Rigid NON-donor pincer ligand complexes of lutetium and lanthanum: synthesis and hydroamination catalysis†

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Reaction of H2XN2 (4,5-bis(2,4,6-triisopropylanilino)-2,7-di-tert-butyl-9,9-dimethylanthene) with [Lu(CH2SiMe3)3(THF)2], and crystallization from O(SiMe3)2, yielded [XN2]Lu(CH2SiMe3)(THF)|[O(SiMe3)2]0.25$^2_x$ 1$-$(O(SiMe3)2)2.5x. Lanthanum complexes of the XN2 dianion were also prepared by salt metathesis; treatment of H2XN2 with excess KH in DME produced the dipotassium salt, [K2(XN2)(DME)]x = 2--10), and subsequent reaction with [LaCl3(THF)]2 afforded [(XN2)LaCl(THF)]2:O(SiMe3)2. Compound 2 reacted with two equivalents of LiCH2SiMe3, to form the dialkyl-ate complex, [Li(THF)][(XN2)La(CH2SiMe3)2]:Toluene LiCl (3-toluene:LiCl; x = 3). Both 1 and 3 (x = 4) were structurally characterized, and were evaluated as catalysts for intramolecular hydroamination; while 3 showed poor activity, 1 is highly active for both intramolecular hydroamination and more challenging intermolecular hydroamination. Reactions with unsymmetrical alkenes yielded Markovnikov products, and the activity of 1 surpassed that of the previously reported yttrium analogue in the reaction of diphenylacetylene with 4-tert-butylbenzylamine.

1 Introduction

A variety of rare earth alkyl and amido complexes have demonstrated high activity for alkene and alkyne hydroamination catalysis, typically exceeding that achievable with transition metal catalysts. However, the ease with which hydroamination reactions occur varies greatly as a function of the substrate. In particular, intramolecular hydroamination reactions are much more facile than intermolecular variants, and intramolecular hydroamination reactions are generally more favorable for (a) alkynes vs. alkenes, (b) substrates leading to 5- vs. 6- vs. 7-membered ring formation, (c) 1-alkylalkenes and 1-alkinoalkenes di-substituted at the 2-position (Thorpe–Ingold effect; bulkier phenyl substituents are more effective than methyl substituents), and (d) aminocycloalkanes without additional alkene substitution (e.g. H2NCH2CPH2CH2CR = CHR$^\prime$ where R = R$^\prime$ = H).1-3 Intermolecular reactions with unactivated alkenes are particularly challenging, and only a handful of effective catalysts have been reported for these substrates, including rare earth anza-cyclopentadienyl complexes (a in Fig. 1) developed by Marks et al.6,7 yttrium complexes of both bidentate and tridentate 1,1′-binaphthalylene-backbone ligands (b and c in Fig. 1) reported by Hultzsch,8,9 and an yttrium complex of a rigid xanthene-backbone NON-donor pincer ligand (d in Fig. 1) reported recently by the Emslie group.10

For trivalent rare earth catalyzed alkene hydroamination, catalytic activity typically increases in parallel with metal ionic radius. For example, in cyclization reactions with H2C=CHCH2CMe2CH2NH2, the activity of [Cp*Ln(CH(SiMe3))]0, [Me5Si(C5Me4)2]0/2Ln(CH(SiMe3))2] (b in Fig. 1) and [[(L)Ln

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Fig. 1 Highly active rare earth catalysts for intermolecular hydroamination of unactivated alkenes (Ln = Nd; Ln$^\prime$ = Y).
Crystals of 1-C_{6}H_{5}O.5 (lattice solvent is residual benzene from the synthesis) were grown by cooling a concentrated O(SiMe_{3})_{2} solution to −30 °C (Fig. 2). In the solid state, the XN_{2} backbone is slightly bent with a 26° angle away from planarity, based on the orientation of the two xanthene aryl rings. This orientation is mirrored in [(XN_{2})Y(CH_{3}SiMe_{3})_{2}(THF)], which displayed a backbone angle of 25° and a very similar geometry at the rare earth metal.\textsuperscript{18} Lutetium is 5-coordinate with the three anionic donors and coordinated THF arranged in an approximate tetrahedron around the metal center. The largest angle in this approximate tetrahedron is the N(1)--Lu--N(2) angle of 130°, and the smallest is the O(2)--Lu--C(54) angle of 97°, while the other angles are between 102° and 110°. The oxygen donor of the xanthene backbone is coordinated on the N(1)/N(2)/C(54) face of the tetrahedron closest to the nitrogen donors. Lutetium lies 0.74 Å out of the plane of the XN_{2} ligand donor atoms, leading to a 53° angle between the NON and the NLuN planes. The neutral oxygen donor on the xanthene backbone is located 0.5 Å out of the N(1)/C(4)/C(5)/N(2) plane in order to coordinate lutetium, with N--Lu--O(1) distances of 69 and 70 Å. Additionally, the oxygen donors on the xanthene backbone are located on the N(2) ligand are bent towards lutetium, illustrated by the C(1)···C(8), C(4)···C(5) and N(1)···N(2) distances of 5.02 Å, 4.57 Å and 4.04 Å respectively, which are comparable with those in [(XN_{2})Y(CH_{3}SiMe_{3})_{2}(THF)] (4.98 Å, 4.56 Å and 4.06 Å).\textsuperscript{18}

