Opportunities and challenges of applying advanced X-ray spectroscopy to actinide and lanthanide N-donor ligand systems

Tim Pruessmann, a Peter Nagel, b Laura Simonelli, c David Batchelor, d Robert Gordon, e,f Bernd Schimmelpfennig, a Michael Trumma and Tonya Vitovaa*

a Institute for Nuclear Waste Disposal, Karlsruhe Institute of Technology, Hermann-von-Helmholtz-Platz 1, 76344 Eggenstein-Leopoldshafen, Germany, b Institute for Quantum Materials and Technologies, Karlsruhe Institute of Technology, 76021 Karlsruhe, Germany, c ALBA Synchrotron Light Facility, Cerdanyola del Vallès 08290, Spain, d Institute for Photon Science and Synchrotron Radiation, Karlsruhe Institute of Technology, Hermann-von-Helmholtz-Platz 1, 76344 Eggenstein-Leopoldshafen, Germany, e PNCSRF, APS Sector 20, Argonne, IL 60439, USA, and f Moyie Institute, Burnaby, BC, Canada. *Correspondence e-mail: tonya.vitova@kit.edu

N-donor ligands such as $n$-Pr-BTP [2,6-bis(5,6-dipropyl-1,2,4-triazin-3-yl)-pyridine] preferentially bind trivalent actinides (An$^{3+}$) over trivalent lanthanides (Ln$^{3+}$) in liquid–liquid separation. However, the chemical and physical processes responsible for this selectivity are not yet well understood. Here, an explorative comparative X-ray spectroscopy and computational ($L_3$-edge) study for the An/Ln $L_3$-edge and the N $K$-edge of [An/Ln($n$-Pr-BTP)$_3$(NO$_3$)$_3$, [Ln($n$-Pr-BTP)$_3$(CF$_3$SO$_3$)$_3$ and [Ln($n$-Pr-BTP)$_3$(ClO$_4$)$_3$ complexes is presented. High-resolution X-ray absorption near-edge structure (HR-XANES) $L_3$-edge data reveal additional features in the pre- and post-edge range of the spectra that are investigated using the quantum chemical codes FEFF and FDMNES. X-ray Raman spectroscopy studies demonstrate the applicability of this novel technique for investigations of liquid samples of partitioning systems at the N $K$-edge.

1. Introduction

A way to significantly reduce the volume and the heat load of a repository after some decades and the long-term radioactivity of spent nuclear fuel is the so-called Partitioning and Transmutation (P&T) strategy (Magill et al., 2003; OECD, 2011; Salvatores & Palmiotti, 2011; González-Romero, 2011). Here the different components are separated (partitioning) and the transuranium elements are converted to short-lived or stable nuclides (transmutation) using neutron-induced fission or capture reactions. A major step in P&T is the separation of 5f elements, especially the minor actinides (An), from their chemically similar 4f counterparts. This separation is necessary, as the lanthanide (Ln) fission products have large neutron absorption cross sections and thereby compromise transmutation efficiency in the nuclear fission process. Selective liquid–liquid extraction of An$^{3+}$ from Ln$^{3+}$ has been demonstrated using soft nitrogen or sulfur-donor ligands (Madic et al., 2007; Ekberg et al., 2008; Kolarik, 2008; Lewis et al., 2010; Harwood et al., 2011; Hudson et al., 2013; Panak & Geist, 2013), especially heterocyclic N-donor ligands, e.g. bistriazinylpyridines (BTP) (Case, 1971; Kolarik et al., 1999a,b) and bistriazinylbipyrindines (BTBP) (Geist et al., 2006) with separation factors (SFs) for Am$^{3+}$ vs Eu$^{3+}$ greater than 100 ($SF = \frac{D_{Am}}{D_{Eu}}$; $D_M = \frac{[M]_{org}}{[M]_{aq}}$). Additionally, the Ln elements exhibit a trend in selectivity as a function of their ionic radii (Steppert et al., 2009). However,
2.6-bis(5,6-dipropyl-1,2,4-triazin-3-yl)-pyridine (n-Pr-BTP) has slow extraction kinetics (Weigl et al., 2006) and low chemical and radiation stability (Hill et al., 2000; Rat & Hérès, 2000). Modifications like different alkyl groups (Panak & Geist, 2013; Kolarik et al., 1999b; Hudson et al., 2006; Trumm, Geist et al., 2011), substitution in the 4-position of the pyridine ring (Trumm, et al., 2011; Benay et al., 2011; Trumm, Wipff et al., 2011) or annulation [e.g. 6,6'-bis(5,5,8,8-tetra-methyl-5,6,7,8-tetrahydrobenzo-1,2,4-triazin-3-yl)-pyridine (CyMe4-BTP) (Hudson et al., 2003; Benay et al., 2010), bis-2,6-(5,6,7,8-tetra-hydro-5,9,9-tri-methyl-5,8-methano-1,2,4-benzotriazin-3-yl)-pyridine (CA-BTP) (Trumm, Wipff et al., 2011)] lead to improvements only of some properties at the cost of others. Structural changes in the lateral rings (Steppert et al., 2009; Halcrow, 2005; Girt et al., 2010; Bremer, Geist et al., 2012; Bremer, Ruff et al., 2012; Beele et al., 2013; Bremer et al., 2013) can lead to drastic changes in extraction properties, e.g. the separation factors (SF) for Am$^{3+}$ over Eu$^{3+}$ change from SF$_{Am/Eu}$ = 130 for n-Pr-BTP (Kolarik et al., 1999a) to SF$_{Am/Eu}$ = 5 for 6-bis(4-ethyl-pyridazin-1-yl)-pyridine (Et-BDP) (Beele et al., 2012). The physical and chemical properties leading to strong differences in the selectivity, kinetics and stability of these related organic molecules are not yet well understood. There are numerous experimental and theoretical structural studies in the literature looking for small electronic and geometric structural variations, which might be associated with differences in bonding affinities for An compared with Ln.

This work is focused on the n-Pr-BTP molecule and the respective An$^{3+}$ and Ln$^{3+}$ complexes with the n-Pr-BTP ligand. We applied XANES and X-ray Raman spectroscopy experimental techniques and computations with the FEFF (Rehr et al., 2009, 2010) and FDMNES (Bunäu & Joly, 2009) quantum chemical codes, which are element- and bulk-sensitive probes and can be used for all Ln and An elements and N in materials in solid and liquid phase, which is not always possible with other methods. Electronic and geometric structural investigations from the ‘point of view’ of the metal and the ligand can facilitate understanding the fundamental principles of the N-donor ligands’ selectivity. We detail experimental and computational approaches pointing out their advantages and limitations. Our aim is to guide related future studies.

