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# Speciation of lanthanide ions in the organic phase after extraction from nitrate media by basic extractants†

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A speciation study was carried out for lanthanide complexes formed in the organic phase after solvent extraction with quaternary ammonium and phosphonium nitrate extractants. These extractants are liquid at room temperature and were applied in their undiluted form. A comparison was made between the quaternary compound trihexyl(tetradecyl)phosphonium nitrate, the nitrate form of the commercial extractant Cyphos IL 101, and Aliquat 336 nitrate, the nitrate form of the commercial trialkylmethylammonium chloride extractant Aliquat 336 (alkyl = mixture of C8 and C10 chains). The structures of the lanthanide complexes across the entire lanthanide series (with the exception of promethium) were determined by a combination of solvent extraction techniques, FTIR, NMR, high-resolution steady-state luminescence spectroscopy, luminescence life time measurements, elemental analysis and EXAFS spectroscopy. The results suggest that the lanthanide ions form an anionic nitrate complex in the organic phase by coordinating with five bidentate nitrate ligands. Charge neutralization is provided by two counter cations of the extractant present in the outer coordination sphere of the complex. Furthermore, it is suggested that the pentanitrate complex is the sole lanthanide species that is formed in significant concentrations in the organic phase.

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## Introduction

The lanthanides comprise a group of 15 chemical elements ranging from lanthanum ( $Z = 57$ ) to lutetium ( $Z = 71$ ). The 4f orbitals, which are sequentially filled from lanthanum to lutetium, are shielded from the nucleus by the fully occupied and more highly energetic 5s and 5p orbitals.<sup>1</sup> Together with scandium and yttrium, the lanthanides form the group of the rare-earth elements (REEs). Due to their similar electronic structure and ion size, all lanthanide ions have comparable chemical and physical properties. Rare-earth ores contain a mixture of lanthanides which need to be separated in order to isolate the pure metals, which are used in applications such as permanent magnets, lamp phosphors, catalysts and optical glass.<sup>2,3</sup> Industrial separation of lanthanides is done by solvent extraction.<sup>4–7</sup> A metal-rich aqueous phase is brought into contact with an immiscible organic phase, containing an extractant, a diluent and often also a modifier.<sup>8–10</sup> The extractant is a metal-coordinating ligand and the diluent is used to increase the solubility of the complex and to decrease the viscosity of the organic phase. The metals distribute between the two phases

based on their affinity for the aqueous phase (interactions with water and/or the complexing agent) or the organic phase (interactions with the extractant and/or the diluent). In general, extractants are divided in three distinct classes: (1) acidic extractants for which the extraction is pH-dependent, (2) neutral extractants, where neutral species containing electron-donating groups such as oxygen coordinate to the metal ion, and (3) basic extractants which extract anionic metal complexes. Due to their similar chemical properties, the separation between neighbouring lanthanides is difficult and often more than 100 extraction stages are required to obtain one lanthanide in high purity (>99.9%).<sup>11</sup> Acidic extractants are preferred by industry to separate the lanthanides into individual elements because they can extract lanthanides from chloride solutions. This has the advantage that HCl can be used to dissolve the mixed lanthanide oxide starting materials, which is cheaper than  $\text{HNO}_3$ .<sup>11</sup> In the case of basic extractants, only nitrate- or thiocyanate-based extractants show significant lanthanide extraction and are therefore less exploited in industry. Nevertheless, they offer a high potential, such as no need for accurate pH control and less consumption of chemicals due to easy stripping with water. Furthermore, our group has recently provided a solution to extract lanthanides also from chloride solution with ammonium and phosphonium nitrate and thiocyanate extractants, based on a concept called “split-anion extraction”.<sup>12</sup>

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In general, the ionic radius of the lanthanides decreases with increasing atomic number (lanthanide contraction) and the charge density increases, so that the electrostatic interaction between the lanthanide and the extractant present in the organic phase increases across the lanthanide series. If conditions are kept constant, this results in higher extraction efficiencies for the heavy lanthanides compared to the lighter lanthanides. This is the so-called “positive sequence” for solvent extraction. There are a few exceptions to this rule. In the case of extraction with carboxylic acids diluted in apolar solvents, the structure of the complex changes across the lanthanide series. This results in an optimum in the plot of extraction efficiency as a function of the atomic number  $Z$ , as shown by Preston and du Preez.<sup>13,14</sup> 2-Bromo-3,5,5-trimethylhexanoic acid extracts heavy lanthanides as monomeric complexes of the form  $\text{Ln}(\text{A})_3(\text{HA})_3$ , while the light lanthanides are extracted as dimeric  $(\text{Ln}(\text{A})_3(\text{HA})_3)_2$  complexes. An optimum in the extraction efficiency along the lanthanide series is also found for the neutral extractants tri-*n*-octylphosphine oxide (TOPO) and tri-*n*-butylphosphate (TBP) and basic trialkylmethylammonium chloride extractants, *e.g.* Aliquat 336.<sup>11,15,16</sup>

A surprising example is the extraction of lanthanide ions by basic extractants having nitrate anions, such as Aliquat 336 in the nitrate form,  $[\text{A336}][\text{NO}_3]$ , where the highest extraction efficiency is observed for  $\text{La}^{3+}$ , and the lowest for  $\text{Lu}^{3+}$ .<sup>11,16</sup> The reverse extraction behaviour is called the “negative sequence”. In contrast, lanthanide extraction with basic extractants in the thiocyanate form, such as Aliquat 336 thiocyanate  $[\text{A336}][\text{SCN}]$ , follow a normal positive sequence. This difference in behaviour could be exploited to separate the lanthanide ions with solvent extraction, as shown by Larsson and Binnemans.<sup>17</sup> Preston and Du Preez suggested that this remarkable negative sequence might be due to steric hindrance of the nitrate in the lanthanide complex, resulting in lower stability constants for the nitrate complexes of the smaller heavy lanthanides compared to the larger light lanthanides.<sup>16</sup> However, they did not provide experimental proof to support their hypothesis. Attempts to gain more insight into the extraction mechanism focus mainly on the structural analysis of the lanthanide complex in the aqueous phase or on slope analysis.<sup>18–20</sup> To the best of our knowledge, a speciation study of the lanthanide complex in the organic phase is not available for this type of extraction systems yet.

In this paper, the structure of the lanthanide complexes formed upon extraction with basic nitrate extractants is investigated. Two common basic extractants are considered: the quaternary compound trihexyl(tetradecyl)phosphonium nitrate  $[\text{P}_{66614}][\text{NO}_3]$  and the quaternary ammonium nitrate extractant Aliquat nitrate  $[\text{A336}][\text{NO}_3]$ .  $[\text{P}_{66614}][\text{NO}_3]$  is the nitrate form of Cyphos IL 101, whereas  $[\text{A336}][\text{NO}_3]$  is the nitrate form of Aliquat 336 (trialkylmethylammonium chloride, with a mixture of  $\text{C}_8$  and  $\text{C}_{10}$  alkyl chains) (Fig. 1). The different lanthanide complexes are studied by solvent extraction techniques, FTIR, NMR, luminescent life time measurements, high-resolution luminescence spectroscopy, elemental analysis and EXAFS.<sup>21</sup> Promethium was not studied due to the lack of stable isotopes.

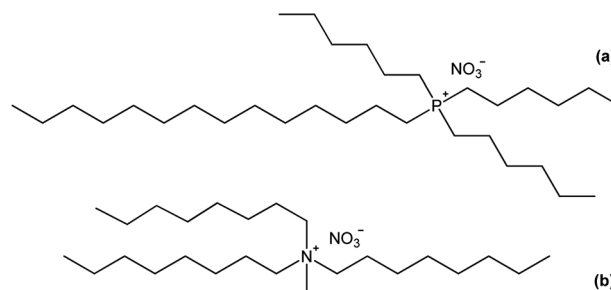


Fig. 1 Chemical structure of (a) trihexyl(tetradecyl)phosphonium nitrate  $[\text{P}_{66614}][\text{NO}_3]$  and (b) trioctylmethylammonium nitrate, the main component of Aliquat 336 in the nitrate form,  $[\text{A336}][\text{NO}_3]$ .

