industry. The already reported complex
Cp*Nd(BH₄)₂(THF)₂1b used in this study was characterized by X-ray analysis, displaying a monomeric
structure including two tridentate borohydride ligands and two THF molecules. When activated in the same
conditions as previously used with the scandium analog, *i.e.* with a borate activator and an aluminum alkyl, this neodymium compound also leads to highly active and *cis*-selective (up to 92%) isoprene polymerisation catalysts. The likely formation of the neodymium cationic species resulting from the reaction of Cp*Nd(BH₄)₂(THF)₂ with the borate activators was established by ¹H NMR experiments. In the second part of this work, scandium, neodymium and samarium half-metallocenes, as well as their precursors Ln(BH₄)₃(THF)ₓ, were associated *in situ* to different types of *N*-heterocyclic carbene ligands, with the aim to modify the polymerisation active species and hence the polymerisation catalysis. The amidocarbene (L²⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻˓
BF₃, Et₂O. The chemical shifts were calibrated by using the residual resonances of the solvent. Elemental analyses were carried out with a Fisons EA 1108 CHON by the Centre d'Analyses de l'Université de Dijon and at the Elemental Analysis Service of London Metropolitan University (UK). Size exclusion chromatography of polymer samples was performed in THF as an eluent at 40 °C (1 mL min⁻¹) with a Waters SIS HPLC pump, a Waters 410 refractometer, and Waters Styragel columns (HR2, HR3, HR4, and HR5E) calibrated with polystyrene standards. A correction factor of 0.5 was applied for the determination of number-average molecular weight of poly(1,4-trans isoprene).⁴⁹

X-ray structure determination of Cp*Nd(BH₄)₂(THF)₂ (1b)

Bright blue X-ray quality crystals were grown from THF/pentane of freshly prepared 1b according to the literature.³³ X-ray analysis of the crystals displayed compound Cp*Nd(BH₄)₂(THF)₂. X-ray data: compound 1b (C₁₃H₁₉B₂NdO₂, Mᵣ = 453.35) crystallizes in the monoclinic space group P2₁/n with a = 8.6110(10), b = 17.3480(3), c = 14.9131(3) Å, β = 105.060(1), V = 2151.3(3) Å³, and ρ = 1.400 g cm⁻³ for Z = 4. Data were collected at 100(1) K on a Bruker Smart Apex CCD 4K system. The structure was solved by charge flipping methods using Superflip software,⁵⁰ and least-square refined with Olex2 software.⁵¹ The model based on 9533 reflections (I > 2.0σ(I), Rᵣᵣ = 0.026), total number of reflections = 59 452, converged to a final R₁ = 2.5% (wR₁ = 5.6%). CCDC 892290.

X-ray structure determination of [Cp*Nd(BH₄)₂]₆ (1b’)

Bright blue X-ray quality crystals were grown from saturated benzene solution of freshly prepared 1b according to the literature.³³ X-ray analysis of the crystals displayed the unsolvated hexamer [Cp*Nd(BH₄)₂]₆. X-ray data: compound 1b’ (C₆₀H₅₀B₁₂Nd₆·2(C₆H₆)), Mᵣ = 1962.8 g mol⁻¹) crystallizes in the triclinic space group P̅₁ with a = 15.979(4), b = 16.981(4), c = 21.7026(6) Å, α = 90.181(5)°, β = 109.502(5)°, γ = 119.054(5)°, V = 4586(2) Å³, and ρ = 1.421 g cm⁻³ for Z = 2. Data were collected at 100(1) K on a Bruker Smart CCD 1K system. All tested crystals were systematically twinned (180° around real 1 0 0 axis); thus the study was undertook using such a sample where twin ratio refined afterward to 49.6%. The structure was solved by charge flipping methods using Superflip software,⁵⁰ and least-square refined with Jana2000 software⁵² including Rigid Body approach. The model based on 9735 reflections (I > 3.0σ(I), Rᵣᵣ = 0.065), total number of reflections = 20 508, converged to a final R₁ = 5.4% (wR₁ = 5.6%). CCDC 892289.