2. Methods

2.1. Materials

The ligand investigated in this work is 2,6-bis(5,6-dipropyl-1,2,4-triazin-3-yl)-pyridine (n-Pr-BTP) (Case, 1971; Kolarik et al., 1999a,b). For quantum chemical calculations 2,6-bis-(1,2,4-triazin-3-yl)-pyridine (H-BTP) (Case, 1971; Boucher et al., 2002) has been used as an approximation for n-Pr-BTP. NO$_3^-$ is the preferred counter ion as the extraction process of An and the separation from Ln takes place in nitric acid solution. However, the NO$_3^-$ N K-edge spectrum has a small contribution in the n-Pr-BTP N K-edge spectrum (see Fig. 18). In ClO$_4^-$ containing solution, precipitation occurs after adding n-Pr-BTP; therefore it is discarded as an alternative to NO$_3^-$.

Complexes with CF$_3$SO$_4^-$ (OTf) as counter ion exhibit good solubility and OTf has a lower bonding affinity to the metal than NO$_3^-$ thereby enhancing the formation of 1:3 complexes. From time-resolved laser fluorescence spectroscopy (TRLFS) experiments (Panak & Geist, 2013; Trumm et al., 2010) on Cm and Eu it can be seen that already above ~0.6 mmol L$^{-1}$ n-Pr-BTP concentration 1:3 complexes are formed exclusively. As an alternative, the [An/Ln(n-Pr-BTP)$_3$](OTf)$_3$ complexes were first crystallized before solving and drying on substrates. Solutions of [An/Ln(n-Pr-BTP)$_3$](OTf/ClO$_4$/NO$_3$)$_3$ complexes have also been dried on substrates in some of the experiments. The Ln salts that have been used in the preparation of the complexes and as reference materials for the L$_Y$-edge measurement are hydrophilic and can contain up to nine H$_2$O in the first coordination shell. Especially for experiments in solution, a high H$_2$O coordination has to be assumed but could not be quantified during the measurements. Calculations have been performed with [Ln(H$_2$O)$_n$(OTf)$_3$]. For the sake of simplicity, coordinating H$_2$O molecules are not explicitly named throughout this work. Pu$^{4+}$ has been shown to be stable in [Pu(n-Pr-BTP)$_3$](NO$_3$)$_3$ complexes (Banik et al., 2010).

2.2. Experiments

High-resolution X-ray absorption near-edge structure (HR-XANES) spectra and core-to-core resonant inelastic X-ray scattering (CC-RIXS) maps at Ln L$_Y$-edges were collected at the ID26 beamline at the European Synchrotron Radiation Facility (ESRF), Grenoble, France. The X-rays emitted from the sample were energy analyzed by a Johann spectrometer in scanning geometry (Glatzel et al., 2009) and detected by an avalanche photodiode. The L$_{a_1}$ or L$_{b_2}$ emission lines of different Ln were diffracted and focused by spherically bent crystals with a 1 m radius of curvature at Bragg angle [Eu L$_{a_1}$: Ge(333)/76.75°; Gd L$_{a_1}$: Si(333)/78.28°; Gd L$_{b_2}$: Ge(620)/77.39°].

Pu and Am L$_Y$-edge HR-XANES spectra have been collected at the INE beamline (Rothe et al., 2012; Walsh et al., 2014; Zimina et al., 2016, 2017) at the Karlsruhe Institute of Technology (KIT) light source (KARA storage ring, KIT Campus North). The X-rays emitted from the sample were energy analyzed by a Johann spectrometer and detected by a Vortex silicon drift detector (Iwanczyk et al., 2003). The L$_{a_1}$ emission lines of Pu (14 282 eV) and Am (14 620 eV) were diffracted and focused by five spherically bent Si(777) crystals with a 1 m radius of curvature at Bragg angle 75.7° and 71.2°, respectively. The samples were measured in solutions of 2 mM and 8 mM Am concentration, respectively.

N K-edge XANES spectra were measured at the UE52-PGM beamline at the BESSY II synchrotron source of the Helmholtz-Zentrum Berlin using partial electron yield detection and a resolving power ($E$/Δ$E$) of ~10000. A plane-grating monochromator (PGM) with a 1200 lines mm$^{-1}$ grating was used to tune the incident X-rays from 380 to 420 eV. The energy of the incident X-ray beam was scanned with 0.05 eV step size. n-Pr-BTP, [Ho(n-Pr-BTP)$_3$](ClO$_4$)$_3$, and [Ho(n-Pr-BTP)$_3$](NO$_3$)$_3$ were solved in isopropanol.
(10 mmol L⁻¹) and ~2 μL were dried on aluminium sample holders.

Additional N K-edge XANES experiments were performed in fluorescence mode at the WERA beamline at the KIT Light Source (KARA storage ring, KIT Campus North). The incident beam was monochromatized by an SGM. The energy of the incident beam was scanned with 0.05 eV step size across the pre-edge and edge region and up to 0.5 eV in the post-edge region. The scanned energy range is from 370 to 485 eV. The spectra have been calibrated using the Ni L₂,₃-edge of a NiO reference. The n-Pr-BTP samples were prepared by solving the ligand in ethanol and drying ~2 μL of the solutions on Al foil.

X-ray Raman spectroscopy experiments were performed at the 20-ID beamline at the Advanced Photon Source (APS) using the Lower Energy Resolution Inelastic X-ray scattering (LERIX) spectrometer (Fister et al., 2006). O K-edge X-ray Raman spectra of water and isopropanol and N K-edge X-ray Raman spectra of crystalline n-Pr-BTP pressed into a pellet were measured with the analyzer energy set at 9890 eV [Si(555), θ = 88.27°]. The incident photon flux was 2 × 10¹² photons s⁻¹ using a double-crystal monochromator with Si(111) crystals. The overall experimental energy resolution was 1.3 eV estimated by measuring the FWHM of the elastically scattered radiation. The liquid samples were measured using a flow-through cell developed in cooperation with Diamond Materials GmbH (Freiburg, Germany). The cell has a diamond window with thickness of 50 μm and was set at 45° or 30° with respect to the incident beam inside a He-filled chamber. A peristaltic pump set to a flow rate of 5 mL min⁻¹ was used to pump the liquids. For the crystalline sample, 18 detectors were employed and the sample was placed on a spinner to reduce radiation damage. Due to the cell blocking part of the scattered energy, only 15 detectors (cell at 30°) or 12 detectors (cell at 45°) out of 18 detectors could be used for liquid measurements.

X-ray Raman spectroscopy experiments were performed at the ID20 beamline at the ESRF using a spectrometer with 72 analyzer crystals. N K-edge X-ray Raman spectra of crystalline n-Pr-BTP and n-Pr-BTP solved in isopropanol (30 mmol L⁻¹) were measured with the analyzer energy set at 9690 eV [Si(660), θ = 88.5°]. The incident photon flux was ~10¹² photons s⁻¹. The cell and pump setup was the same as used at the APS with the cell set at 45° to the incident beam. Due to the cell blocking part of the scattered X-rays only 24 detectors could be used for liquid measurements. The crystalline n-Pr-BTP powder was pressed into a depression in an Al plate set at 30° to the incident beam. This allowed the use of 48 crystals.