## Experimental

### Chemicals

Praseodymium(III) nitrate hexahydrate (99.9%), samarium(III) nitrate hexahydrate (99.9%), europium(III) nitrate hexahydrate (99.9%), gadolinium(III) nitrate hexahydrate (99.9%), dysprosium(III) nitrate hexahydrate (99.9%), holmium(III) nitrate hydrate (99.9%), ytterbium(III) nitrate pentahydrate (99.9%), lutetium(III) nitrate hydrate (99.9%) and lutetium(III) chloride hexahydrate (99.9%) were obtained from STREM Chemicals (Newburyport, MA, US). Cerium(III) nitrate hexahydrate (99.9%) and terbium(III) nitrate pentahydrate (99.99%), Aliquat® 336 (88–90% purity), ammonium nitrate (99.9%), nitric acid (65%), Triton-X 100 and deuterium oxide (99.9%) were obtained from Sigma-Aldrich (Overijse, Belgium). Lanthanum(III) nitrate hexahydrate (99.9%), neodymium(III) nitrate hexahydrate (99.9%) and thulium(III) nitrate pentahydrate (99.9%) were purchased from Alfa Aesar (Karlsruhe, Germany) and erbium(III) nitrate pentahydrate (99.9%) from GFS Chemicals (Powell, OH, US). Trihexyl(tetradecyl)phosphonium chloride (Cyphos® IL 101,  $[\text{P}_{66614}]\text{Cl}$ ) was purchased from Cytec Industries (Faulquemont, France). ICP standard solutions ( $\text{Ce}$ ,  $\text{Pr}$ ,  $\text{Nd}$ ,  $\text{Sm}$ ,  $\text{Ho}$ ,  $\text{Er}$ ,  $\text{Y}$ :  $1000 \mu\text{g mL}^{-1}$  in 3–5%  $\text{HNO}_3$  and the rare-earth mixture of  $50 \mu\text{g mL}^{-1}$  in 3–5%  $\text{HNO}_3$ ) were obtained from Chem-Lab (Zedelgem, Belgium). Toluene, hydrochloric acid (~37%) and absolute ethanol were obtained from Fisher Scientific (Hampton, NH, US). The silicone solution in isopropanol was ordered from SERVA Electrophoresis (Heidelberg, Germany). Ultrapure water ( $18.2 \text{ M}\Omega \text{ cm}$ ) was used for all dilutions and produced using a Sartorius Arium Pro Ultrapure Water System.

### Extraction experiments

Trihexyl(tetradecyl)phosphonium nitrate,  $[\text{P}_{66614}][\text{NO}_3]$ , and Aliquat 336 nitrate (trialkylmethylammonium nitrate,  $[\text{A336}][\text{NO}_3]$ ) were synthesized *via* a metathesis reaction with  $\text{NH}_4\text{NO}_3$  starting from Cyphos IL 101 and Aliquat 336, respectively. The Cyphos IL 101 and Aliquat 336 were each contacted three times with a  $10 \text{ mol L}^{-1}$   $\text{NH}_4\text{NO}_3$  solution in a 1 : 1 aqueous-to-organic (A/O) volume ratio while stirring vigorously. Subsequently, the organic phase was washed with water to remove the remaining salt impurities. The presence of chloride in the water-saturated organic phase was ruled out by



total-reflection X-ray fluorescence (TXRF) spectroscopy.<sup>22,23</sup> All extraction experiments were performed by mixing an aqueous lanthanide nitrate solution (various concentrations) with water-saturated  $[P_{66614}][NO_3]$  or  $[A336][NO_3]$  in a 1 : 1 A/O volume ratio. Higher lanthanide loadings in the organic phase were obtained by applying higher A/O volume ratios. Extraction equilibrium was reached by shaking the mixtures at 70 °C for 4 h in a TMS-200 Thermoshaker (Nemus Life, Sweden) operating at 1700 rpm. A Heraeus Megafuge 1.0 centrifuge was used to speed up phase disengagement. The glass vials used for extraction were always first pretreated by cleaning with HCl (37%) for 1 h at 60 °C to avoid leaching of metal ions present in the glass (Fe, Pb) during the extractions. The lanthanide concentration in the aqueous phase was measured with inductively coupled plasma optical emission spectroscopy (ICP-OES) and the concentration in the organic phase was calculated using the mass balance. The extraction of the lanthanides was quantified by calculating the distribution ratio  $D_{Ln}$ :

$$D_{Ln} = \frac{[Ln]_{org}}{[Ln]_{aq}} \quad (1)$$

where  $[Ln]_{org}$  and  $[Ln]_{aq}$  are the lanthanide concentration in the organic and aqueous phase, respectively. The separation between two lanthanides is calculated using the separation factor  $\alpha_{Ln_1}^{Ln_2}$ :

$$\alpha_{Ln_1}^{Ln_2} = \frac{D_{Ln_1}}{D_{Ln_2}} \quad (2)$$

### Analysis techniques

The lanthanide concentration in the aqueous phase of the extraction experiments was measured with ICP-OES on a Perkin-Elmer Optima 8300 spectrometer. The samples were diluted 100 times in 2 vol%  $HNO_3$ . Calibration solutions of 0.01 and 1 mg  $L^{-1}$  were prepared by diluting the ICP standard rare-earth mixture (50  $\mu g mL^{-1}$  in 3–5%  $HNO_3$ ) in 2 vol%  $HNO_3$ . Scandium and indium (5 mg  $L^{-1}$ ) were added as internal standard to the samples and calibration solutions to check for matrix effects. Total reflection X-ray fluorescence (TXRF) spectroscopy was performed on a Bruker Picofox S2 spectrometer to determine the chloride content of the synthesized  $[P_{66614}][NO_3]$  and  $[A336][NO_3]$  and to measure the lanthanide concentration in the organic phase for determination of the maximum loading. A 20 mg aliquot was dissolved in 1 mL of absolute ethanol. A 100  $\mu L$  aliquot of this solution was mixed with 100  $\mu L$  of an internal standard solution of a neighbouring lanthanide (1000  $\mu g mL^{-1}$ ) and diluted to 1 mL with ethanol. The quartz sample carriers were pretreated with 30  $\mu L$  of silicone solution in isopropanol and dried for 30 min at 60 °C, prior to addition of the sample on the carrier (1  $\mu L$ ) and again drying for 30 min at 60 °C. The samples were measured for 500 s. The carbon, hydrogen and nitrogen

content (wt%) of the metal loaded  $[P_{66614}][NO_3]$  phase was determined by CHN analysis using a Thermo Scientific Inter-science Flash 2000 elemental analyser. Quantitative  $^{31}P$  NMR was performed to determine the phosphorus content in the metal-loaded  $[P_{66614}][NO_3]$  phase by adding benzyl-triphenylphosphonium chloride as a standard to the sample, diluted in DMSO.  $^{31}P$  NMR spectra were recorded on a Bruker Ascend 400 spectrometer operating at 162 MHz. The d1 parameter of the measurement was optimized to 20 s. The P concentration was calculated based on the integration of the peaks of  $[P_{66614}][NO_3]$  ( $\delta = 33.76$  ppm) and the standard ( $\delta = 23.16$  ppm). Photoluminescence spectroscopy was performed on a Horiba Jobin Yvon Fluorolog-3 spectrofluorimeter equipped with a double-grating monochromator in both the excitation and emission sides, coupled to a R928P Hamamatsu photomultiplier and a 450 W Xe arc lamp as the excitation source. The luminescence lifetimes were measured by a pulsed Xe lamp with variable repetition rate and elaborated with standard software fitting procedures. Fourier-transform infrared spectra were recorded on a Bruker Vertex 70 spectrometer and analysed with the Bruker OPUS software (version 7.5). The liquid samples were measured directly, without sample preparation, using the attenuated total reflectance module (Platinum ATR). Extended X-ray Absorption Fine Structure (EXAFS) spectra were collected at the Dutch-Belgian Beamline (DUBBLE, BM26A) of the European Synchrotron Radiation Facility (ESRF) in Grenoble (France). The energy of the X-ray beam was tuned by a double-crystal monochromator operating in fixed-exit mode using a Si(111) crystal pair for measurements on the  $L_{III}$ -lines of Nd, Sm, Eu, Gd, Tb, Dy, Ho, Er, Tm, Yb and Lu and a Si(311) crystal pair for measurements on the K-edge of La (38 925 eV) and Ce (40 443 eV). The measurements were done in transmission mode using Ar/He gas filled ionization chambers at ambient pressure for the  $L_{III}$  edges and Ar/Kr filled ionization chambers for the K edges. A brass sample holder with Kapton windows and a flexible polymeric spacer (VITON) was used. Standard procedures were applied for pre-edge subtraction and data normalization in order to isolate the EXAFS function ( $\chi$ ). The EXAFS oscillations, isolated by a smoothing spline using the program VIPER, were  $k^4$ -weighed and Fourier transformed using a Gaussian rounded ends window function.<sup>24</sup> The data were fitted in  $R + \Delta$  ( $\text{\AA}$ ) space using the *ab initio* code FEFF 7.0, which was used to calculate the theoretical phase and amplitude functions of the model that was subsequently used in the non-linear least-squares refinement of the experimental data.<sup>25</sup> The input used for FEFF 7.0 was manually constructed and consisted of the targeted lanthanide in the centre position, with one bidentate nitrate in its first coordination sphere at a distance corresponding to lanthanide pentanitrato crystal structures.<sup>26</sup> Estimated errors are shown between parentheses and calculated by VIPER. The amplitude reduction factor  $S_0$  was fixed at 0.9 for the La and Ce samples and at 1.1 for all other samples.  $[P_{66614}]_3[LuCl_6]$  was used as a reference compound and was prepared by dissolving 100 mg of  $LuCl_3 \cdot 6H_2O$  in 5 mL Cyphos IL 101 and drying at a Schlenk line for two days at 60 °C.