The Lu–N distances of 2.221(2) Å and 2.228(2) Å are slightly shorter than those in [(XN_{2})Y(CH_{3}SiMe_{3})_{2}(THF)] (2.252(3) Å),\textsuperscript{18} consistent with the smaller ionic radius of Lu\textsuperscript{III} compared to Y\textsuperscript{III} (0.861 Å vs. 0.900 Å).\textsuperscript{18} Additionally, the Lu–N distances in 1 fall within the range reported for related compounds such as [(2:ArN--CMC)(6-ArNCMe_{2})C_{6}H_{5}N(Lu(CH_{3}SiMe_{3})_{2})] (Ar = C_{6}H_{5}Pr_{2}-2,6; 2.188(4) Å)\textsuperscript{19} and [(1,8-(Pz)_{2})C_{2}Lu(CH_{3}SiMe_{3})_{2}] (Pz = 1-iso-propylpyrazolyl; Cz = 3,6-dimethylcarbazole; 2.233(1) Å).\textsuperscript{20} The Lu-C(54) distance of 2.326(2) Å is shorter compared to that in [(XN_{2})Y(CH_{3}SiMe_{3})_{2}(THF)] (2.364(3) Å), also in keeping with the relative sizes of yttrium and lutetium (vide supra). Additionally, this distance falls within the range of Lu–C distances reported in the literature. For example, the Lu–C distances in the aforementioned alkyl complexes range from 2.329(6) Å to 2.374(3) Å,\textsuperscript{10} and Lu–C in [(2:2-NAr)(6-Xyl)C_{5}H_{3}N_{2}Lu(CH_{3}SiMe_{3})_{2}(THF)] (Ar = C_{6}H_{5}Pr_{2}-2,6; Xyl = o-xylolyl) is 2.323(14) Å.\textsuperscript{13} [(XN_{2})Lu(CH_{3}SiMe_{3})_{2}(THF)] (O(SiMe_{3})_{2})_{1.5} (1·O(SiMe_{3})_{2})_{1.5} was tested as an ethylene polymerization catalyst at 24 °C and 80 °C (toluene, 1 atm ethylene, 1 h) but exhibited negligible activity. Compound 1 was also investigated as a catalyst for both intra- and inter-molecular hydroamination and the results are summarized in Tables 1 and 2. Compound 1 catalyzed intramolecular hydroamination of a range of substrates in benzene at 24 °C, proceeding to >99% completion in all cases. The time required to reach >99% completion was slightly increased compared to reactions catalyzed by the yttrium complex, [(XN_{2}) Y(CH_{3}SiMe_{3})_{2}(THF)], which is consistent with the majority of previous reports (vide supra), in which hydroamination activity increases with increasing rare earth metal size.\textsuperscript{19} This is particularly evident in entries 2 and 4 in Table 1, as [(XN_{2}) Y(CH_{3}SiMe_{3})_{2}(THF)] achieved >99% conversion after 1.5 h and
34 h,\textsuperscript{10} whereas 1 required 2.75 h and 48 h respectively. Nevertheless, the ability of 1 and the yttrium analogue to catalyze these more challenging intramolecular hydroamination reactions at room temperature stands these catalysts apart from most others.\textsuperscript{10}

Compound 1 also catalyzed intermolecular hydroamination with 4-tert-butylaniline, 4-tert-butylbenzylamine and octylamine in combination with 1-octene and diphenylacetylene, and in all reactions with 1-octene, the Markovnikov product was formed selectively. These reactions were performed in toluene at 110 °C and the degree of conversion was determined by GC-MS (Table 2). Over a 24 h time period, the reaction of 1-octene with octylamine (entry 3) resulted in a turnover frequency ($N_t$) of 0.41 h$^{-1}$, which is greater than that obtained for the reaction with 4-tert-butylbenzylamine (entry 2, 0.35 h$^{-1}$), which in turn is significantly greater than that obtained for the reaction with 4-tert-butylaniline (entry 1, 0.04 h$^{-1}$). These results are consistent with the increased donor ability and reduced steric bulk of the former amines. The same trend was previously observed for [(XN$_2$)$_2$Y(CH$_2$SiMe$_3$)(THF)],\textsuperscript{10} and the ability of 1 to catalyze intermolecular hydroamination of 1-octene (an unactivated alkene) places it in a select group of catalysts with this capability (vide supra). The intermolecular hydroamination activity of 1 closely mirrors that of the yttrium analogue, although for entries 2 and 5 in Table 2, compound 1 afforded lower and higher activities, respectively ($N_t$ = 0.35 vs. 0.40 and 0.42 vs. 0.33). The intermolecular reactions with the largest conversions after 24 h at 110 °C (10 mol% catalyst) were those utilizing diphenylacetylene (entries 4–6), as the amounts of unreacted 4-tert-butylaniline, 4-tert-butylbenzylamine and octylamine were below the detection limit of the GC instrument.