2.3. Computations

2.3.1. Structure optimization. Ln(H-BTP)₃ and An(H-BTP)₃ structures were optimized imposing the D₃ point group at the density functional theory (DFT) level with the BP86 functional (Becke, 1988; Perdew, 1986) employing the resolution-of-the-identity (RI) routines as implemented in TURBOMOLE (TURBOMOLE, 2012). Def2-TZVP basis sets (Weigend et al., 1998) were used for the light elements whereas the def-TZVP basis sets (Eichkorn et al., 1997) and small-core Stuttgart-PP were taken for the f-elements. Structures optimized in this way are denoted structure I throughout this work. H-BTP is used as an approximation for n-Pr-BTP as it has been proved that the substituents on the triazine rings hardly affect the complex structure (Berthet et al., 2002; Denecke et al., 2005; Petit et al., 2006).

Additionally Gd(H-BTP)₃ structures in gas and aqueous phase (called structure 2 and structure 3, respectively) have been calculated. The structures were optimized on the DFT level in C₃ symmetry using a SVP basis set and the BH-LYP functional (Becke, 1993). For the aqueous phase the structure has been solvated with COSMO-RS (Klamt, 1995).

2.3.2. Calculation of Ln/An L₃-edge spectra. To compare different codes and methods, the spectrum of [Eu(n-Pr-BTP)₃][(OTf)₃] was calculated with FEFF9.6 (Rehr et al., 2009, 2010) and FDMNES (Bunû & Joly, 2009). The structural data of [Eu(n-Pr-BTP)₃][(OTf)₃] were taken from unpublished X-ray diffraction data. With FEFF, the self-consistent field (SCF) was calculated for a cluster of 28 atoms and the full multiple scattering (FMS) for a cluster of 46 atoms. A Hedin–Lundqvist-type exchange correlation potential was used. FDMNES FMS and full potential (FP) calculations were performed on a cluster of 28 atoms. In all calculations, quadrupole transitions were taken into account.

To compare different structure optimizations the experimental spectrum of [Gd(n-Pr-BTP)₃][(OTf)₃] was compared with FDMNES FMS spectra calculated from Gd(H-BTP)₃ optimized in the different ways described above, i.e. optimization with D₃ symmetry, with C₁ symmetry and with C₁ point group and solvation shell.

The [Gd(n-Pr-BTP)₃][(OTf)₃] spectrum and the f and d angular momentum projected density of states (f-, d-DOS) were calculated using the ab initio multiple-scattering theory based FEFF9.5 code (Rehr et al., 2009, 2010). The SCF and FMS calculations were performed on a cluster of 152 atoms corresponding to one [Gd(n-Pr-BTP)₃][(OTf)₃] molecule. A Hedin–Lundqvist-type exchange correlation potential was used. The f-DOS was calculated self-consistently by using the UNFREEZE/ENZEF card. The Fermi level was set to 3 eV below the calculated value of ~9.6 eV in order to reproduce the pre-edge structure in the spectrum. To reach convergence, the core-hole type was set to random phase approximation (RPA card) and the core-hole potential was calculated for a cluster of 47 atoms (SCREEN card).

3. Results and discussion

3.1. Ln/An L₃-edge

The electronic and geometric structures of [An/Ln(n-Pr-BTP)₃][(NO₃)₃], [Ln(n-Pr-BTP)₃][(CF₃SO₃)₃] and [Ln(n-Pr-BTP)₃][(ClO₄)₃] complexes are probed from the metal ‘point-of-view’ using An/Ln L₃-edge XRANES and CC-RIXS as well as HR-XANES ab initio quantum chemical calculations.
with the FEFF and FDMNES codes. Benchmark calculations using different codes and structural models are carried out.

3.1.1. Ln/An $L_3$-edge HR-XANES and CC-RIXS spectra. Fig. 1 shows a simplified molecular orbital (MO) scheme of An/Ln bound to N and the excitations relevant for X-ray absorption spectroscopy (XAS) at the An/Ln $L_3$-edge. At the white line (WL) electrons are excited from An/Ln 2$p_{3/2}$ states to unoccupied molecular orbitals mixtures of An/Ln 5$d$/5$6d$ and N 2$p$ atomic orbitals.

The WL is mainly used to observe changes in the oxidation states (Centeno et al., 2000), i.e. the relative energy shift of the absorption edge due to the change in valence orbital occupation and the resulting change in electron density and screening of the core-hole. The WL is also sensitive to the local structure, and the density of occupied and unoccupied states, that depend also on other factors than the oxidation state, e.g. the chemical environment, i.e. type of bonding partner (Asakura et al., 2014), coordination number, local symmetry (Asakura et al., 2015), changes due to induced pressure (Rueff, 2009; Heathman et al., 2010), etc. This can lead to changes in energy position, width, shape and intensity of the WL. From a quantum chemical point of view $L_3$-edge spectra describe the angular momentum projected density of states (DOS) of unoccupied $d$ and $s$ (less important) like states (Mott, 1949), $5d$/5$6d$ states in the case of Ln/An $L_3$-edge spectra following the dipole selection rules $\Delta l = \pm 1$ ($\Delta J = 0, \pm 1$).

After the first experimental observation of a weak pre-edge structure in Ce XANES (Bianconi et al., 1987), band structure calculations showed a correspondence to quadrupole transitions (Finkelstein et al., 1992). Even though quadrupole transitions (2$p_{3/2} \rightarrow 4f$) are considerably weaker than dipole transitions (2$p_{3/2} \rightarrow 5d$/5$6d$), they have been observed for a variety of Ln (2$p_{3/2} \rightarrow 4f$ transitions) and An (2$p_{3/2} \rightarrow 5f$ transitions) materials using X-ray emission spectrometers with an instrumental energy bandwidth similar to the core-hole lifetime broadening (Vitova et al., 2010, 2013, 2015, 2018; Krisch et al., 1995; Kvashnina et al., 2011; Hämäläinen et al., 1991; Carra & Altarelli, 1990; Tanaka et al., 1994; Carra et al., 1995; Bartolomé et al., 1997, 1999; Gallet et al., 1999; Dallera et al., 2000, 2003, 2004, 2006; Journel et al., 2002; Nakazawa et al., 2002; Rueff et al., 2004, 2006; Glatzel et al., 2005; Sham et al., 2005; Yamaoka et al., 2006; Brouder et al., 2008; Kotani, 2008). Atomic multiplet calculations of the pre-edge spectral features of Ln $L_3$ HR-XANES demonstrated that they are shaped by electron–electron interactions and their shape, intensity and energy position on the excitation and emission energy depend on the number of available $f$ electrons in the systems (Kvashnina et al., 2011; Severing et al., 1989; Hansmann et al., 2008). The intensity of the pre-edge can also increase in the case of hybridization of Ln $f$ and $d$ states.