## Results and discussion

### Extraction

The distribution ratios for the trivalent lanthanide ions are known to show a decreasing trend across the lanthanide series upon extraction with the extractants  $[P_{66614}][NO_3]$  and  $[A336][NO_3]$  from a nitrate feed solution.<sup>11,16</sup> As shown in Fig. 2, the distribution ratios indeed decreased when going from low atomic numbers (La,  $Z = 57$ ) to high atomic numbers (Lu,  $Z = 71$ ), confirming the negative sequence. The decreasing trend was more pronounced at high  $NH_4NO_3$  concentrations in the aqueous phase. At high nitrate concentrations, the distribution ratios showed a clear tetrad effect.<sup>27,28</sup> This behaviour is typically seen in several lanthanide extraction system and divides the lanthanides into four tetrads, with Gd ( $Z = 64$ ) being the common element of the two central tetrads. Furthermore, it can be seen that the distribution ratios increased with an increasing nitrate concentration in the aqueous phase, with a larger increase for the lanthanides with low atomic number compared to the ones with a high atomic number. Consequently, the slope of the plot changed with increasing nitrate concentration, resulting in improved selectivity for extraction of the lanthanides with a low atomic number over those with a high atomic number, as a function of the nitrate concentration.

### Systematic research in the CSD

A systematic research of the Cambridge Structural Database (CSD) was performed to get a first idea of the coordination of nitrate to the extracted lanthanide ions in the organic phase. Lanthanide crystal structures containing at least four bidentate nitrate ligands, including structures with five and six nitrates and/or other ligands, were extracted from the database. An overview of the different structures is presented in the ESI (Fig. S2†). One must be aware of the fact that crystal structures are not necessarily representative for the structure of the complexes in the liquid state. For  $La^{3+}$ ,  $Ce^{3+}$ ,  $Pr^{3+}$  and  $Nd^{3+}$ , the

majority of the isolated nitrate complexes contained six bidentate nitrate ligands with a coordination number of twelve. For  $Sm^{3+}$ ,  $Eu^{3+}$ ,  $Gd^{3+}$  and  $Dy^{3+}$  complexes with six nitrate ions have been isolated, but the number of complexes with four or five nitrates was larger. For  $Tb^{3+}$ ,  $Ho^{3+}$ ,  $Er^{3+}$ ,  $Yb^{3+}$ ,  $Tm^{3+}$  and  $Lu^{3+}$ , only complexes with four or five nitrate ions have been reported. In general, the lighter lanthanides mostly occur as hexanitrate complexes,  $[Ln(NO_3)_6]^{3-}$ , while the heavier lanthanides mainly form pentanitrate complexes  $[Ln(NO_3)_5]^{2-}$ .

### Fourier-transform infrared (FTIR) spectroscopy

FTIR spectroscopy was used to determine the mode in which the nitrate anions coordinate to the extracted trivalent lanthanide ion in the organic phase. Nitrate can coordinate to a metal ion in three different ways: monodentate, bidentate bridging and bidentate chelating (Fig. 3).

The transmittance of the organic phase was recorded in the range between 4000 and 400  $cm^{-1}$  as a function of the  $Pr^{3+}$ ,  $Nd^{3+}$ ,  $Ho^{3+}$  and  $Er^{3+}$  loading in the organic phase. Similar bands were observed for all four lanthanides. Therefore, the spectrum of  $Nd^{3+}$  is discussed in more detail. The full spectrum of  $Nd^{3+}$  can be found in the ESI (Fig. S3†). Bands corresponding to the vibration modes of the cation of the extractant did not show an increase in intensity or shift in wavenumber upon metal loading, indicating that the extractant cation is not directly coordinating to the metal centre. The vibration modes of the cation include the  $CH_2$  stretching mode in the region 2850–2950  $cm^{-1}$  and the  $CH_2$  rocking mode at 719  $cm^{-1}$  (Fig. 4). A band attributed to the bending mode of  $H_2O$  can be found at 1630  $cm^{-1}$ . The intensity of this band decreased with increasing metal loading, due to the decrease of the water content in the organic phase at higher metal loadings. This decrease in water content is attributed to the formation of an anionic lanthanide complex with a lower charge density than the nitrate ion, leading to a weaker anion hydration in the organic phase. The majority of the remaining bands can be attributed to the vibration of the nitrate anion.<sup>29–34</sup> A combination band of the symmetrical stretch and in-plane bending of the nitrate can be found in the region 1730–1770  $cm^{-1}$ . This band is split upon loading of the organic phase with  $Nd^{3+}$  into two bands, at 1770  $cm^{-1}$  and 1720  $cm^{-1}$ . This is attributed to the reduction of the local symmetry of the nitrate anion from  $D_{3h}$  in the non-coordinated form, to  $C_{2v}$  or lower for the metal-coordinated nitrate. The splitting of this doubly degenerate combination band into two separate bands is a measure for the coordination mode. In this case, the size of the gap was 43  $cm^{-1}$ , which is an

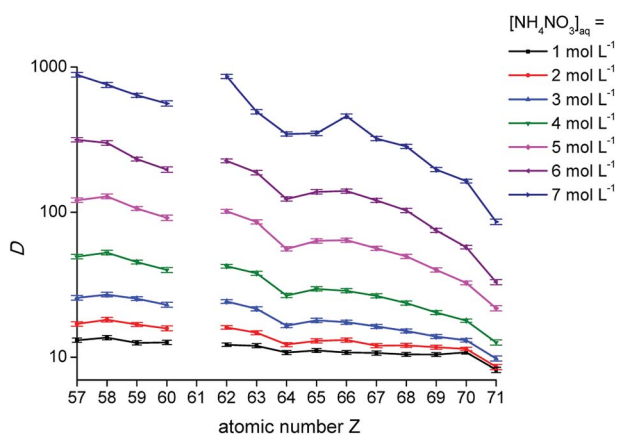


Fig. 2 Distribution ratios ( $D$ ) as a function of the atomic number of the lanthanides for the extraction with  $[P_{66614}][NO_3]$  for different  $NH_4NO_3$  concentrations in the aqueous phase.  $[Ln^{3+}]_{aq,in} = 0.01 \text{ mol L}^{-1}$ , A/O 1 : 1.

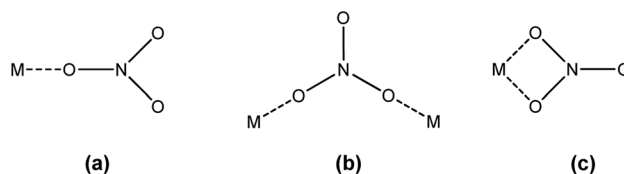


Fig. 3 Different coordination modes of the nitrate ion: (a) monodentate, (b) bidentate bridging and (c) bidentate chelating.