In order to further explore the effectiveness of the XN$_2$ ligand for rare earth coordination, and the impact of metal ionic radius on hydroamination activity, the synthesis of lanthanum XN$_2$ complexes was undertaken. As the lanthanum trialkyl compound, [La(CH$_2$SiMe$_3$)$_3$(THF)] is not readily accessible,\textsuperscript{34}

\begin{table}[h]
\centering
\begin{tabular}{|c|c|c|c|c|c|}
\hline
Entry & Reagent & Product & Mol\% & Time & Temp. (°C) & Product formation$^a$ & $N_t$ (h$^{-1}$) \\
\hline
1 & \begin{tabular}{c}
\includegraphics[width=0.2\textwidth]{fig1.png}
\end{tabular} & \begin{tabular}{c}
\includegraphics[width=0.2\textwidth]{fig1.png}
\end{tabular} & 1 & <10 min & 24 & >99\% & \geq 600 \\
2 & \begin{tabular}{c}
\includegraphics[width=0.2\textwidth]{fig1.png}
\end{tabular} & \begin{tabular}{c}
\includegraphics[width=0.2\textwidth]{fig1.png}
\end{tabular} & 1 & 2.75 h & 24 & >99\% & \sim 36 \\
3 & \begin{tabular}{c}
\includegraphics[width=0.2\textwidth]{fig1.png}
\end{tabular} & \begin{tabular}{c}
\includegraphics[width=0.2\textwidth]{fig1.png}
\end{tabular} & 10 & <20 min & 24 & >99\% & \geq 30 \\
4 & \begin{tabular}{c}
\includegraphics[width=0.2\textwidth]{fig1.png}
\end{tabular} & \begin{tabular}{c}
\includegraphics[width=0.2\textwidth]{fig1.png}
\end{tabular} & 10 & 48 h & 24 & >99\% & \sim 0.2 \\
\hline
\end{tabular}
\caption{Intramolecular hydroamination reactivity with 1 in C$_6$D$_6$}
\label{tab:intramolecular}
\end{table}

$^a$ Conversion of reactant to product determined by $^1$H NMR spectroscopy. $^b$ Turnover frequency.

\begin{figure}[h]
\centering
\includegraphics[width=0.8\textwidth]{fig2.png}
\caption{Two views of the X-ray crystal structure for 1·(C$_6$H$_5$)$_0.5$. Ellipsoids are set to 50%. Hydrogen atoms and lattice solvent are omitted. The tert-butyl groups are rotationally disordered over two positions, and only one is shown for clarity. In view B the 2,4,6-triisopropylphenyl groups are depicted in wire-frame format. Selected bond lengths [Å] and angles [°]: Lu–N(1) 2.221(2), Lu–N(2) 2.228(2), Lu–C(54) 2.326(2), Lu–O(1) 2.299(1), Lu–O(2) 2.264(1), N(1)–Lu–N(2) 130.41(6), N(1)–Lu–C(54) 106.09(7), N(2)–Lu–C(54) 110.41(7), O(1)–Lu–C(54) 104.38(6), O(2)–Lu–C(54) 97.56(7), O(2)–Lu–N(2) 105.39(6), O(2)–Lu–N(1) 101.76(6).}
\label{fig:crystal}
\end{figure}
salt metathesis was employed for ligand attachment in the place of alkane elimination. Stirring H$_2$XN$_2$ with excess KH in DME at 24°C produced the dipotassium salt of the 4,5-bis(2,4,6-triisopropylanilido)-2,7-di-tert-butyl-9,9-dimethylxanthene ligand, [K$_2$(XN$_2$)(DME)$_x$] ($x = 2$–2.5), as a beige solid in 80% isolated yield. [K$_2$(XN$_2$)(DME)$_2$] was reacted with [LaCl$_3$(THF)$_3$] in THF at 24°C, and a recrystallization from O(SiMe$_3$)$_2$, [(XN$_2$)LaCl(THF)$_x$]$_{O(SiMe_3)_2}$, was obtained in 51% yield as an off-white solid (Scheme 2).

