SUPPLEMENTARY INFORMATION

List of Figures

Figure S1: Characterization of $[La(L^{Et})_3]$: (a) IR spectrum; (b) ¹H NMR spectrum at 303K in C₆D₆; (c) ¹⁹F{¹H} NMR spectrum at 303K in C₆D₆; and (d) ¹⁹F{¹H} NMR spectrum at 203K in C₇D₈.^{*}

Figure S2: Characterization of $[Ce(L^{Me})_3]$: (a) IR spectrum; (b) ¹H NMR spectrum at 303K in C₇D₈; (c) ¹H NMR spectrum at 343K in C₇D₈; (d) ¹⁹F{¹H} NMR spectrum at 303K in C₇D₈ and (e) ¹⁹F{¹H} NMR spectrum at 343K in C₇D₈.^{*}

Figure S3: Characterization of $[Ce(L^{Me})_2F]_3$: (a) ¹H NMR spectrum at 373K in C₇D₈; (b) ¹⁹F{¹H} NMR spectrum at 373K in C₇D₈.^{*}

Figure S4: Characterization of $[Nd(L^{Me})_3]$: (a) IR spectrum; (b) ¹H NMR spectrum at 303K in C₇D₈; and (c) ¹⁹F{¹H} NMR spectrum at 343K in C₇D₈.^{*}

S5: GC /MS analyses of reaction mixture forming $[Ce(L^{Me})_3]$ and $[CeF(L^{Me})_2]_3$:

*Due to the high sensitivity of the complexes to moisture, traces of ligand HL^{Me}/HL^{Et} were present in the NMR spectra of the $[Ln(L^{Me/Et})_3]$ or $[Ln(L^{Me})_2F]_n$ products, as shown from their spectra, e.g. the two singlet or two multiplet resonances at -140 to -142 ppm and -160 to -162 ppm in the ${}^{19}F{}^{1}H{}$ NMR spectrum, belong to F3,5 and F2,6 respectively of HL^{Me}/HL^{Et} .

If both products $[Ln(L^{Me/Et})_3]$ and $[Ln(L^{Me})_2F]_n$ were isolated from the same reaction, one or the other was sometimes present as a minor contaminant in the isolated $[Ln(L^{Me/Et})_3]$ or $[Ln(L^{Me})_2F]_n$ products, e.g. resonances at -142.8 and -177.6 ppm, which belong to $[Ce(L^{Me})_3]$, are visible in the ¹⁹F{¹H} NMR spectrum of $[Ce(L^{Me})_2F]_3$ (Figure S3(b)).

Figure S1: Characterization of $[La(L^{Et})_3]$: (a) IR spectrum; (b) ¹H NMR spectrum at 303K in C₆D₆; (c) ¹⁹F{¹H} NMR spectrum at 303K in C₆D₆; and (d) ¹⁹F{¹H} NMR spectrum at 203K in C₇D₈.^{*}





Figure S2: Characterization of $[Ce(L^{Me})_3]$: (a) IR spectrum; (b) ¹H NMR spectrum at 303K in C₇D₈; (c) ¹H NMR spectrum at 343K in C₇D₈; (d) ¹⁹F{¹H} NMR spectrum at 303K in C₇D₈ and (e) ¹⁹F{¹H} NMR spectrum at 343K in C₇D₈.^{*}

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(b)



(d)



Figure S3: Characterization of $[Ce(L^{Me})_2F]_3$: (a) ¹H NMR spectrum at 373K in C₇D₈; (b) ¹⁹F{¹H} NMR spectrum at 373K in C₇D₈.^{*}





Figure S4: Characterization of $[Nd(L^{Me})_3]$: (a) IR spectrum; (b) ¹H NMR spectrum at 303K in C₇D₈; and (c) ¹⁹F{¹H} NMR spectrum at 343K in C₇D₈.^{*}



(a)

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(c)

S5: GC /MS analyses of reaction mixture forming $[Ce(L^{Me})_3]$ and $[CeF(L^{Me})_2]_3$:



File	:C:\msdchem\1\DATA\sally\RK1_73A.D
Operator	: VDG
Acquired	: 28 May 2010 15:56 using AcqMethod gen5sl a.M
Instrument	: Agilent GCMS
Sample Name	: RK1.73a (R. Kelly)
Misc Info	: C16H12F8N2 = 384, C10H13f3N2 = 218
Vial Number	: 11







File :C:\msdchem\1\DATA\sally\RK1_73A.D Operator : VDG Acquired : 28 May 2010 15:56 using AcqMethod gen5sl_a.M Instrument : Agilent GCMS Sample Name: RK1.73a (R. Kelly) Misc Info : C16H12F8N2 = 384, C10H13f3N2 = 218 Vial Number: 11



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File :C:\msdchem\l\DATA\sally\RK1_73A.D
Operator : VDG
Acquired : 28 May 2010 15:56 using AcqMethod gen5sl_a.M
Instrument : Agilent GCMS
Sample Name: RK1.73a (R. Kelly)
Misc Info : C16H12F8N2 = 384, C10H13f3N2 = 218
Vial Number: 11
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File :C:\msdchem\l\DATA\sally\RK1_73A.D Operator : VDG Acquired : 28 May 2010 15:56 using AcqMethod gen5sl_a.M Instrument : Agilent GCMS Sample Name: RK1.73a (R. Kelly) Misc Info : C16H12F8N2 = 384, C10H13f3N2 = 218 Vial Number: 11



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