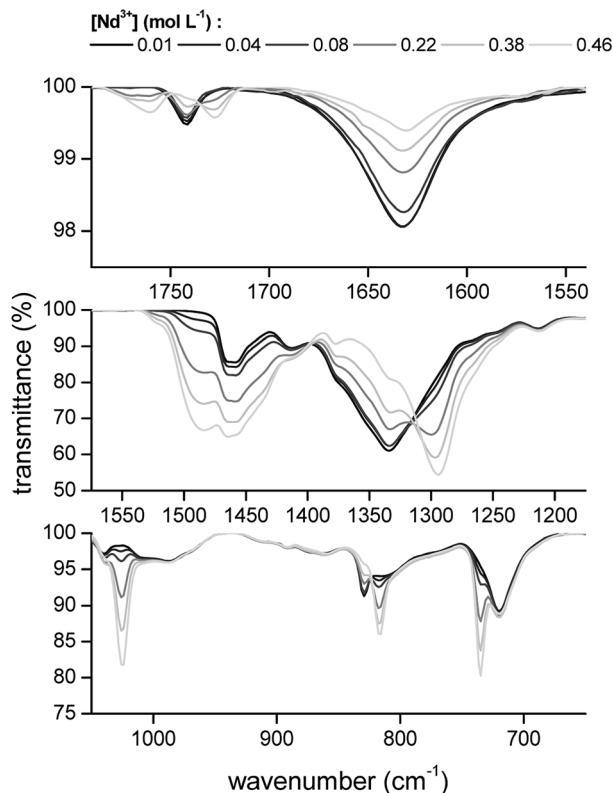


Fig. 4 FTIR spectra of the  $[P_{66614}][NO_3]$  phase after extraction of increasing concentrations of  $Nd^{3+}$  from a nitrate solution.

indication for the bidentate coordination of the nitrate anion to the metal cation. A monodentate coordination would be characterized by a gap in the order of only  $10\text{ cm}^{-1}$ .<sup>30</sup> Additionally, a comparison was made with the spectra of  $Er^{3+}$  and  $Ho^{3+}$  at the highest loading (Fig. 5). Both bands were shifted towards lower wavenumbers with increasing atomic number of the lanthanide. This might be explained by a weakening of the O–N bond in the nitrate ligand due to an increase in electron transfer to the  $Ln^{3+}$  metal ion with increasing charge density. Furthermore, two bands appear in the spectrum of  $Nd^{3+}$  upon increasing

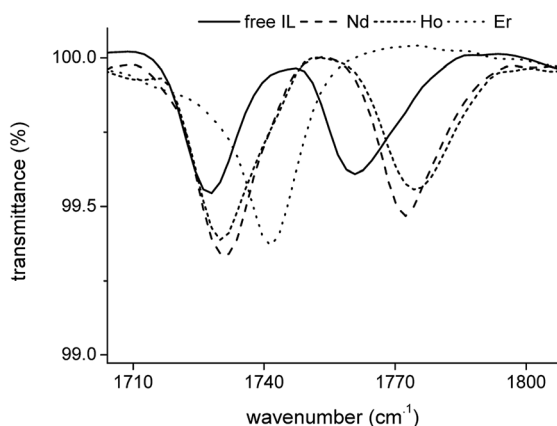


Fig. 5 FTIR spectra of the fully loaded  $[P_{66614}][NO_3]$  phase after extraction of  $Nd^{3+}$ ,  $Ho^{3+}$  and  $Er^{3+}$  from a nitrate solution.

loading, at  $1150\text{ cm}^{-1}$  and  $725\text{ cm}^{-1}$  and their intensity increased with increasing metal loading (Fig. 4). They correspond to the symmetrical stretching and symmetrical bending of  $NO_2$  respectively and are not IR active in the free, non-coordinated nitrate ligand. Upon coordination, the symmetry of the nitrate decreases and they become IR active.<sup>29</sup> The band at  $1335\text{ cm}^{-1}$  was shifted to  $1295\text{ cm}^{-1}$  upon coordination and was attributed to the asymmetric stretching of  $NO_2$ . The band of the out-of-plane rocking of  $NO_2$  appeared at  $829\text{ cm}^{-1}$  in the non-coordinated nitrate and is shifted to  $816\text{ cm}^{-1}$  upon loading of the organic phase. The band in the region  $1464\text{--}1480\text{ cm}^{-1}$  was attributed to the N–O stretching vibration. Finally, several isosbestic points were observed in the spectra upon increasing metal loading, e.g. at  $825$ ,  $1333$  and  $1392\text{ cm}^{-1}$ . An isosbestic point indicates the presence of one single equilibrium between two species, in this case the non-coordinated and the coordinated nitrate ligand. Consequently, only one species is formed upon coordination of nitrate to the extracted lanthanide ion at high metal loadings. Notice that at low metal loadings, differences in the FTIR spectra were rather difficult to distinguish. Thus, based on these results one must be careful when drawing conclusions about the number of different lanthanide-nitrate species present at low metal loadings.

### Photoluminescence spectroscopy

Complementary to FTIR,  $Eu^{3+}$  luminescence emission spectra were recorded to gain insight into the local symmetry and structure of the  $Eu^{3+}$  nitrate complex in the organic phase. A comparison was made between  $[P_{66614}][NO_3]$  and  $[A336][NO_3]$  and between the water-saturated and the dried organic phase. Furthermore, the influence of the metal loading in the organic phase on the spectrum was investigated. All spectra showed the same transitions and relative intensities, both in the  $Eu^{3+}$  loaded  $[A336][NO_3]$  and  $[P_{66614}][NO_3]$  and as a function of the  $Eu^{3+}$  loading (Fig. 6). This indicates that the same species is

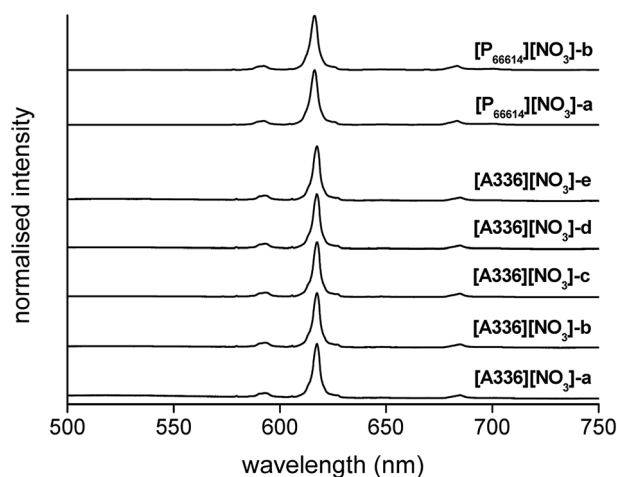


Fig. 6 Emission spectra ( $\lambda_{ex} = 464\text{ nm}$ ) of  $[A336][NO_3]$  loaded with increasing concentrations of  $Eu^{3+}$  ( $0.1$  to  $0.3\text{ mol L}^{-1}$ , sample  $[A336][NO_3]$ -a to  $[A336][NO_3]$ -e), and  $[P_{66614}][NO_3]$  loaded with  $Eu^{3+}$  ( $0.5\text{ mol L}^{-1}$ ), water-saturated ( $[P_{66614}][NO_3]$ -a) and dry ( $[P_{66614}][NO_3]$ -b), by extraction from a nitrate solution.