Attempts to synthesize a lanthanum monoalkyl complex were undertaken via reactions of trimethylsilylmethyl lithium and methyl lithium with [(XN$_2$)LaCl(THF)$_x$]$_{O(SiMe_3)_2}$, 2(O(SiMe$_3$)$_2$)$_{0.25x}$; $x = 1$ or 2). NMR-scale reactions were performed in d$_8$-THF at 24°C and both resulted in the formation of a dialkyl-‘ate’ complex, based on the integrations of the respective alkyl peaks, regardless of whether one equivalent (per La) or an excess of the alkali metal-alkyl reagent was added. The reaction utilizing trimethylsilylmethyl lithium was pursued

<table>
<thead>
<tr>
<th>Entry</th>
<th>Amine or alkyn$^c$</th>
<th>Product</th>
<th>Temp. (°C)</th>
<th>Time (h)</th>
<th>Product formation$^{b,c}$</th>
<th>% Markovnikov product$^c$</th>
<th>$N_f^d$ (h$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Bu$_3$N-CH$_2$Bu</td>
<td></td>
<td>24</td>
<td>110</td>
<td>11%</td>
<td>97</td>
<td>0.04</td>
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<tr>
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<td>83%</td>
<td>&gt;99</td>
<td>0.35</td>
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<tr>
<td>3</td>
<td>H$_2$N-CH$_2$Ph</td>
<td></td>
<td>24</td>
<td>110</td>
<td>98%</td>
<td>&gt;99</td>
<td>0.41</td>
</tr>
<tr>
<td>4</td>
<td>Bu$_3$N-CH$_2$Ph</td>
<td></td>
<td>24</td>
<td>110</td>
<td>&gt;99%</td>
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<td>0.42</td>
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<tr>
<td>5</td>
<td>Bu$_3$N-CH$_2$Ph</td>
<td></td>
<td>24</td>
<td>110</td>
<td>&gt;99%</td>
<td>N/A</td>
<td>0.42</td>
</tr>
<tr>
<td>6</td>
<td>H$_2$N-CH$_2$Ph</td>
<td></td>
<td>24</td>
<td>110</td>
<td>&gt;99%</td>
<td>N/A</td>
<td>0.42</td>
</tr>
</tbody>
</table>

$^a$ Alkene/alkyne present in 20 fold excess relative to the amine. $^b$ Conversion determined by product: unreacted amine ratio. $^c$ Determined by GC-MS. $^d$ Turnover frequency. $^e$ In entry 4 the product is formed as a single isomer, whereas in entries 5 and 6 the products are formed as 1 : 0.35 and 1 : 0.24 mixtures of the E and Z isomers (based on literature assignments for similar compounds), respectively.

**Table 2** Intermolecular hydroamination reactivity with 1 (10 mol%) in toluene

Scheme 2 Synthesis of [K$_2$(XN$_2$)(DME)$_x$] and complexes 2 and 3.
further, as it provides a direct comparison with the lutetium and yttrium trimethylsilylmethyl complexes of the XN₂ ligand. \[[[\text{XN}_2]\text{LaCl(THF)}_2]_2 \cdot (\text{O(SiMe}_3)_2)_0.255, 2 \cdot (\text{O(SiMe}_3)_2)_0.255; x = 1 \text{ or } 2\] reacted with 2 equivalents of trimethylsilylmethyl lithium in THF at 24 °C, and after removal of the salts by centrifugation in toluene and layering with hexanes at -30 °C, \[[\text{Li(THF)}_4]_2[[\text{XN}_2]_2\text{La(CH}_2\text{SiMe}_3)_2 \cdot \text{toluene-LiCl} (3 \cdot \text{toluene-LiCl}; x = 3)\] was isolated as a pale yellow solid in 55% yield (Scheme 2). Compound 3 is sparingly soluble in benzene and other non-polar solvents such as hexanes and pentane, so all characterization was carried out in d₄-THF. The 1H NMR spectrum of 3 revealed the expected signals for the XN₂ ligand backbone and only one set of signals for the two alkyl groups was observed between 25 and -80 °C (a singlet with an integration of 18 for LaCH₂SiMe₃ and a singlet with an integration of 4 for LaCH₂SiMe₃). Crystals of [Li(THF)]_2[[XN]_2]La(CH₂SiMe₃)_2 · THF were grown from a concentrated THF solution of 3 · tolune · LiCl (x = 3), layered with pentane and cooled to -30 °C (Fig. 3).