In a standard measurement, the core-hole lifetime broadening affects the XANES spectrum leading to a significant spectral broadening compared with HR-XANES (see Fig. 2). Most notably, in HR-XANES, a pre-edge (feature A) can be resolved, the WL is sharper and has higher intensity and post-edge features (B) are well separated from the WL. The HR-XANES pre-edge and WL provide details on the electronic structure not available with conventional methods. Post-edge features close to the WL are sensitive to multiple scattering of the photoelectron with the surrounding shells and therefore the local atomic geometry around the absorbing atom, e.g. bonding angles, become accessible. It can complement the information relative to interatomic distances, local structural disorder, and number and kind of atoms surrounding the absorber, which can be quantitatively obtained by analyzing the extended X-ray absorption fine-structure (EXAFS) region.

By recording HR-XANES selecting different emission lines with similar energy resolution, it is possible to obtain HR spectra with different sensitivity, due to the different intrinsic final state core-hole lifetime broadening [e.g. U $L_3$ HR-XANES collected at the $L_{\beta5}$ emission line (Kvashnina et al., 2014)]. It can also be possible to detect additional spectral features due to different screening of core-holes at different energy levels. In the case of Ln $L_3$-edge spectra the $L_{\beta2}$ emission line can be used instead of the conventional high-intensity $L_{\alpha1}$ emission line. However, the $4d$ core-hole...
resulting from the $L_{g2}$ emission has a higher core-hole lifetime broadening (2 eV) (McGuire, 1972) than the $3d$ core-hole resulting from the $L_{a1}$ emission (1.18 eV) (McGuire, 1974) hence no increase in resolution can be achieved. In Fig. 3 the spectra of [Gd(n-Pr-BTP)$_3$](OTf)$_3$, measured recording the emitted fluorescence by fixing the outcome energy at the maximum of the $L_{a1}$ or $L_{g2}$ emission lines, are compared.

The WL of the spectrum measured at the $L_{g2}$ emission line is slightly narrower and the pre-edge has only one instead of two peaks. To further investigate the differences in the pre-edge region, CC-RIXS maps were recorded. Fig. 4 shows 2p$3d$ ($L_{a1}$) and 2p$4d$ ($L_{g2}$) RIXS maps of the pre-edge of [Gd(n-Pr-BTP)$_3$](OTf)$_3$ with a red line marking the emission energy at which the HR-XANES spectra shown in Fig. 3 were measured.

The 2p$3d$ RIXS map has two features at the same excitation energy at $\sim$1180.5 eV and $\sim$1182.5 eV energy transfer. The HR-XANES spectrum cuts through both features leading to the observed double structure in the pre-edge. In contrast, the 2p$4d$ RIXS map has one feature at $\sim$137.5 eV energy transfer and additional intensity at $\sim$140 eV that could be the overlap of pre-edge and WL tails or a second feature comparable with 2p$3d$ RIXS, but with lower intensity. The HR-XANES spectrum only intersects the maximum of the first feature resulting in a single pre-edge peak. Because the differences in the CC-RIXS maps appear along the energy transfer scale, they are related to final state effects. In this case these are different splitting of the 2p$3d^5$5d$^1$ and 2p$4d^5$5d$^1$ final states, respectively; changes in the screening of the core-hole cause an energy shift of the pre-edge resonance relative to the normal emission, i.e. the maximum of the emission line in the post-edge range. Since the 4$f$–4$f$ electron–electron interactions have the same influence on both spectra, the appearance of two resonances along the energy transfer scale is likely related to differences in the 3$d$–4$f$ and 4$d$–4$f$ electron–electron interactions. It was shown that they have the highest influence on the spectrum after the 4$f$–4$f$ electron–electron interactions (Kvashnina et al., 2011).

3.1.2. Calculations of Ln $L_{y}$-edge HR-XANES spectra of [Ln(n-Pr-BTP)$_3$](OTf)$_3$. Ln $L_{y}$-edge HR-XANES spectra of Ln(H-BTP)$_3$ and [Ln(n-Pr-BTP)$_3$](OTf)$_3$ were calculated with different codes and various structural models to find the best agreement between experimental and theoretical spectra. In Fig. 5 the effects of using different codes (FEFF and FDMNES) and methods (FMS and FP) to calculate an Eu $L_{y}$-edge spectrum of [Eu(n-Pr-BTP)$_3$](OTf)$_3$ are shown and compared with experimental data. The calculated spectra are less broadened than the experimental spectrum to be able to see all features clearly. Therefore, the pre-edge (A) and the first post-edge feature (B) of the calculated spectra show significantly larger intensity compared with the experimental spectrum. The FEFF and FDMNES calculations reproduce well the A, D (FEFF) and B (FDMNES) features, respectively. Between the FDMNES FMS and FP calculations there are intensity differences for the A and B features and a small energy shift for the D peak. Since the differences are small and an FP calculation takes significantly longer than an FMS calculation (FMS: 1 h; FP: 1 week), FP calculations were not further applied.
Fig. 6 depicts the differences in the calculated spectra resulting from different structures used in the calculation. Both \textit{FEFF} and \textit{FDMNES} calculations were performed for a DFT optimized Eu(H-BTP)$_3$ structure. They are compared with the \textit{FEFF} and \textit{FDMNES} FMS spectra shown also in Fig. 5 calculated with the experimental crystalline [Eu(n-Pr-BTP)$_3$](OTf)$_3$ structure and the experimental spectrum.

The B and D absorption resonances are shifted to lower energies for the spectrum calculated with \textit{FEFF} using the DFT optimized structure compared with the spectrum using the experimental structure; the former spectrum has less agreement with the experimental spectrum. This effect is probably due to the 5 pm larger Eu—N bond length in the DFT optimized structure. There is no effect on the pre-edge. For the \textit{FDMNES} calculations on the other hand, the pre-edge of the spectrum using the experimental structure has larger intensity compared with the spectrum using the optimized structure, perhaps due to changes in the electronic density associated with the shorter bond length. B and C exhibit only small differences, suggesting that \textit{FDMNES} treats multiple-scattering effects less accurately than \textit{FEFF}.

Fig. 7 shows the spectra of differently optimized Gd(H-BTP)$_3$ structures calculated with \textit{FEFF} and \textit{FDMNES}. Structures were optimized using DFT with symmetry restrictions (structure 1), without symmetry restrictions (structure 2) and without symmetry restrictions and a solvation sphere (structure 3). Geometric differences between the structures are summarized in Table 1, Fig. 8.
It can be seen that the average Gd—N bond length changes as well as the orientation of the triazine rings with respect to the pyridine ring. In the spectra calculated with FEFF a shift to higher energies of B and D can be observed from structure 1 to structure 3 showing that structure 3 is closest to the real structure. The same effect is visible in the spectra for structures 2 and 3 calculated with FDMNES. In the spectrum of structure 1, however, B and D are significantly shifted to lower energies and the pre-edge shows a double structure that is not visible in the other spectra. For the spectra compared here, the spectrum of structure 3 has the best agreement with the experimental spectrum indicating that structure 3 is closer to the real structure than the other two. Further improvements can be expected using higher-level quantum chemical methods, which are not always applicable for large molecules.