present at both low and high metal loadings and in both organic phases, which confirms that the cation of the studied extractants did not affect the speciation. Furthermore, the  $\text{Eu}^{3+}$  emission spectrum in dried  $[\text{P}_{66614}][\text{NO}_3]$  is the same as in water-saturated  $[\text{P}_{66614}][\text{NO}_3]$ . Most likely, no water molecules are directly coordinated to the  $\text{Eu}^{3+}$  ion, assuming that all water is evaporated from the sample. The  $^5\text{D}_0 \rightarrow ^7\text{F}_0$  transition is very weak and shows only one peak, indicating that only one species is present. Due to the non-degeneracy of the  $^5\text{D}_0$  and  $^7\text{F}_0$  levels, only one peak is expected for a single species in solution. The peak for the  $^5\text{D}_0 \rightarrow ^7\text{F}_1$  transition is weak and split. In an icosahedral environment, this transition is not split, while in a hexagonal, tetragonal and trigonal symmetry, it is split into one non-degenerate and two degenerate levels.<sup>35</sup> The hexanitrate complex ( $\text{CN} = 12$ ) is expected to have an octahedral or icosahedral-like geometry, while the pentanitrate complex ( $\text{CN} = 10$ ) would show a trigonal bipyramidal type of symmetry.<sup>36</sup> The hypersensitive transition  $^5\text{D}_0 \rightarrow ^7\text{F}_2$  is very intense and sharp, which indicates that the  $\text{Eu}^{3+}$  complex does not contain an inversion centre. The high intensity of this transition thus indicates the presence of a trigonal bipyramidal pentanitrate europate(III) complex with bidentate coordinating nitrate ligands in a  $D_{3h}$  symmetry. Furthermore, the fact that this transition shows a very narrow peak and is not split gives additional evidence for a  $D_{3h}$  symmetry. The  $^5\text{D}_0 \rightarrow ^7\text{F}_3$  transition is very weak and the  $^5\text{D}_0 \rightarrow ^7\text{F}_4$  transition is weak. For five bidentate nitrate ions to be compatible with a trigonal bipyramid ( $\text{CN} = 5$ ) as an idealized coordination polyhedron, each nitrate ion must be considered to occupy one of the five coordination places in the trigonal bipyramid and the total coordination number of the europium complex would be  $\text{CN} = 10$  (Fig. 7).

The hydration number  $q$  of the metal complex in the organic phase was determined using luminescence lifetime decay measurements of  $\text{Eu}^{3+}$  and  $\text{Tb}^{3+}$ . The luminescence decay curves were recorded for the loaded organic phase in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  medium, at low and high metal loadings,  $0.05 \text{ mol L}^{-1}$  and  $0.5 \text{ mol L}^{-1}$ , respectively (Fig. S4–S6†). All curves were fitted well

**Table 1** Lifetimes of  $\text{Eu}^{3+}$  and  $\text{Tb}^{3+}$  luminescence in the organic phase by extraction with  $[\text{P}_{66614}][\text{NO}_3]$  in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$

Sample	$\tau_{\text{H}_2\text{O}}$ (ms)	$\tau_{\text{D}_2\text{O}}$ (ms)	$q$
$\text{Eu}^{3+}$ $0.05 \text{ mol L}^{-1}$	1.13	1.26	$0.1 \pm 0.5$
$\text{Tb}^{3+}$ $0.05 \text{ mol L}^{-1}$	1.30	1.36	$0.1 \pm 0.5$
$\text{Tb}^{3+}$ $0.5 \text{ mol L}^{-1}$	1.43	1.44	$0.0 \pm 0.5$

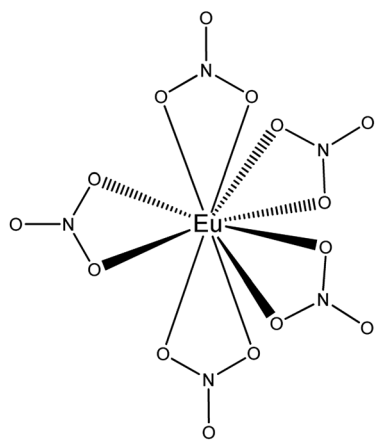
by a mono-exponential decay function, indicating that each sample contained only one species. The calculated lifetimes (ms) are presented in Table 1. The hydration number  $q$  was calculated using the Horrocks–Sudnick formula (3):

$$q = A \left[ \frac{1}{\tau_{\text{H}_2\text{O}}} - \frac{1}{\tau_{\text{D}_2\text{O}}} \right] \quad (3)$$

with  $\tau_{\text{H}_2\text{O}}$  and  $\tau_{\text{D}_2\text{O}}$  the luminescence lifetimes in the  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  sample, respectively, and  $A$  equal to 1.05 (molecules ms) for  $\text{Eu}^{3+}$  and 4.2 (molecules ms) for  $\text{Tb}^{3+}$ .<sup>35,37</sup> Supkowski and Horrocks derived a modified formula, based on literature data of several  $\text{Eu}^{3+}$  complexes in solution, which takes into account additional quenching of the  $^5\text{D}_0$  state of  $\text{Eu}^{3+}$  by other interactions apart from the coordinated water molecules in the first coordination sphere, including bulk water in the sample.<sup>38</sup> However, the samples studied in this work are ionic liquid samples, hence they do not contain water in the bulk. Therefore, the hydration number in these samples is correctly estimated by the original Horrocks–Sudnick equation.

### Maximum loading

Further evidence for the exact stoichiometry of the lanthanide nitrate complex in the organic phase was provided by analyzing the elemental composition of the organic phase at maximum loading. First, the conditions required for maximum loading of the organic phase were estimated by contacting  $[\text{P}_{66614}][\text{NO}_3]$  in a 4 : 1 A/O phase ratio with an aqueous solution containing an excess of lanthanide ( $0.5 \text{ mol L}^{-1}$ ) at varying nitrate concentrations (between 0 and  $8 \text{ mol L}^{-1}$ ).  $\text{La}^{3+}$  and  $\text{Ho}^{3+}$  were arbitrarily selected as representatives for the light and heavy lanthanides, respectively. A comparison between  $\text{Ca}(\text{NO}_3)_2$  and  $\text{NH}_4\text{NO}_3$  was made to evaluate the influence of the salting-out effect of the cation. The results are summarized in Fig. 8. The theoretical lanthanide concentration in the organic phase in case of pentanitrate complex formation was calculated to be approximately  $0.7 \text{ mol kg}^{-1}$ . For the formation of hexanitrate complexes, the concentration would be around  $0.5 \text{ mol kg}^{-1}$ . The loading curve of  $\text{La}^{3+}$  extracted from  $\text{NH}_4\text{NO}_3$  does not approach the line of either the penta- or hexanitrate complex which means that  $\text{NH}_4\text{NO}_3$  is not a good salting-out agent to fully extract the lanthanide ions. Ammonium ions have only weak salting-out or even salting-in properties.<sup>39</sup> Therefore, the results were compared with extraction from  $\text{Ca}(\text{NO}_3)_2$  solution, since  $\text{Ca}^{2+}$  is a much better salting-out agent. The metal loading in the organic phase indeed increased significantly. A plateau value was reached for  $\text{Ho}^{3+}$ , but not for  $\text{La}^{3+}$ , indicating that a maximum might not be reached for  $\text{La}^{3+}$ . The maximum loading of  $\text{Ho}^{3+}$  reached the theoretical value of the formation



**Fig. 7** Schematic representation of the trigonal bipyramidal pentanitrateeuropium complex  $\text{Eu}(\text{NO}_3)_5^{2-}$ . The bipyramid is formed by the position of the nitrogen atoms.



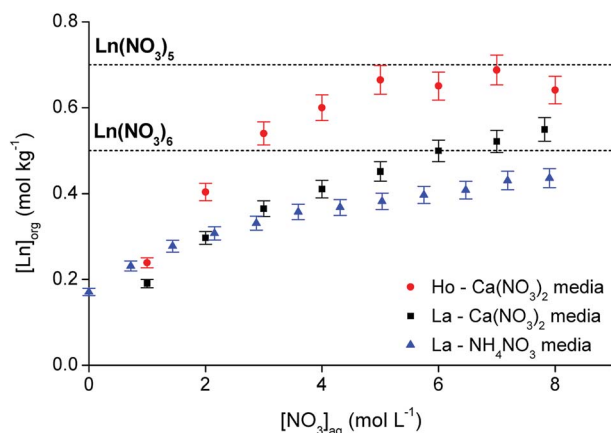


Fig. 8 Loading of  $[P_{66614}][NO_3]$  by extraction of  $Ho^{3+}$  and  $La^{3+}$  from a  $Ca(NO_3)_2$  and  $NH_4NO_3$  solution as a function of the nitrate concentration. 4 : 1 A/O phase ratio,  $[Ln^{3+}]_{aq,in} = 0.5 \text{ mol L}^{-1}$ ,  $[NO_3]_{aq} = 0\text{--}8 \text{ mol L}^{-1}$ . The two dashed lines indicate the theoretical maximum loading in case of  $Ln(NO_3)_5^{2-}$  or  $Ln(NO_3)_6^{3-}$  complexes.

of a pentanittrato complex, while the loading of  $La^{3+}$  was slightly higher than the theoretical value for the formation of hexanittrato complexes. In any case, the results show that conditions close to maximum loading of both lanthanide ions were reached by salting-out with a  $Ca(NO_3)_2$  solution with a concentration higher than  $3 \text{ mol L}^{-1}$ .