In the solid state, lanthanum is 5-coordinate with the two amido donors and two alkyl groups arranged in a distorted tetrahedron around the metal center. The largest angle in this approximate tetrahedron is the N(1)–La–N(2) angle of 118°, and the smallest is the C(54)–La–C(58) angle of 100°, while the other angles are between 101° and 106°. Lanthanum lies 0.96 Å out of the plane of the XN₂ ligand donor atoms, leading to a 50° angle between the NON and NLAN planes. The XN₂ backbone is bent with a 35° angle away from planarity, based on the orientation of the two aryl rings of the xanthene backbone. The neutral oxygen donor of the xanthene donor is located 0.64 Å out of the N(1)/(C(4)/(C(5)/(N(2) plane in order to coordinate to lanthanum, resulting in N-La-O(1) angles of 63°-64°. In addition, it is of note that the nitrogen donors on the ligand are not bent towards lanthanum to the extent that they are in 1, illustrated by the C(1)–C(8), C(4)–C(5) and N(1)–N(2) distances in 3 of 4.90 Å, 4.57 Å and 4.21 Å respectively, compared to 5.02 Å, 4.57 Å and 4.04 Å in 1. The La–N distances of 2.462(5) Å and 2.445(5) Å are substantially lengthened relative to those in [[XN]_2]Y(CH₂SiMe₃)_2[THF] [2.392(4) Å]³⁹ and [[XN]_2]Lu(CH₂SiMe₃)_2[THF] · (O(SiMe₃)_2)_0.15 (1) [2.22(1) Å and 2.28(1) Å], consistent with the large ionic radius of lanthanum compared to yttrium and lutetium, combined with increased steric hindrance and an overall negative charge in 3, resulting in a less electrophilic metal centre. For analogous reasons, the La–C(54) and La–C(58) distances of 2.573(7) Å and 2.613(7) Å are also greatly elongated compared to those in [[XN]_2]Y(CH₂SiMe₃)_2[THF] [2.364(3) Å]³⁹ and 1 [2.362(2) Å]. However, both the La–N and La–C distances in 3 are significantly shorter than those previously reported for [(R)-Binap(NCyp)]_2La[(μ-Me)]Li(THF)][(μ-Me)Li(THF)]_2 (Binap = 2,2’-disubstituted-1,1’-binaphthyl; Cyp = cyclopentyl) and [La = 2.626(7)–2.677(8) Å; La–C = 2.704(8)–2.832(11) Å,]²⁹,³⁷

Reaction of [Li(THF)]_2[[XN]_2]La(CH₂SiMe₃)_2 · tolune-LiCl (3 · tolune-LiCl) with [[XN]_2]LaCl[THF]_2 · (O(SiMe₃)_2)_0.255, 2 · (O(SiMe₃)_2)_0.255; x = 1 or 2) did not provide access to the neutral XN₂ lanthanum alkyl; at 24 °C no reaction was observed, and heating to 70 °C resulted only in thermal decomposition of 3 · tolune-LiCl.

Rare earth alkyl-‘ate’ complexes have been reported to catalyze intramolecular hydroamination as well as intermolecular hydroamination. A few examples include, [(R)-Binap(NCyp)]_2La[(μ-Me)]Li(THF)][(μ-Me)Li(THF)]_2 which catalyzed asymmetric intramolecular hydroamination of amino-1,3-dienes,²⁹ [Li(THF)]_4[[(R)-Binap(NCyp)]_2Y(CH₂SiMe₃)_2] (Binap = 2,2’-disubstituted-1,1’-binaphthyl; Cyp = cyclopentyl) which catalyzed intramolecular hydroamination of secondary amino-alkenes,³⁶ [Li(THF)]_4[[(ArNC(Me)=C(Me)NAr)]_2Y(CH₂SiMe₃)_2] (Ar = C₆H₄(Ph₂)C₆H₄) which catalyzed the intermolecular hydroamination reaction of styrene and pyrrolidine,³⁹ and [(R)-Binap(NCyp)]_2[(μ-Me)]Li(THF)][(μ-Me)Li(THF)]_2 (Binap = 2,2’-disubstituted-1,1’-binaphthyl; Cyp = cyclopentyl) which catalyzed 1-amino-2,2-diphenyl-4-pentene cyclization, requiring 1.9 h at 25 °C with 6 mol% catalyst loading to reach 100%.

Fig. 3  Two views of the X-ray crystal structure for 3 · THF (x = 4). Ellipsoids are set to 50%. Hydrogen atoms and lattice solvent are omitted. The tert-butyl groups are rotationally disordered over two positions, and only one is shown for clarity. In view A the [Li(THF)]_4⁺ cation is omitted for clarity. In view B the 2,4,6-trisopropylphenyl groups and the THF molecules are depicted in wire-frame format. Selected bond lengths [Å] and angles [°]: La–N(1) 2.462(5), La–N(2) 2.445(5), La–C(54) 2.573(7), La–C(58) 2.613(7), La–O(1) 2.643(4), N(1)–La–N(2) 118.22(17), N(1)–La–C(54) 108.9(2), N(1)–La–C(58) 100.9(2), N(2)–La–C(54) 106.3(2), N(2)–La–C(58) 120.7(2), O(1)–La–C(54) 97.63(18), O(1)–La–C(58) 159.7(2), C(54)–La–C(58) 100.0(2), N(1)–La–O(1) 63.59(15), N(2)–La–O(1) 62.72(15).
conversion. Complex 3-toluene–LiCl, (x = 3) was tested as a catalyst for intramolecular hydroamination with 1-amino-2,2-diphenyl-4-pentene in d8-THF at 24 °C. However, the time required to reach >99% completion (45 h) was significantly increased compared to that required for [[(THF)2(CH2SiMe3)] (THF)] and [[(THF)Lu(CH2SiMe3)(THF)] (1–O(SiMe3)2)].<10 min] under analogous conditions in benzene (or in THF for the yttrium complex). Consequently, the catalytic activity of 3 was not further investigated.