Fig. 9(a) shows FEFF shell-by-shell calculations of Gd(H-BTP)3 in a simplified approach to correlate the spectral features with specific groups of atoms surrounding the absorbing atom. The spectra of Gd(H-BTP)3 with structure 1 were calculated with SCF and FMS radius increasing from 3 Å to 6 Å in 1 Å steps. A scheme of the used shells is shown in Fig. 9(b). The spectrum only including the contributions from the first shell, i.e. the bonding N atoms, is nearly featureless because only a few multiple-scattering paths are available. Adding the non-bonding neighbors of the bonding N atoms to the calculation as a second shell leads to clearly resolved features B and D. When adding additional atoms to the calculation, B is shifted to lower energies, farther away from the experimental energy position, and D is shifted to higher energies, closer to the experimental energy position. These shifts are due to the interference of additional scattering signals and indicate that B is mainly influenced by first- and second-shell atoms while D is related to third- and fourth-shell atoms. The pre-edge A also exhibits small changes probably due to changes in the scattering potentials of the atoms surrounding the Gd.

3.1.3. Comparison between Gd L3-edge HR-XANES of [Gd(n-Pr-BTP)]3[NO3]3 and [Gd(NO3)3](OTf)3. Gd L3-edge HR-XANES spectra of [Gd(n-Pr-BTP)]3[NO3]3 and [Gd(NO3)3]3 are shown in Fig. 10(a). The [Gd(n-Pr-BTP)]3[NO3]3 spectrum is shifted 0.4 eV to lower energies compared with the [Gd(NO3)3]3 spectrum indicating higher electronic density on the cation in the [Gd(n-Pr-BTP)]3[NO3]3 complex. This trend is preserved if NO3− is exchanged with an OTf− anion. The [Gd(n-Pr-BTP)]3[OTf]3 and [Gd(OTf)3]3 spectra are plotted in Fig. 10(b) showing a 0.5 eV energy shift of [Gd(n-Pr-BTP)]3[OTf]3 to lower energies. The energy shifts were validated by ensuring that the measured emission lines coincide in energy position. The additional pre-edge intensity at about 7241 eV (marked with A in Fig. 10), more noticeable for [Gd(n-Pr-BTP)]3[NO3]3 compared with [Gd(NO3)3], can be assigned to 2p5/2 photoelectron transitions to 4f and/or 5d final states.

The origin of the pre-edge feature is revealed by calculations with the FEFF9.5 and the FDMNES codes. The calculated Gd(H-BTP)3 spectrum and the f- and d-DOS are plotted in Fig. 11. The f-DOS has high intensity at the energy position of the pre-edge feature, whereas the 5d states have minor contributions. This result suggests that this feature arises from electronic transitions to orbitals with major 4f and minimal 5d Ln participations. Even though the direct bonding partners of Gd change from N in [Gd(n-Pr-BTP)]3[OTf]3 to O in [Gd(OTf)3], hardly any difference between the areas of the pre-edges...
and Wls is detectable, indicating that the relative electronic populations of the Gd 4f and 5d states are not significantly influenced by bonding with the n-Pr-BTP molecule. Nevertheless, the −0.4 eV relative energy shift of the WL for [Gd(n-Pr-BTP)3](OTf)3 over Gd(OTf)3 is a clear indication of better screening of the 2p core-hole due to higher electron density on Gd in [Gd(n-Pr-BTP)3](OTf)3 than in Gd(OTf)3. The pre-edge feature B is at lower energy in [Gd(n-Pr-BTP)3](OTf)3 than the corresponding feature C in Gd(OTf)3. In Fig. 10(a) the spectra of [Gd(n-Pr-BTP)3](NO3)3 and Gd(NO3)3 are compared. The same features and energy shifts are noticeable; they are less pronounced due to the different experimental setup resulting in lower energy resolution for [Gd(n-Pr-BTP)3](NO3)3 and Gd(NO3)3. Due to the specific energy resolution of different beamlines/experimental setups and the varying intrinsic broadening of different elements it is not possible to quantitatively compare the areas of the peaks for the different compounds.

3.1.4. Comparison between An L3-edge HR-XANES of [Am(n-Pr-BTP)3](NO3)3 and Am(NO3)3.

The direct comparison of experimental [Pu(n-Pr-BTP)3](NO3)3 and [Am(n-Pr-BTP)3](NO3)3 spectra [Fig. 14(a)] shows the slightly broader WL of [Am(n-Pr-BTP)3](NO3)3. Feature D has different intensity in both spectra due to problematic normalization. In addition, the low signal-to-noise ratio in this range prevents the detection of the expected pre-edge feature A arising from excitations to a mixture of d and f states.

In Fig. 13, features A and B are not resolved in the Am L3-edge HR-XANES Am(NO3)3 and [Am(n-Pr-BTP)3](NO3)3 spectra due to the higher core-hole lifetime (Am: 3.87 eV; Pu: 3.74 eV) and experimental broadening (Am: 3.88 eV; Pu: 4.14 eV) for Am compared with Pu. The WL is broader in the [Am(n-Pr-BTP)3](NO3)3 spectrum compared with Am(NO3)3. For both Pu and Am the spectra have similar feature D 35 eV above the WL and the spectra of the complexes are shifted to lower energies (<0.5 eV). This indicates a better screening of the 2p core-hole due to higher charge density on the metal in [Pu/Am(n-Pr-BTP)3](NO3)3 than in Pu/Am(NO3)3 similar to the results for the Ln compounds.

The direct comparison of experimental [Pu(n-Pr-BTP)3](NO3)3 and [Am(n-Pr-BTP)3](NO3)3 spectra [Fig. 14(a)] shows the slightly broader WL of [Am(n-Pr-BTP)3](NO3)3. Feature D has different intensity in both spectra due to problematic normalization. In addition, the low signal-to-noise ratio in this range prevents the detection of the expected pre-edge feature A arising from excitations to a mixture of d and f states.

In Fig. 13, features A and B are not resolved in the Am L3-edge HR-XANES Am(NO3)3 and [Am(n-Pr-BTP)3](NO3)3 spectra due to the higher core-hole lifetime (Am: 3.87 eV; Pu: 3.74 eV) and experimental broadening (Am: 3.88 eV; Pu: 4.14 eV) for Am compared with Pu. The WL is broader in the [Am(n-Pr-BTP)3](NO3)3 spectrum compared with Am(NO3)3. For both Pu and Am the spectra have similar feature D 35 eV above the WL and the spectra of the complexes are shifted to lower energies (<0.5 eV). This indicates a better screening of the 2p core-hole due to higher charge density on the metal in [Pu/Am(n-Pr-BTP)3](NO3)3 than in Pu/Am(NO3)3 similar to the results for the Ln compounds.