Cp*Nd(BH₄)₂(THF)₂,1b from [Cp*Nd(BH₄)₂]₆

Complex [Cp*Nd(BH₄)₂]₆ was stirred overnight in THF. The blue solution was evaporated to dryness without any further drying under vacuum.¹¹H NMR in C₆D₆ of the compound after treatment was consistent with Cp*(BH₄)₂(THF)₂ showing that the clustering is reversible.¹¹H NMR (C₆D₆, 293 K, δ in ppm): 93.65 (br, 8H, BH₂); 8.66 (br, 15H, Me-Cp*), −1.95 (br, 8H, THF); −5.32 (br, 8H, THF).¹¹B NMR (C₆D₆, 298 K, calibrated against BF₃·Et₂O, δ in ppm): 174.3 (br, BH₄).
**1H NMR experiments**

**Nd(BH₄)₃(THF)₂b with one equivalent of HL.** At the addition of 1 equiv. HL to 2b (6.7 mg, 16.4 μmol) in C₆D₆ in an NMR tube equipped with a Teflon valve (Young™) an immediate bubbling was observed along with formation of a partially soluble compound. **¹H NMR analysis recorded immediately showed resonances of a paramagnetic compound fully compatible with the formula (BH₄)₂NdOCMe₂CH₂[1-C{N(CHCH)NPr}].** ¹H NMR (C₆D₆, 293 K, δ in ppm): 41.3 (br, 2H, CH₂), 39.1 (br, 6H, CMe₂), ca. 35 ppm (elapsed in the baseline, 4H, BH₄), 8.6 (br, 1H, C–H), −8.4 (s, 6H, ¹Pr), −10.4 (br, 1H, ¹Pr). A tentative synthesis carried out at bulk scale in THF failed to obtain the desired product.

**Nd(BH₄)₃(THF)₂b with one equivalent of [H₂L]⁺[I]⁻.** The reaction of 2b (8.1 mg, 20 μmol) with 1 equiv. [H₂L]⁺[I]⁻ (6.2 mg) in C₆D₆–THF mixture in an NMR Young tube was very bubbly, giving a major set of signals similar to the above. ¹H NMR (C₆D₆, 293 K, δ in ppm): 60 (vbr, 8H, BH₄), 29.8 (vbr, 2H, CH₂), 14.6 (s, 9H, ¹Bu), 11.1 (s, 2H, CH₂), 6.7 (s, 1H, C–H), 3.6 (s, 9H, ¹Bu), 2.8 (br, THF), 1.48 (br, THF) which could be compatible with the formula (BH₄)₂NdOCMe₂CH₂[1-C{N(CHCH)NPr}].

**Nd(BH₄)₃(THF)₂b with one equivalent of HL²⁻Bu/0.5 DBM.** No gas evolution was observed on the addition of 1 equiv. HL²⁻Bu (LiBr adduct, 4.8 mg) to 2b (5 mg, 12.3 μmol) in C₆D₆, the poorly soluble HL²⁻Bu remaining unchanged at the bottom of the tube. Addition of 0.5 equiv. DBM afforded a clean spectrum with paramagnetic resonances. ¹H NMR (C₆D₆, 293 K, δ in ppm): 68 (vbr, 8H, BH₄), 30.2 (vbr, 2H, CH₂), 14.6 (s, 9H, ¹Bu), 12.0 (s, 2H, CH₂), 6.26 (s, 1H, C–H), 4.03 (s, 9H, ¹Bu), 2.59 (br, 8H, THF), 1.71 (br, 8H, THF). The scale up syntheses of (L²⁻Bu)Nd(BH₄)₂(THF)₂ and (L²⁻Bu)Nd(BH₄)₂3’ are described further down in the Experimental part.

**Cp*Nd(BH₄)₃(THF)₂b with one equivalent of L.** 10 mg of Cp*Nd(BH₄)₃(THF)₂b (2.2 × 10⁻⁵ mol) were mixed with 4 mg of L (2.2 × 10⁻⁵ mol) in C₆D₆ in an NMR tube equipped with a young valve. ¹H NMR (C₆D₆, 293 K, δ in ppm): 82.93 (br, 8H, BH₄); 8.41 (br, 15H, Me-Cp*), 6.80 (br, 2H, free H-L¹); 1.51 (br, 18H, free tBu-L¹), −0.67 (br, 4H, THF); −1.74 (br, 4H, THF); additional peaks (attributable to coordinated-L¹): 13.02, 11.02, 9.99, 7.97, 4.69, 0.85, −1.41, −4.85. Some free THF is present in the spectrum. ¹¹B NMR (C₆D₆, calibrated against BF₃, Et₂O): 162 (br, BH₄).