3 Summary and conclusion

The rigid NON-donor pincer ligand, XN₃, has been employed for the synthesis of a lutetium monoalkyl complex (1), a lanthanum chloride complex (2) and a lanthanum dialkyl-‘ate’ complex (3). Complex 3 was tested as a catalyst for intramolecular hydroamination, but showed low activity. By contrast, the neutral lutetium alkyl complex, 1, is highly active for both intra- and inter-molecular hydroamination with a variety of substrates. For intramolecular alkene hydroamination, the time required to reach >99% completion was slightly increased compared to the previously reported yttrium analogue. By contrast, the inter-molecular hydroamination reaction between 4-tert-butylbenzylamine and diphenylacetylene afforded a higher turnover number than the yttrium analogue.

4 Experimental

General details

An argon-filled MBraun UNIlab glove box equipped with a −30 °C freezer was employed for the manipulation and storage of all air-sensitive compounds, and reactions were performed on a double manifold high vacuum line using standard techniques. Residual oxygen and moisture was removed from the argon stream by passage through an Oxisorb-W scrubber from Matheson Gas Products. A Fisher Scientific Ultrasonic FS-30 bath was used to sonicate reaction mixtures where indicated. A VWR Clinical 200 Large Capacity Centrifuge (with 28’ fixed-angle rotors that hold 12 × 15 mL or 6 × 50 mL tubes) in combination with 15 mL. A Kimble Chase glass centrifuge tubes was used when required (inside the glovebox). Diethyl ether, THF, toluene, benzene and hexanes were initially dried and distilled at atmospheric pressure from Na/Ph₂CO. Hexamethyldisiloxane (O(SiMe₃)₂) was dried and distilled at atmospheric pressure from Na/Ph₂CO. Hexane, hexamethyldisiloxane (O(TMS)₂/O(SiMe₃)₂) = Na/Ph₂CO/tetra-glyme; Et₂O = Na/Ph₂CO and introduced to reactions via vacuum transfer with condensation at −78 °C. The deuterated solvents (ACP Chemicals) C₆D₆, THF-d₈ and toluene- d₈ were dried over Na/Ph₂CO. 

The H₂XN₃₄₈, [Lu(CH₂SiMe₃)₂(THF)]_₂ and the commercially available intramolecular hydroamination reagents<6,46> were prepared according to literature procedures. 1-Amino-5-hexene was purchased from GFS Chemicals, dried over CaH₂ and distilled prior to use. 1,3,5-Triisopropylbenzene, xanthone, KH (30 wt% in mineral oil), LiCh₂SiMe₃ (1.0 M in pentane), MeLi (1.6 M in Et₂O), nBuLi (1.6 M in hexanes), Br₂, NaH, NaO'Bu, Pd(OAc)₂, DPEPhos, [bis(2-diphenylphosphino)phenyl]ether, diphenylacetylene and MgSO₄ were purchased from Sigma-Aldrich, LuCl₃ and LaCl₃ were purchased from Strem Chemicals. Solid LiCh₂SiMe₃ and MeLi were obtained by removal of the solvent in vacuo, and solid KH was obtained by filtration and washing with hexanes. [LuCl₃(THF)]₃ and [LaCl₃(THF)]₃ were obtained by refluxing the anhydrous lutetium/lanthanum trihalide in THF for 24 h followed by removal of the solvent in vacuo. 4-tert-Butyl-aniline, 4-tert-butylbenzylamine, n-octylamine and 1-ocetone were purchased from Sigma-Aldrich, dried over molecular sieves and distilled prior to use. Argon (99.999% purity) and ethylene (99.999% purity) were purchased from Praxair.