The direct comparison of experimental [Pu(n-Pr-BTP)3](NO3)3 and [Am(n-Pr-BTP)3](NO3)3 spectra [Fig. 14(a)] shows the slightly broader WL of [Am(n-Pr-BTP)3](NO3)3. Feature D has different intensity in both spectra due to problematic normalization. In addition, the low signal-to-noise ratio in this range prevents the detection of the expected
small variations. The spectra of Pu(H-BTP)₃ and Am(H-BTP)₃ calculated with FDMNES [Fig. 14(b)] show differences in the pre-edge (A) due to the changing occupation of the 5f orbitals (5f⁵ for Pu and 5f⁴ for Am). Feature D in Am(H-BTP)₃ is shifted to higher energies due to the shorter An–N bond length in Am(H-BTP)₃ compared with Pu(H-BTP)₃. Both these effects are not resolved experimentally.

3.2. N K-edge XANES experiments – feasibility studies

Fig. 15 shows N K-edge spectra of [Ho(n-Pr-BTP)₃](NO₃)₃ measured at the WERA beamline in partial electron yield (PEY), total electron yield (TEY) and fluorescence mode. PEY detection measures electrons emitted from the sample and discriminates between photo- and Auger electrons. This method is surface sensitive up to a depth of 1–2 nm. TEY detection measures all emitted electrons by counting the current necessary to neutralize the sample. Both electron detection methods are susceptible to charge build-up in non-conducting samples like n-Pr-BTP and its complexes. They are also not element selective, which is especially problematic for N K-edge spectra in the presence of C due to a large non-linear background from the C K-edge. Thus, fluorescence detection is generally preferred for materials for N K-edge investigations in the presence of C, even though it has worse signal-to-noise ratio and possible self-absorption artifacts compared with the electron detection methods. At the UE52-PGM beamline, however, the fluorescence detector was unavailable for the beamtime, so PEY has been used.

Fig. 16 shows N K-edge spectra of n-Pr-BTP measured in PEY at the UE52-PGM beamline at BESSY and at the WERA beamline at KARA. Both spectra have a similar signal-to-noise ratio. The spectrum collected at the UE52-PGM beamline is shifted to higher energies compared with the spectrum collected at the WERA beamline, because at the UE52-PGM beamline no reference spectra could be collected to calibrate the energy scale. In both spectra the FWHM is \( \sim 1 \) eV and spectral features are equally visible.

During the measurements at BESSY II, changes of pre-edge features are observed after irradiation of the samples (Fig. 17). Feature A loses intensity while the intensity of feature B is increased. These effects are attributed to radiation damage, i.e. ionization and breaking of bonds by the incident X-ray beam, and were not observed at the WERA beamline (KARA) due to the lower photon flux density impinging on the sample.

Fig. 18(a) shows an N K-edge spectrum of Ho(NO₃)₃ compared with [Ho(n-Pr-BTP)₃](NO₃)₃. The main intensity of the Ho(NO₃)₃ spectrum lies at higher energies than the [Ho(n-Pr-BTP)₃](NO₃)₃ pre-edge, which is barely influenced. This is confirmed by direct comparison between NO₃⁻-containing and NO₃⁻-free ([Ho(n-Pr-BTP)₃](NO₃)₃ and [Ho(n-Pr-BTP)₃](ClO₄)₃ complexes, see Fig. 18(b).
3.3. X-ray Raman spectroscopy – feasibility studies

The process of separation of An from Ln takes place in a liquid phase. However, the N K-edge low photon energy (~400 eV) requires ultra-high-vacuum investigations challenging for liquid samples. First XAS tests with a liquid, constant flow, pump-through cell equipped with 150 nm SiC window (Blum et al., 2009) at beamline 8.0.1 at the Advanced Light Source (ALS) resulted in fast formation of X-ray induced radiolysis, i.e. radiation damage, coloring the n-Pr-BTP liquid sample from light orange to dark brown within 2 to 3 minutes. An alternative technique for K-edge XANES investigations of low-Z elements is X-ray Raman spectroscopy. An incident beam with energies above 10 keV facilitates use of liquid sample cells and double containments necessary for investigations of radioactive materials. The high energy of the X-rays reduces the radiation damage and allows penetration through windows in sample cells made from materials such as Kapton, which has high chemical and X-ray stability and therefore is often used as window material. Disadvantages of the technique are: (1) the low cross section of the process [0.13 cm² g⁻¹ inelastic scattering cross section compared with 3 × 10²⁰ cm² g⁻¹ photo-absorption cross section (Henke et al., 1993)] requiring at least 3 at% concentration of N atoms in the sample for 2 × 10¹² photons s⁻¹ incident beam, 18 analyzer crystals, (2) the reduced experimental energy resolution when favoring the flux compared with a dedicated soft X-ray beamline leading to broadening of the spectral features. Only a few X-ray Raman spectroscopy measurements of highly concentrated liquid samples are reported in the literature (Bowron et al., 2000; Bergmann et al., 2002, 2007; Näslund et al., 2005; Juurinen et al., 2013, 2014; Wernet et al., 2004; Pylkkänen et al., 2011; Sahle et al., 2013, 2016; Niskanen et al., 2015).

At the 20-ID beamline at the APS the feasibility studies carried out in November 2012 presented here were the first measurements of liquids performed with the LERIX spectrometer (Fister et al., 2006); therefore they contributed to the development program of the beamline to extend the range of possible samples. Fig. 19(a) shows the O K-edge isopropanol spectrum after averaging two scans measured 45 minutes per scan. We observed formation of gas bubbles during the measurements probably formed by radiolysis or heating of the isopropanol. The angle of the cell with respect to the incoming X-ray beam was changed from 45° to 30° to allow the bubbles to escape from the cell without disturbing the measurement. This cell arrangement reduced the number of usable analyzers from 15 to 12. In the O K-edge water spectrum [Fig. 19(b)], details for the edge region, similar to those reported for water ice in the literature (Fister et al., 2009; Zubavichus et al., 2006), are well distinguishable. The N K-edge spectrum of crystalline n-Pr-BTP measured for 30 minutes by averaging over 18 detectors is shown in Fig. 19(c). The spectrum has a lower signal-to-noise ratio than the standard XANES spectrum (Fig. 18). After one scan (30 min) the changed surface color of the sample indicated damage by the beam. A second scan was consistently showing a lower signal. The n-Pr-BTP solution revealed no visible color changes after several hours exposure to the beam. However, due to the low concentration no discernible N signal was measured.