**Sc(BH₄)₃(THF)₁₂a with one equivalent of L.** 10 mg of Sc(BH₄)₃(THF)₁₂a (5 × 10⁻⁵ mol) were mixed with 9 mg of L (5 × 10⁻⁵ mol) in C₆D₆ in an NMR tube equipped with a young valve. ¹H NMR (C₆D₆, 293 K, δ in ppm): 6.80 (s, 2H, free H-L¹); 6.45 (s, 2H, coordinated H-L¹); 3.64 (m, 4H, THF); 1.59 (s, 18H,
coordinated ‘Bu-L’), 1.51 (s, 18H, free ‘Bu-L’), 1.41 (m, 4H, THF); 2.7–0.5 (br, 8H, BH₄); peaks which could be attributed to coordinated-L¹ ligand: 13.02, 11.02, 9.99, 7.97, 4.69, 0.85, −1.41, −4.85. ¹¹B NMR (C₆D₆, calibrated against BF₃·Et₂O, 293 K, δ in ppm): −21.35 and −31.89 (br, BH₄). Non-attributed peaks (all minor except the last one): 8.09, 7.47, 7.35, 1.22, 0.90. Initial data for 2a (2 THF). ¹H NMR (C₆D₆, 293 K, δ in ppm): 3.89 (m, 8H, THF); 1.14 (m, 8H, THF); 2–0.25 (br, 12H, BH₄); ¹¹B NMR (C₆D₆, calibrated against BF₃, Et₂O, 293 K, δ in ppm): −18.79 (br, BH₄).

NMR scale synthesis of [Cp*Sm(BH₄)(THF)ₓ][B(C₆F₃)₄]. In an NMR tube equipped with a Teflon valve 3 mg of Cp*Nd(BH₄)ₓ(THF)ₓ 1b (6.61 × 10⁻⁶ mol) and 6.1 mg of [CPh₃][B(C₆F₃)₄] (6.61 × 10⁻⁶ mol) were mixed in THF-d₈. The solution instantly turns from pale blue to pale yellow. Formation of a single product with δBH₄ = 100.3 ppm (δ BH₄ in Cp*Nd(BH₄)ₓ(THF)ₓ initially at 67.9 ppm) and δ Cp* = 8.96 ppm (δ Cp* in Cp*Nd(BH₄)(THF)ₓ, initially at 7.46 ppm) corresponding to [Cp*Nd(BH₄)(THF)ₓ][B(C₆F₃)₄]. Deuterated THF was polymerized after standing one week at room temperature. When the reaction was conducted with [HNMe₂Ph][B(C₆F₃)₄] instead of [CPh₃][B(C₆F₃)₄], slightly different chemical shifts was observed (δ BH₄ = 100.4 ppm and δ Cp* = 9.21 ppm) along with the formation of free amine PhNMe₂ (δ = 7.83; 7.44; 7.32; 3.12 ppm).

Synthesis of (L²-dBu)Nd(BH₄)₂(THF)₂ (3)