Besides  $La^{3+}$  and  $Ho^{3+}$ , also the extraction of  $Nd^{3+}$ ,  $Pr^{3+}$ ,  $Eu^{3+}$ ,  $Er^{3+}$  and  $Lu^{3+}$  with  $[P_{66614}][NO_3]$  was characterized at maximum loading conditions ( $3.5 \text{ mol L}^{-1} Ca(NO_3)_2$ ). The composition of the dried organic phase after extraction was determined by three techniques: TXRF for the lanthanide concentration, CHN elemental analysis for the wt% carbon, hydrogen and nitrogen in the sample and quantitative  $^{31}P$  NMR for determining the relative ratio P/Ln. All these results were compared with the theoretical values of both the formation of lanthanide pentanittrato and lanthanide hexanittrato in the organic phase (Table S1† and Fig. 9). The experimental P/Ln ratio was close to a value of 2 for all tested lanthanides, except for  $La^{3+}$  (2.7). A P/Ln ratio of 2 means that two extractant cations are required to neutralize the lanthanide nitrate complex. Thus, the lanthanide should be coordinated by five nitrate ligands. It is likely that the higher ratio found for La should not be explained by the formation of a nitrato complex with a higher charge, but by the fact that the organic phase was not maximally loaded with  $La^{3+}$ . CHN analysis showed that the mass fraction of carbon and hydrogen are very close to the theoretical value of a pentanittrato complex. The mass fraction of nitrogen is scattered between the theoretical value for the pentanittrato and hexanittrato complexes.

## EXAFS

EXAFS was used as a complementary technique to confirm the presence of lanthanide pentanittrato complexes  $[Ln(NO_3)_5]^{2-}$  in the organic phase. X-ray absorption spectra were recorded on the  $L_{III}$ -edge of the lanthanides Nd to Lu, and on the K-edge of La and Ce. The EXAFS function was extracted from the data and Fourier transformed to obtain the pseudo-radial distribution

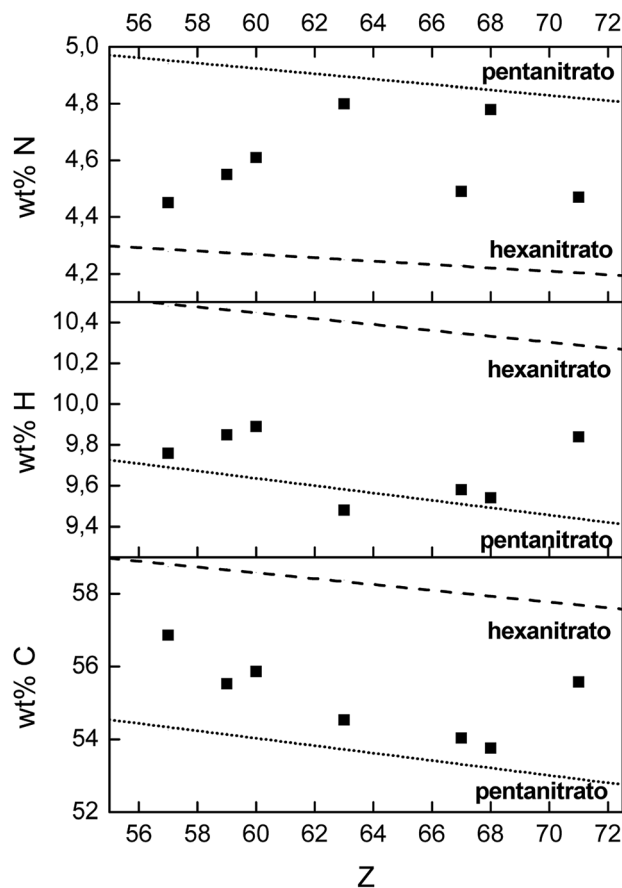


Fig. 9 CHN elemental analysis of the dried  $[P_{66614}][NO_3]$  phase, after full loading by extraction of  $La^{3+}$ ,  $Pr^{3+}$ ,  $Nd^{3+}$ ,  $Eu^{3+}$ ,  $Ho^{3+}$ ,  $Er^{3+}$  and  $Lu^{3+}$  from a  $3.5 \text{ mol L}^{-1} Ca(NO_3)_2$  solution (4 : 1 A/O phase ratio,  $[Ln^{3+}]_{aq,in} = 0.5 \text{ mol L}^{-1}$ ). The dashed line and the dotted line indicate the theoretical value of hexanittrato and pentanittrato complexes, respectively.

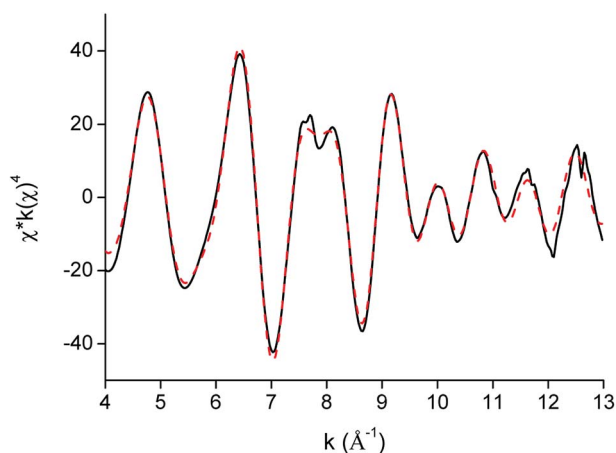
function. The Fourier transform consisted of two shells, at around  $R + \Delta = 2 \text{ \AA}$  and  $4 \text{ \AA}$ , respectively. The first shell, closest to the lanthanide centre, corresponded to the coordinating oxygen atom  $O_c$  and the nitrogen atom N of the nitrate ligand, and is mainly formed by the single scattering paths  $Ln-O_c$  and  $Ln-N$ . The second shell corresponded to the third, non-coordinating, oxygen atom of the nitrate ligand,  $O_{nc}$  and contained contributions of the  $Ln-O_{nc}$  single scattering path, the three-leg scattering path  $Ln-O_{nc}-N$  and the four-leg scattering path  $Ln-O_{nc}-N-O_{nc}$ . The Fourier transform was first fitted using a model based on one bidentate nitrate ligand coordinated to the lanthanide center and the five scattering paths described above,  $Ln-O_c$ ,  $Ln-N$ ,  $Ln-O_{nc}$ ,  $Ln-O_{nc}-N$  and  $Ln-O_{nc}-N-O_{nc}$ . The results for the Lu sample are summarized in Table 2, Fig. 10 and 11.

The data quality of the lighter lanthanides is lower than that of lutetium, due to the lower energies of the  $L_{III}$ -edge (Fig. 12 and Fig. 13). This complicates obtaining a good fit of the second coordination shell in the Fourier transform. However, the  $Ln-O$  and  $Ln-N$  bond distances of the coordinating oxygen atoms and the nitrogen atom of the nitrate

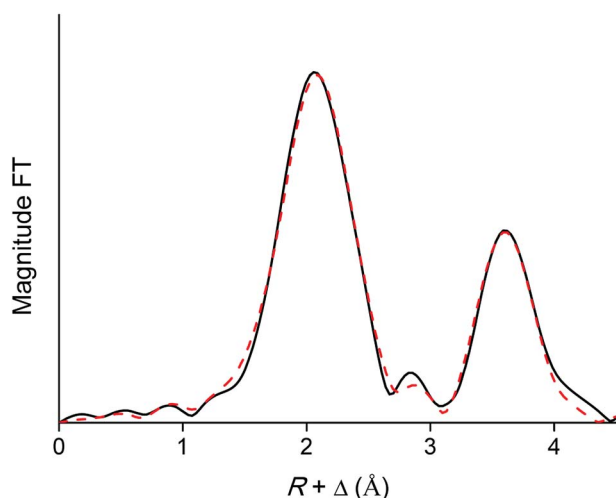


**Table 2** Fitting results of the  $[\text{Lu}(\text{NO}_3)_5]^{2-}$  complex in the organic phase obtained by fitting the five-path model to the two coordination shells of the Fourier transform, between  $R + \Delta = 0$  and  $4.46 \text{ \AA}$ . Errors are mentioned in between brackets