At the ID20 beamline at the ESRF (Huotari et al., 2017) it was possible to measure n-Pr-BTP solutions due the higher number of available analyzer crystals and the higher incident photon flux (ID20: 10¹⁴ photons s⁻¹; 20-ID: 10¹² photons s⁻¹). The experiments were performed in November 2014. Fig. 20 shows the N K-edge spectra of n-Pr-BTP crystalline and in solution. The crystalline sample has been measured with 48 crystals for 3 h in an evacuated chamber, while the solution sample was measured with 24 crystals for 10 h. The crystalline sample showed similar discoloration as the sample investigated at the 20-ID beamline. A part of the isopropanol in the solution sample evaporated during the measurements. The
pre-edge and WL of the spectra have similar energy positions, but different intensities due to difficulties with the normalization. Even though the signal-to-noise ratio is much worse for the sample in solution, the general structure of the spectrum can be easily recognized and it can be assumed that the structure in solved and crystalline state does not differ significantly. The signal-to-noise ratio of the spectrum of the liquid sample can be improved by utilizing more analyzer crystals and an improved cell design, e.g. using a capillary. In addition, the solution should be temperature controlled to reduce evaporation and avoid precipitation of the complex due to temperature difference between preparation lab and beamline.

4. Conclusions

The Ln/An L\textsubscript{3}-edge HR-XANES technique reveals higher charge density on the metal Ln/An atoms for the [Ln(\textit{n}-Pr-BTP)\textsubscript{3}](OTf/NO\textsubscript{3})\textsubscript{3} compared with the Ln/An(OTf/NO\textsubscript{3})\textsubscript{3} complexes. The high energy resolution allows resolving a pre-edge feature for the [Ln(\textit{n}-Pr-BTP)\textsubscript{3}](OTf)\textsubscript{3} complexes and it is shown that, as indicated in previous studies (Vitova et al., 2010, 2013, 2015, 2018; Finkelstein et al., 1992; Krisch et al., 1995; Kvashnina et al., 2011), it originates from electronic transitions to orbitals with predominant 4\textit{f} character. The shape and energy positions of these pre-edges do not change noticeably for the [Ln(\textit{n}-Pr-BTP)\textsubscript{3}](OTf)\textsubscript{3} and Ln(OTf)\textsubscript{3} complexes. This strongly suggests that the 4\textit{f} states are localized on the metal atom and do not participate in bonding. The shapes of the pre-edges vary and depend on the number of 4\textit{f} electrons. The 4\textit{f} electrons induce electron–electron interactions leading to a complex structure of the 4\textit{f} states, i.e. multiplets. Due to the large core-hole lifetime broadening effects pre-edges are not resolved for the [An(\textit{n}-Pr-BTP)\textsubscript{3}](NO\textsubscript{3})\textsubscript{3} complexes. Use of the L\textsubscript{\textit{ps}} emission lines for measurements of An L\textsubscript{3}-edge HR-XANES spectra would lead to reduced broadening effects. The 5\textit{f} states can be also directly probed by An M\textsubscript{\textit{4,5}}-edge HR-XANES and their level of participation in the chemical bond elucidated.

It is demonstrated that the HR-XANES technique allows resolving post-edge features not visible in the conventional spectra. The correlation of their energy positions to specific structural changes, i.e. interatomic distances and bonding angles, are revealed with the help of FEFF and FDMNES XANES quantum chemical calculations and simulations. Both codes are used for detailed analyses of the spectra. Optimized structures with additions of ‘no symmetry restrictions’ and interactions with the solvation sphere improve the agreement between theory and experiment.

The benchmark calculations using various codes and input structures helped to select the most appropriate conditions for the simulations. The input parameters for the FDMNES XANES calculations were tested and defined. We found that the best agreement between theory and experiment was achieved using structure 3 [aqueous Gd(H-BTP)\textsubscript{3}], i.e. structure 3 is closer to the real structure than the other two.

The benchmark X-ray Raman spectroscopy studies demonstrate the applicability of this novel technique for investigations of liquid samples of partitioning systems at the N K-edge and K-absorption edges of other low-Z elements. No significant differences between the N K-edge spectrum for

![Figure 19](image1.png)

Figure 19
O K-edge spectra of isopropanol (a) and water (b). N K-edge spectrum of \textit{n}-Pr-BTP (c).

![Figure 20](image2.png)

Figure 20
N K-edge spectra of \textit{n}-Pr-BTP crystalline and in isopropanol solution.
the n-Pr-BTP molecule in the solid phase or solved in isopropanol are found. This result strongly suggests that the results obtained for the solid state complexes and ligands are relevant also for their liquid forms. Specific practical suggestions for further improvement of the experimental set-up are given based on experience gained over several experiments using the state-of-the-art spectrometers installed at the brightest synchrotrons. It is shown that N K-edge X-ray Raman spectroscopy investigations of N-donor ligands in solution are in general possible at the ID20 beamline, ESRF. However, the sample setup has to be further improved by using a capillary to allow a high signal-to-noise ratio; temperature-controlled cooling of the sample will reduce the evaporation of the solvent and possible radiation damage.

Acknowledgements

We would like to thank APS, BESSY and ESRF for granting beamtime. We thank the KIT light source for provision of beamtime at the WERA beamline operated by the Institute for Quantum Materials and Technologies (IOMT) and the INE-Beamline operated by the Institute for Nuclear Waste Disposal. We acknowledge the Karlsruhe Nano Micro Facility for Information (KNMFi) for provision of beamtime at the WERA beamline. We would like to thank the Institute for Beam Physics and Technology (IBPT) for the operation of the WERA beamline. We would like to thank the Institute for Nuclear Waste Disposal. We acknowledge the Karlsruhe Nano Micro Facility (KNMFi) for provision of beamtime at the WERA beamline. We thank K. Kvaschnina and J. Rothe for assistance with the experiments. We thank U. Müllerich for the synthesis of the 118-actinide complexes and ligands. Open access funding enabled and organized by Projekt DEAL.

Funding information

The following funding is acknowledged: Helmholtz Association, Grant VH-NG-734.

References

Asakura, H., Shishido, T., Fuchi, S., Teramura, K. & Tanaka, T. (2014). J. Phys. Chem. C, 118, 20881–20888.

Asakura, H., Shishido, T., Teramura, K. & Tanaka, T. (2015). J. Phys. Chem. C, 119, 8070–8077.

Banik, N. L., Schimmelpfennig, B., Marquardt, C. M., Brendebach, B., Geist, A. & Denecke, M. A. (2010). Dalton Trans. 39, 5117–5122.

Bartolomé, F., Krisch, M. & Kao, C. C. (1997). J. Chem. Soc. Dalton Trans., 463–483.

Bergmann, U., Wernet, P., Glatzel, P., Cavalleri, M., Pettersson, L. G. M., Nilsson, A. & Cramer, S. P. (2002). Phys. Rev. B, 66, 092107.

Berthet, J.-C., Miquel, Y., Iveson, P. B., Nierlich, M., Thürer, P., Madic, C. & Ephritikhine, M. (2002). J. Chem. Soc. Dalton Trans. pp. 3265–3272.

Bianconi, A., Marcelli, A., Dexpert, H., Karnatak, R., Kotani, A., Jo, T. & Petiau, J. (1987). Phys. Rev. B, 35, 806–812.