Nd(BH₄)₂(THF)₃ (442 mg, 1.09 × 10⁻³ mol) and HL²-dBu (LiBr adduct, 338 mg, 1.09 × 10⁻³ mol) were weighed in a double-neck flask in a glove box. The vessel was connected to the vacuum line and 20 mL of toluene were distilled into the flask. BEM (0.5 equiv., 300 mg of a 20% hexane solution) diluted in toluene (10 mL) was added dropwise at room temperature to the mixture of 2b and HL²-dBu. Immediately a deep blue–green solution was formed, along with a white precipitate. The mixture was allowed to stir overnight at room temperature and was filtered and concentrated to ca. 10 mL. No crystals were obtained after 24 h at room temperature. Pentane(10 mL) was added by distillation, leading to precipitation of a light blue powder which expanded on drying, after isolation of this blue powder from the mother liquor. 0.248 g (43% yield) of (L²-dBu)Nd(BH₄)₂(THF)₂ 3 was finally obtained as a blue–red dichroic solid. ¹H NMR (C₆D₆, 293 K, δ in ppm): 28.8 (vbr, 2H, CH₂), 15.2 (s, 9H, 'Bu), 11.96 (s, 2H, CH₂), 6.1 (s, 1H, C–H), 3.87 (s, 9H, 'Bu), 2.73 (br, 8H, THF), 1.44 (br, 8H, THF), one CH signal from the NHC carbene and BH₄ signals could not be observed. After prolonged drying, unsolvated complex (L²-dBu)Nd(BH₄)₂ 3 was obtained: elemental analysis calculated (%): (NdC₆H₁₂B₂N₂); 396.28 g mol⁻¹): C 39.40, H 8.14, N 10.60; found: C 39.29, H 8.19, N 10.55. Recrystallization attempts in THF/pentane at various temperatures (room temperature, freezer) failed to afford suitable crystals for X-ray studies.

Synthesis of Cp*Sm(BH₄)₂(THF)₂ (1c)

[Cp*Sm(BH₄)₂]ₓ has already been described as an unsolvated compound.⁴⁸ Sm(BH₄)₂(THF)₃ (1.3 g, 3.16 × 10⁻³ mol) and KCp* (0.54 g, 3.10 × 10⁻³ mol) were weighed in a vessel in a glove box. The vessel was
connected to the vacuum line and 75 mL of THF were distilled into the flask. After overnight stirring at room temperature, an orange solution was obtained with white salts. The solution was filtered and THF was removed under vacuum. The red solid was then washed with 30 mL of pentane and dried under dynamic vacuum at room temperature for 2 h. 1.314 g (92% yield) of Cp*Sm(BH$_4$)$_2$(THF)$_2$ (according to $^1$H NMR) were obtained as a red solid. $^1$H NMR (C$_6$D$_6$, 293 K, δ in ppm): −12.1 (broad, 8H, BH$_4$); 0.61 (broad, 8H, THF); 0.83 (s, 15H, Cp*); 2.02 (broad, 8H, THF).

1.314 g (92% yield) of Cp*Sm(BH$_4$)$_2$(THF)$_2$ were obtained as a red solid.

Typical isoprene polymerisation experiment with 1b/[CPh$_3$][B(C$_6$F$_5$)$_4$]/Al(iBu)$_3$ (run 1.12, Table 2)

In a glove box under argon (H$_2$O and O$_2$ <2 ppm), 2.4 mg of complex Cp*Nd(BH$_4$)$_2$(THF)$_2$ 1b (5.3 × 10$^{-6}$ mol) are dissolved in 0.5 mL of dry and degassed toluene. Two equivalents of [CPh$_3$][B(C$_6$F$_5$)$_4$] (9.5 mg, 1.06 × 10$^{-3}$ mol) were then added under stirring and after several minutes an orange solution was obtained. 10 equivalents of Al(iBu)$_3$ ($d = 0.786, 14 \mu$L, 5.5 × 10$^{-5}$ mol) were then added with a microsyringe and after several minutes the solution turns to pale yellow. 3.2 mL of isoprene (0.032 mol) were then added, and the reaction was carried out at 20 °C for 3 h. At the end of the reaction the medium is viscous. The reaction was quenched by addition of some methanol drops. The polymer was dissolved in toluene, poured in a large amount of methanol, filtered and dried under vacuum. The yield reaches 40%, the average molecular weight of the obtained polymer is $M_n = 104$ 500 g mol$^{-1}$, and the polydispersity index of 2.0. $^1$H and $^{13}$C NMR analyses in CDCl$_3$ showed that the polyisoprene obtained was up to 91.7% 1,4-cis stereoregular.