Combustion elemental analyses were performed both at McMaster University on a Thermo EA1112 CHNS/O analyzer and by Midwest Microlab, LLC, Indianapolis, IN, USA. NMR spectroscopy (¹H, ¹³C) DEPT-Q, COSY, HMQC, HMB, All ¹H NMR and ¹³C NMR spectra were referenced relative to SiMe₄ through a resonance of the employed deuterated solvent or proteo impurity of the solvent; C₆D₆ (7.16 ppm), d₈-Tol (2.08, 6.97, 7.01, 7.09 ppm), d₈-THF (1.72, 3.58 ppm) for ¹H NMR; and C₆D₆ (128.0 ppm), d₈-Tol (20.43, 125.13, 127.96, 128.87, 137.48 ppm), d₈-THF (25.31, 67.21 ppm) for ¹³C NMR. Herein, numbered proton and carbon atoms refer to the positions of the xanthene backbone, as shown in Scheme 1. Inequivalent ortho isopropyl protons are labeled A and B, while inequivalent aryl ring protons and inequivalent methyl protons are labeled ‘a’ and ‘b’, so that the corresponding carbon resonances can be identified.

X-ray crystallographic analyses were performed on suitable crystals coated in Paratone oil and mounted on a SMART APEX II diffractometer with a 3 kW sealed tube Mo generator in the McMaster Analytical X-ray (MAX) Diffraction Facility. In all cases, non-hydrogen atoms were refined anisotropically and hydrogen atoms were generated in ideal positions and then updated with each cycle of refinement. GC-MS analyses were performed using an Agilent 6890N gas chromatograph (Santa Clara, CA, USA), equipped with a DB-17ht column (30 m × 0.25 mm i.d. × 0.15 μm film, J & W Scientific) and a retention gap (deactivated fused silica, 5 m × 0.53 mm i.d.), and coupled to an Agilent 5973 MSD single-quadrupole mass spectrometer. One microliter of sample was injected using Agilent 7683 autosampler in splitless mode. The injector temperature was 230 °C and carrier gas (helium) flow was 0.7 mL min⁻¹. The transfer line was 250 °C and the MS source temperature was 230 °C. The column temperature started at 50 °C and was raised to 300 °C at 8 °C min⁻¹. It was then held at 300 °C for 15 min to give a total run time of 46.25 min. Full scan mass spectra between m/z 50 and 800 were acquired after a five minute solvent delay.
CH$_4$), 6.23 (br s, 2H, Xanth-CH$_3$), 4.24 (sept, 2H, $^3$J$_{HH}$ 6.73 Hz, A-ortho-CH$_3$Me$_2$), 3.38 (sept, 2H, $^3$J$_{HH}$ 6.78 Hz, B-ortho-CH$_3$Me$_2$), 2.83 (sept, 2H, $^3$J$_{HH}$ 6.74 Hz, para-CH$_3$Me$_2$), 2.73 (s, 4H, 1 equiv. THF-C$_3$H$_2$), 1.88 (s, 3H, C$_5$Me$_3$), 1.72 (s, 3H, C$_5$Me$_3$), 1.49 (d, 6H, $^3$J$_{HH}$ 6.73 Hz, A-ortho-CH$_3$Me$_2$), 1.47 (d, 6H, $^3$J$_{HH}$ 6.73 Hz, A-ortho-CH$_3$Me$_2$), 1.26 (s, 18H, C$_5$Me$_3$), 1.24 (d, 6H, $^3$J$_{HH}$ 6.78 Hz, B-ortho-CH$_3$Me$_2$), 1.21 (m, 12H, para-CH$_3$Me$_2$), 1.01 (d, 6H, $^3$J$_{HH}$ 6.78 Hz, B-ortho-CH$_3$Me$_2$), 0.83 (s, 4H, 1 equiv. THF-C$_3$H$_2$), 0.35 (s, 9H, LuCh$_2$SiMe$_3$), −0.40 (s, 2H, LuCh$_2$SiMe$_3$).

$^{13}$C NMR (CD$_2$Cl$_2$, 126 MHz): $^\delta$ 147.83 (Xanth-$^\delta$C), 147.72 (Xanth($^\gamma$C), 147.38 (A-ortho-CH$_3$Me$_2$), 146.17 (B-ortho-CH$_3$Me$_2$), 145.33 (para-CH$_3$Me$_2$), 141.82 (Ar-Cipso), 141.30 (Xanth($^\alpha$C)), 130.56 (Xanth($^\alpha$C)), 122.20 (Ar-CH$_2$), 121.85 (Ar-CH), 109.35 (Xanth-$^\gamma$C), 106.47 (Xanth-$^\beta$C), 70.85 (THF-C$_3$H$_2$), 37.99 (LuCh$_2$SiMe$_3$), 35.57 (Xanth-$^\alpha$C$_2$Me$_2$), 35.25 (CMe$_3$), 35.05 (CMe$_3$), 34.50 (para-CH$_3$Me$_2$), 31.91 (CMe$_3$), 28.39 (B-ortho-CH$_3$Me$_2$), 27.67 (A-ortho-CH$_3$Me$_2$), 27.12 (A-ortho-CH$_3$Me$_2$), 25.85 (B-ortho-CH$_3$Me$_2$), 25.48 (B-ortho-CH$_3$Me$_2$), 25.04 (CMe$_3$), 24.85 (THF-C$_3$H$_2$), 24.62 (A-ortho-CH$_3$Me$_2$), 24.51 (para-CH$_3$Me$_2$), 4.13 (LuCh$_2$SiMe$_3$). Anal. calcd for C$_{129}$H$_{208}$N$_{25}$O$_{25}$Si$_{25}$LaCl: C, 66.41; H, 8.36; N, 2.68%. Found: C, 66.89; H, 8.83; N, 2.60%.