Scattering path	$N$	$r \text{ (\AA)}$	$\sigma^2 \text{ (\AA}^2\text{)}$
Lu–O <sub>c</sub>	9.9(2)	2.395(2)	0.011(1)
Lu–N	5.0(2)	2.852(4)	0.008(1)
Lu–O <sub>nc</sub>	5.0(2)	4.049(3)	0.008(1)
Lu–N–O <sub>nc</sub>	9.9(2)	4.049(3)	0.008(1)
Lu–N–O <sub>nc</sub> –N	9.9(2)	4.059(3)	0.008(1)

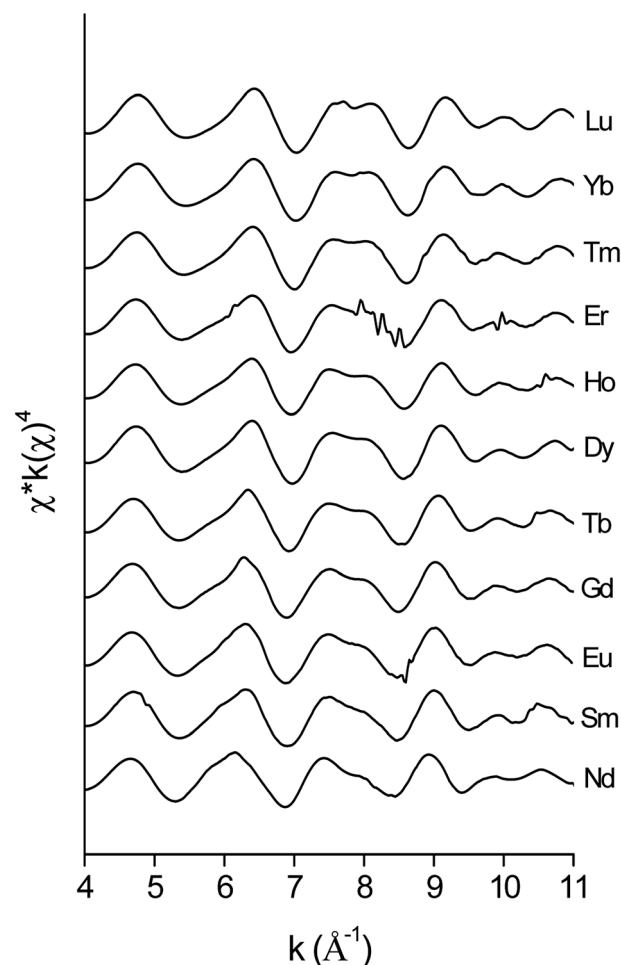


**Fig. 10** EXAFS function (black) and five-path model (red dotted) of the  $[\text{Lu}(\text{NO}_3)_5]^{2-}$  complex extracted to the organic phase.



**Fig. 11** Fourier transform (black) and model (red dotted) of the EXAFS function of the  $[\text{Lu}(\text{NO}_3)_5]^{2-}$  complex extracted to the organic phase. The five-path model was fitted to the Fourier transform between  $R = 0$  and  $4.46 \text{ \AA}$ .

ligands, together with the coordination number can be accurately deduced by fitting only the first coordination shell of the Fourier transform (Fig. 13), using only the single scattering paths Ln–O<sub>c</sub> and Ln–N. The fitting results of all measured lanthanides are summarized in Table S2.† No data are



**Fig. 12** EXAFS functions, recorded on the  $L_{\text{III}}$ -edge of the  $[\text{Ln}(\text{NO}_3)_5]^{2-}$  complexes of various lanthanides in the organic phase.

available for Pr, since the signal-to-noise ratio of the Pr data set was too poor for a correct analysis due to limitations of the beam line. For the heavy lanthanides, a coordination number of 10 was found, while the fitting results of the lighter lanthanides suggest the higher coordination number of 12.

The fitted coordination number is highly dependent on the amplitude reduction factor  $S_0$ ,<sup>40</sup> which was set arbitrarily to 1.1 for all lanthanides, except La and Ce, based on the EXAFS fitting results of the  $[\text{P}_{66614}]_3[\text{LuCl}_6]$  reference compound. Since La and Ce were measured on the K-edge,  $S_0$  was arbitrarily set to 0.9. Consequently, the fitted coordination number is not a good reflection of the number of nitrate ligands coordinated to the lanthanide center. On the other hand, the Ln–O<sub>c</sub> and Ln–N bond distances also provide information on the coordination number. Therefore, the fitted bond distances were compared to literature values of crystal structures of pentanitrate and hexanitrate lanthanide complexes (Table S3†).<sup>26,41–49</sup> The crystal structures were extracted from the Cambridge Structural Database (CSD). Average distances and their corresponding standard deviations were calculated with the Mercury software. A graphical representation of the experimental and literature data is presented in Fig. 14. For all measured lanthanide ions except La, our experimental values corresponded best to the bond distances of



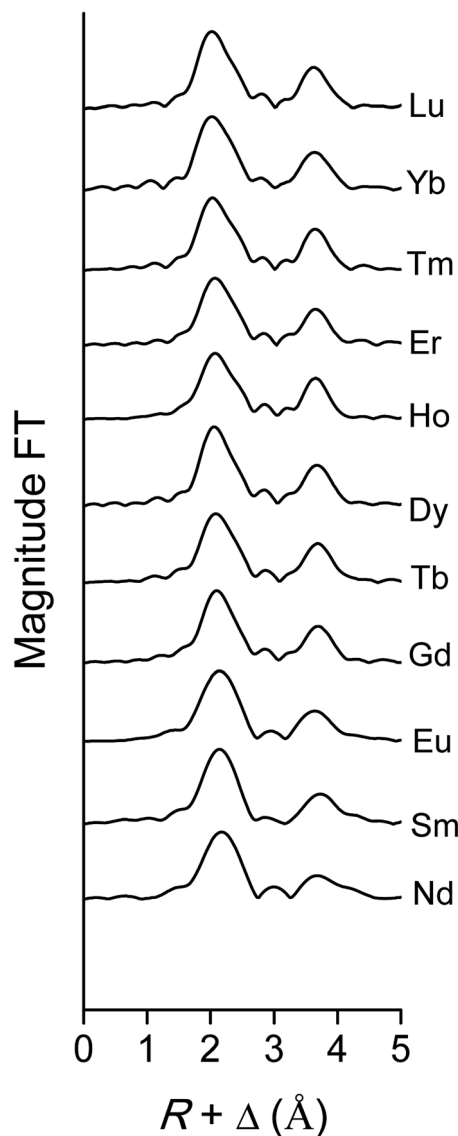


Fig. 13 Fourier transform of the EXAFS function, recorded on the  $L_{III}$  edge, of the  $[Ln(NO_3)_5]^{2-}$  complexes of various lanthanides in the organic phase.

pentanitrato crystal structures. La seems to deviate from this general trend. The value of both the La–O<sub>c</sub> and La–N distances are close to the literature value of the hexanitrato complex. Unfortunately, a crystal structure containing a lanthanum pentanitrato moiety could not be found in the CSD.