Typical isoprene polymerisation experiment with 1a/L$^1$/[CPh$_3$][B(C$_6$F$_5$)$_4$]/Al(iBu)$_3$ (run 2.11, Table 3)

In a glove box under argon (H$_2$O and O$_2$ <2 ppm), 2.8 mg of complex Cp*Sc(BH$_4$)$_2$(THF) 1a (1 × 10$^{-5}$ mol) are dissolved in 0.5 mL of dry and degassed toluene. 1 equivalent of L$^1$ ligand (1.8 mg, 1 × 10$^{-5}$ mol) was added under stirring. After several minutes, 2 equivalents of [CPh$_3$][B(C$_6$F$_5$)$_4$] (18 mg, 2 × 10$^{-5}$ mol) were then added also under stirring and an orange solution was obtained. 10 equivalents of Al(iBu)$_3$ ($d = 0.786, 28 \mu$L, 1 × 10$^{-4}$ mol) were then added with a microsyringe and after several minutes the solution turns pale orange. 0.5 mL of isoprene (5 × 10$^{-3}$ mol) was then added, and the reaction was carried out at 20 °C. After 3 min the reaction medium is completely solid. The reaction was quenched by adding a THF solution containing some methanol drops. The polymer was poured in a large amount of methanol, filtered and dried under vacuum. The yield reaches 100%, the average molecular weight of the obtained polymer is $M_n = 13$ 900 g mol$^{-1}$, and the polydispersity index of 5.1. $^1$H and $^{13}$C NMR analyses in CDCl$_3$ showed that the polyisoprene obtained was 55.8% 3,4-stereoregular.
Typical isoprene polymerisation experiment with 1b/HL\textsubscript{2Mes}/BuEtMg/[CPh\textsubscript{3}][B(C\textsubscript{6}F\textsubscript{5})\textsubscript{4}]/Al(Bu\textsubscript{3}) (run 2.10, Table 3)

In a glove box under argon (H\textsubscript{2}O and O\textsubscript{2} <2 ppm), 4.5 mg of complex Cp\textsuperscript{+}Nd(BH\textsubscript{4})\textsubscript{2}(THF)\textsubscript{2}1b (1 × 10\textsuperscript{-5} mol) are dissolved in 0.5 mL of dry and degassed toluene. 1 equivalent of HL\textsubscript{2Mes} ligand (3.6 mg, 1 × 10\textsuperscript{-5} mol) was added under stirring, followed by 0.5 equivalent of BuEtMg (BEM, 20 wt% in hexanes, 4 μL, 5 × 10\textsuperscript{-6} mol). After several minutes, two equivalents of [CPh\textsubscript{3}][B(C\textsubscript{6}F\textsubscript{5})\textsubscript{4}] (18 mg, 2 × 10\textsuperscript{-5} mol) were then added also under stirring and a green–brown solution was obtained. 10 equivalents of Al(Bu\textsubscript{3}) (d = 0.786, 28 μL, 1 × 10\textsuperscript{-4} mol) were then added with a microsyringe. 0.5 mL of isoprene (5 × 10\textsuperscript{-3} mol) was then added, and the reaction was carried out at 20 °C for 65 h. At the end of the reaction the medium was viscous. The reaction was quenched by adding some methanol drops, and then dissolved in THF. The polymer was poured in a large amount of methanol, filtered and dried under vacuum. The yield reaches 100%, the average molecular weight of the obtained polymer is $M_n = 17\ 100\ g\ mol^{-1}$, and the polydispersity index of 1.9. \textsuperscript{1}H and \textsuperscript{13}C NMR analyses in CDCl\textsubscript{3} showed that the polyisoprene obtained was 68.9% trans-stereospecific.
Notes and references

[H NMR signals of Cp*Nd(BH₄)₂(THF)ₓ: when freshly synthesized, Cp*Nd(BH₄)₂(THF)ₓ is a monomer bearing two THF molecules (¹H NMR, THF-d₈, δ Cp* = 7.4 ppm, δ BH₄ = 69 ppm, ref. 33). With time, this complex tends to clustering as a desolvated hexamer leading to the appearance of a new BH₄ signal at 96 ppm either in THF-d₈ or benzene-d₆. This cannot be attributed to Nd(BH₄)₃(THF)₃ which gives a BH₄ signal at 92 ppm in THF-d₈. The X-ray structure of the hexamer 1b′ [Cp*Nd(BH₄)₂]₆ is available in the ESI (CCDC 892289). A similar hexamer [Cp(Me₄nPr)Nd(BH₄)₂]₆ was already reported.³⁰


S. Li, D. Cui, D. Li and Z. Hou, Organometallics, 2009, 28, 4814.