$\mathrm{[Li(THF)\{\{XN$_2\}\{La(CH$_2$SiMe$_3$)$_2$\}\}] (3-toluene-LiCl).}$

$\mathrm{[Li(THF)\{\{XN$_2\}\{La(CH$_2$SiMe$_3$)$_2$\}\}] (x = 1 or 2)}$ (0.086 g, 0.82 mmol mol of THF) was dissolved in 4 mL of THF and added to LiCH$_2$SiMe$_3$ (0.015 g, 0.165 mmol), which was stirred at 24 °C for 5 days. The solvent was removed in vacuo and the yellow solid was dissolved in toluene (1 mL), centrifuged, and the mother liquors were separated with hexanes and stored at −30 °C. Very small pale yellow crystals of $\{\{XN$_2\}\{Li(CH$_2$SiMe$_3$)$_2$\}\}$-toluene-LiCl were obtained (0.064 g, 55%). X-ray quality crystals of $\{\{XN$_2\}\{Li(CH$_2$SiMe$_3$)$_2$\}\}$ were grown from a concentrated THF solution of 3, layered with pentane and cooled to −30 °C.

$^1$H NMR (d$_6$-THF, 600 MHz): $^\delta$ 7.00 (s, 4H, Ar-CH$_2$), 6.28 (2H, $^3$J$_{HH}$ 2.16 Hz, Xanth-$^\gamma$C), 5.62 (2d, 2H, $^3$J$_{HH}$ 2.18 Hz, Xanth-$^\gamma$C), 3.55 (sept, 4H, $^3$J$_{HH}$ 6.84 Hz, ortho-CH$_3$Me$_2$), 2.86 (sept, 2H, $^3$J$_{HH}$ 6.93 Hz, para-CH$_3$Me$_2$), 1.61 (s, 6H, CMe$_3$), 1.25 (d, 12H, $^3$J$_{HH}$ 6.9 Hz, para-CH$_3$Me$_2$), 1.24 (d, 12H, $^3$J$_{HH}$ 6.8 Hz, A-ortho-CH$_3$Me$_2$), 1.10 (s, 18H, CMe$_3$), 1.00 (d, 12H, $^3$J$_{HH}$ 6.8 Hz, B-ortho-CH$_3$Me$_2$), −0.39 (s, 18H, LaCH$_2$SiMe$_3$), −1.17 (br. s, 4H, LaCH$_2$SiMe$_3$).

$^{13}$C NMR (d$_6$-THF, 126 MHz): $^\delta$ 146.81 (Xanth-$^\delta$C), 146.17 (ortho-CH$_3$Me$_2$), 145.26 (para-CH$_3$Me$_2$), 140.45 (Ar-Cipso), 128.34 (Xanth($^\alpha$C)), 122.57 (Ar-CH), 108.43 (Xanth($^\gamma$C)), 107.41 (Xanth($^\beta$C)), 67.22 (THF-C$_3$H$_2$), 35.07 (CMe$_3$), 34.96 (para-CH$_3$Me$_2$), 33.05 (CMe$_3$), 31.81 (CMe$_3$), 29.19 (ortho-CH$_3$Me$_2$), 25.84 (B-ortho-CH$_3$Me$_2$), 25.25 (THF-C$_3$H$_2$), 24.52 (para-CH$_3$Me$_2$), 24.16 (A-ortho-CH$_3$Me$_2$).

General procedure for intramolecular hydroamination

In the glove box, the appropriate amounts of 1 or 3 and the hydroamination substrate were weighed into separate vials, dissolved in C$_6$D$_6$ or d$_6$-THF and placed in a teflon-valved J-young NMR tube. The reactions were monitored at 24 °C by $^1$H NMR spectroscopy and the expected products were confirmed by their agreement to reported literature spectra.22

General procedure for intermolecular hydroamination

In the glove box, the appropriate amounts of 1, amine, and the alkene/alkyne were weighed into separate vials, dissolved in d$_a$-THF, selected for $^1$H NMR spectroscopy and the expected products were confirmed by their agreement to reported literature spectra.22

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toluene, placed in a Teflon-valved J-young NMR tube and then placed into a preheated oil bath at 110 °C. After heating for the designated amount of time, NMR spectra were obtained and the sample was submitted for analysis by GC-MS.

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