Blum, M., Weinhardt, L., Fuchs, O., Bärr, M., Zhang, Y., Weigand, M., Krause, S., Poopkantrata, S., Hofmann, T., Yang, W., Denlinger, J. D., Umbach, E. & Heske, C. (2009). Rev. Sci. Instrum. 80, 123102.

Boucher, C., Drew, M. G. B., Giddings, P., Harwood, L. M., Hudson, M. J., Iveson, P. B. & Madic, C. (2002). Inorg. Chem. Commun. 5, 596–599.

Bowron, D. T., Krisch, M. H., Barnes, A. C., Finney, J. L., Kaprolat, A. & Lorenzen, M. (2000). Phys. Rev. B, 62, R9223–R9227.

Bremner, A., Geist, A. & Panak, P. J. (2012). Dalton Trans. 41, 7582–7589.

Bremner, A., Geist, A. & Panak, P. J. (2013). Radiochim. Acta, 101, 285–292.

Bremner, A., Ruff, C. M., Girnt, D., Mülllich, U., Rothe, J., Roesky, P. W., Panak, P. J., Karpov, A., Mülller, T. J., Denecke, M. A. & Geist, A. (2012). Inorg. Chem. 51, 5199–5207.

Brouder, C., Juhn, A., Bordage, A. & Arrio, M.-A. (2008). J. Phys. Condens. Matter, 20, 455205.

Bunau, O. & Joly, Y. (2009). J. Phys. Condens. Matter, 21, 345501.

Carra, P. & Altarelli, M. (1990). Phys. Rev. Lett. 64, 1286–1288.

Carra, P., Fabrizio, M. & Thole, B. T. (1995). Phys. Rev. Lett. 74, 3700–3703.

Case, F. H. (1971). J. Heterocycl. Chem. 8, 1043–1046.

Centeno, M. A., Malet, P., Carrizosa, I. & Ondrizola, J. A. (2000). J. Phys. Chem. B, 104, 3310–3319.

Dallera, C., Annese, E., Rueff, J. P., Palenza, A., Vankö, G., Braicovich, L., Shukla, A. & Orioni, M. (2003). Phys. Rev. B, 68, 245114.

Dallera, C., Grioni, M., Palenza, A., Taguchi, M., Annese, E., Ghiringhelli, G., Tagliaferri, A., Brookes, N. B., Neisius, T. & Braicovich, L. (2004). Phys. Rev. B, 70, 085112.

Dallera, C., Krisch, M. H., Rogalev, A., Gauthier, C., Goulon, J., Sette, F. & Sole, A. (2000). Phys. Rev. B, 62, 7093–7097.

Dallera, C., Wessely, O., Colarieti-Tosti, M., Eriksson, O., Ahuja, R., Johansson, B., Katsnelson, M. I., Annese, E., Rueff, J. P., Vankö, G., Braicovich, L. & Grioni, M. (2006). Phys. Rev. B, 74, 081101.

Denecke, M. A., Rossberg, A., Panak, P. J., Weigl, M., Schimmel- pfennig, B. & Geist, A. (2005). Inorg. Chem. 44, 8418–8425.

Ekberg, C., Weigend, F., Treutler, O. & Ahrlehrs, R. (1997). Theoret. Chim. Acta, 97, 119–124.

Ekberg, C., Fernvik, A., Retegan, T., Skarnemark, G., Foreman, M. R. S., Hudson, M. J., Englund, S. & Nilsson, M. (2008). Radiochim. Acta, 96, 225–233.

Finkelstein, L. D., Postnikov, A. V., Efremova, N. N. & Kurmaev, E. Z. (1992). Mater. Lett. 14, 115–118.

Fister, T. T., Seidler, G. T., Hamner, C., Cross, J. O. & Rehr, J. J. (2009). Phys. Rev. B, 79, 174117.

Fister, T. T., Seidler, G. T., Wharton, L., Battle, A. R., Ellis, T. B., Cross, J. O., Macrander, A. T., Elam, W. T., Tyson, T. A. & Qian, Q. (2006). Rev. Sci. Instrum. 77, 063901.

Gallet, J. J., Mariot, J. M., Journel, L., Hague, C. F., Rogalev, A., Ogasawara, H., Kotani, A. & Sacchi, M. (1999). Phys. Rev. B, 60, 14128–14131.

Geist, A., Hill, C., Modolo, G., Foreman, M. R. S. J., Weigl, M., Gompper, K. & Hudson, J. M. (2006). Solvent Extr. Ion Exch. 24, 463–483.

Girnt, D., Roesky, P. W., Geist, A., Ruff, C. M., Panak, P. J. & Denecke, M. A. (2010). Inorg. Chem. 49, 9627–9635.

Glatzel, P., de Groot, F. M. F., Manoïlova, O., Grandjean, D., Weckhuysen, B. M., Bergmann, U. & Barrea, R. (2005). Phys. Rev. B, 72, 014117.
Vitova, T., Pidchenko, I., Fellhauer, D., Pruessmann, T., Bahl, S., Dardenne, K., Yokosawa, T., Schimmelpfennig, B., Altmaier, M., Denecke, M., Rothe, J. & Geckeis, H. (2018). Chem. Commun. 54, 12824–12827.

Walshe, A., Prüßmann, T., Vitova, T. & Baker, R. J. (2014). Dalton Trans. 43, 4400–4407.

Weigend, F., Häser, M., Patzelt, H. & Ahlrichs, R. (1998). Chem. Phys. Lett. 294, 143–152.

Weigl, M., Geist, A., Müllich, U. & Gompper, K. (2006). Solvent Extr. Ion Exch. 24, 845–860.

Wernet, P., Nordlund, D., Bergmann, U., Cavalleri, M., Odelius, M., Ogasawara, H., Näslund, L. A., Hirsch, T. K., Ojamae, L., Glatzel, P., Pettersson, L. G. M. & Nilsson, A. (2004). Science, 304, 995–999.

Yamaoka, H., Taguchi, M. M., Vlaicu, A., Oohashi, H., Yokoi, K., Horiguchi, D., Tochio, T., Ito, Y., Kawatsura, K., Yamamoto, K., Chainani, A., Shin, S., Shiga, M. & Wada, H. (2006). J. Phys. Soc. Jpn, 75, 034702.

Zimina, A., Dardenne, K., Denecke, M. A., Doronkin, D. E., Huttel, E., Lichtenberg, H., Mangold, S., Pruessmann, T., Rothe, J., Spangenberg, T., Steininger, R., Vitova, T., Geckeis, H. & Grunwaldt, J. D. (2017). Rev. Sci. Instrum. 88, 113113.

Zimina, A., Dardenne, K., Denecke, M. A., Grunwaldt, J. D., Huttel, E., Lichtenberg, H., Mangold, S., Pruessmann, T., Rothe, J., Steininger, R. & Vitova, T. (2016). J. Phys. Conf. Ser. 712, 012019.

Zubavichus, Y., Zharnikov, M., Yang, Y. J., Fuchs, O., Umbach, E., Heske, C. & Grunze, M. (2006). Langmuir, 22, 7241–7247.