### Combination of the results of the different techniques

A systematic review of the crystal structures in the Cambridge Structural Database indicated that the heavy lanthanides generally tend to form pentanitrato complexes in the solid form, while the majority of the reported complexes with the lighter lanthanides are hexanitrato complexes. However, the crystal structures do not necessarily reflect the coordination in liquid form. Lanthanide speciation after extraction to the organic phase could be different from what is observed in crystal structure. Unfortunately, no single analysis technique allows to

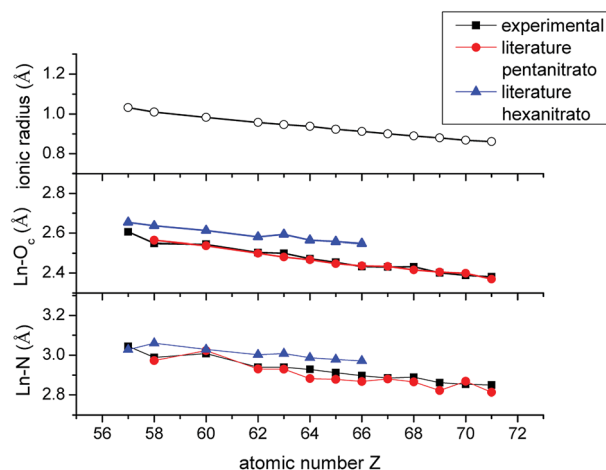


Fig. 14 Comparison between the ionic radius of the trivalent lanthanide ions  $Ln^{3+}$  (coordination number 6), the Ln–O<sub>c</sub> and Ln–N interatomic distances determined experimentally with EXAFS and the corresponding literature values of  $[Ln(NO_3)_5]^{2-}$  and  $[Ln(NO_3)_6]^{3-}$  crystal structures. Error bars are not shown since they are too small. A detailed list of errors can be found in the ESI (Table S2†).

unambiguously determine the exact speciation of all lanthanides in the organic phase and a combination of complementary analytical techniques was applied. FTIR indicated that the cation of the extractant has little effect on the speciation of  $Nd^{3+}$  (light lanthanide), so the cation does not intrude the first coordination sphere of the lanthanide ion. The wavenumber, shifts and splitting patterns of the absorption bands in the FTIR spectra indicate that all nitrate ions coordinate to the lanthanide ions in a bidentate fashion. Furthermore, it was shown that only one species is formed at high metal loadings, as was evident from the presence of several isosbestic points in the FTIR spectra. No conclusions could be drawn on the number of species at low loadings since the differences in intensities were too small. The FTIR spectra of  $Ho^{3+}$  and  $Er^{3+}$ , both heavy lanthanides, showed similar features as  $Nd^{3+}$ . This is a first indication that the speciation of the light and heavy lanthanides is rather similar. Photoluminescence spectroscopy provided further evidence for the presence of only one species at low loadings. The  $^5D_0 \rightarrow ^5F_0$  emission line of the  $Eu^{3+}$  nitrate complex was not split, and the same emission spectrum was recorded for both high and low  $Eu^{3+}$  loadings. Furthermore,  $Eu^{3+}$  and  $Tb^{3+}$  emission lifetime measurements showed a mono-exponential decay. This is a clear indication that only one species is present in the organic phase, regardless of the lanthanide concentration. The comparison between the  $[P_{66614}][NO_3]$  and  $[A336][NO_3]$  emission spectra confirmed that the cation of the extractant has little influence on the speciation of the lanthanide ion. Furthermore, very similar spectra were observed for both wet and dry samples. This indicated that removal of water had no influence on the speciation of the lanthanide and that no water molecules are coordinating to the lanthanide ion. From the lifetime measurements, the hydration number of  $Eu^{3+}$  and  $Tb^{3+}$  was calculated to be close to zero, again confirming that no water is coordinated to the lanthanide ion. From the intensities and the splitting pattern of the  $Eu^{3+}$



emission lines, it was elucidated that  $\text{Eu}^{3+}$  is most probably present as a bidentate pentanitrate complex in the organic phase, which is an important conclusion. Unfortunately, this conclusion applies to  $\text{Eu}^{3+}$  only and cannot simply be extrapolated to the other lanthanides. Elemental analysis (CHN analysis, quantitative  $^{31}\text{P}$  NMR and TXRF analysis) of the organic phase at maximum loading by extraction further confirmed the presence of a pentanitrate complex for the lanthanides Pr, Nd, Ho, Er and Lu. The ratio P/Ln was very close to a value of 2. The pentanitrate complex would have a charge of  $-2$  and should thus be neutralized by two phosphonium cations. The relative carbon, hydrogen and nitrogen contents of the maximally loaded organic phases were close to the theoretical values of the pentanitrate complex  $[\text{P}_{66614}]_2[\text{Ln}(\text{NO}_3)_5]$ . On the other hand, the results on La were inconclusive, probably because maximum loading was not achieved. EXAFS measurements provided confirmation that the pentanitrate complex is formed in the organic phase for all lanthanides, except La. A large systematic error exists for the measured number of scattering paths (neighbours) due to the arbitrary estimation of the amplitude reduction factor. Therefore, the measured bond distances are a much better indication of the coordination number of the complex. These match best with the reported bond distances for pentanitrate complexes for all tested lanthanides except La. Furthermore, the measured bond distances confirmed that all nitrate ligands coordinate in a bidentate fashion. Duvail and coworkers suggested a Ln–N distance of 3.70 Å and 3.57 Å for  $\text{Nd}^{3+}$  and  $\text{Dy}^{3+}$  respectively in case of monodentate coordination and 3 Å and 2.88 Å in case of bidentate coordination.<sup>19</sup> The latter corresponds best to our measured distances of 3.008 Å and 2.897 Å for  $\text{Nd}^{3+}$  and  $\text{Dy}^{3+}$  respectively.

As indicated in the above discussion, it is not possible to draw a final conclusion about the stoichiometry of lanthanum. However, the extraction experiments in Fig. 2 showed a steady decrease in  $D$  value when going from left to right across the lanthanide series. This suggests the same speciation for all investigated lanthanides. When a change in speciation would occur across the lanthanide series, this would be clearly visible as a discontinuity in the separation factors and  $D$  values as a function of the atomic number. On the other hand, the elemental composition of the loaded organic phase with CHN,  $^{31}\text{P}$  NMR and TXRF suggested the presence of 6 nitrate anions per extracted  $\text{La}^{3+}$ . It could be argued that this is caused by the fact that the organic phase was not maximally loaded. Moreover, the bond distances La–O<sub>c</sub> and La–N deduced from the EXAFS experiments corresponded well to the bond distances in the crystal structures of hexanitrate complexes. However, no crystal structure of a lanthanide pentanitrate complex could be found for comparison. Finally, in previous work of our group, it was suggested that the stoichiometry of the  $\text{La}^{3+}$  complex was different from that of  $\text{Sm}^{3+}$  based on a different viscosity of the organic phase upon extraction of both metal ions.<sup>50</sup> In retrospect, this viscosity difference was more likely due to the experimental conditions rather than to a difference in speciation of the metal ions. Although having a significant influence on the viscosity, the water content in the organic phase was not controlled or monitored. In addition, the co-extraction of

other metal ions was not taken into account. Because of the many contradictory arguments, the speciation of  $\text{La}^{3+}$  in the organic phase remains inconclusive, but it is expected to be the same as the other lanthanides.

In the introduction, it was mentioned that the negative sequence of the extraction of the lanthanides with nitrate is assumed to be due to steric effects. Our study did not aim at verifying this hypothesis, but it might provide a first insight into the matter. In Fig. 13, the decrease in the bond distances Ln–O<sub>c</sub> and Ln–N is plotted as a function of the atomic number and compared with the ionic radii of the trivalent lanthanide ions  $\text{Ln}^{3+}$  with coordination number 6.<sup>51</sup> The plot of the bond distances is continually parallel with the plot of the ionic radii. If steric effects would influence the coordination of nitrate ions to the lanthanides, it is expected that the bond distance plot would most likely diverge from the ionic radius plot when steric crowding becomes relevant. Based on these observations, the speciation of the lanthanides in the organic phase is most likely not influenced by steric effects and hence not the cause of the negative sequence. From this, it is clear that this work was only the start and much more research will be required to finally give an answer to the remarkable extraction behaviour of the lanthanides in basic nitrate extractants.

## Conclusions

The structure of the lanthanide complexes extracted by an undiluted quaternary phosphonium nitrate and a quaternary ammonium nitrate extractant was investigated by a combination of complementary speciation techniques including FTIR, photoluminescence spectroscopy, luminescence lifetime measurements, elemental analysis (CHN and TXRF), quantitative  $^{31}\text{P}$  NMR and EXAFS. It was suggested that the speciation is the same for the entire lanthanide series. The results for  $\text{La}^{3+}$  are inconclusive, but the speciation is expected to be identical to that of the other trivalent lanthanides. The lanthanide ions are suggested to form a sole species in the organic phase, namely the pentanitrate complex  $[\text{Q}]_2[\text{Ln}(\text{NO}_3)_5]$ , in which five nitrate ligands are bidentately coordinated to the lanthanide ion. The anionic complex is neutralized by two extractant cations  $\text{Q}^+$ . The lanthanide complexes do not contain any coordinating water ligands. This work provides a first insight into the negative lanthanide extraction sequence.

## Conflicts of interest

There are no conflicts to declare.

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