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# Thermodynamics of the Eu(III)–Mg–SO<sub>4</sub>–H<sub>2</sub>O and Eu(III)–Na–SO<sub>4</sub>–H<sub>2</sub>O systems. Part I: solubility experiments and the full dissociation Pitzer model†

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The solubility of Eu(III) was investigated under undersaturated conditions in acidic, dilute to concentrated MgSO<sub>4</sub> and Na<sub>2</sub>SO<sub>4</sub> solutions at  $T = (22 \pm 2)$  °C. After attaining equilibrium conditions, solid phases were characterized by a multi-method approach, including X-ray diffraction (XRD), Raman and infrared (IR) spectroscopy, quantitative chemical analysis (ICP-OES) and thermogravimetric analysis (TG-DTA). A total of 45 solubility samples were investigated for the systems Eu<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>–MgSO<sub>4</sub>–H<sub>2</sub>O (19 samples) and Eu<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>–Na<sub>2</sub>SO<sub>4</sub>–H<sub>2</sub>O (26 samples). Eu<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>·8H<sub>2</sub>O(cr) was found to control the solubility of Eu(III) in all investigated MgSO<sub>4</sub> solutions, as well as in dilute Na<sub>2</sub>SO<sub>4</sub> systems. The transformation of Eu<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>·8H<sub>2</sub>O(cr) into the double salt Na<sub>2</sub>Eu<sub>2</sub>(SO<sub>4</sub>)<sub>4</sub>·2H<sub>2</sub>O(cr) was observed at  $m\text{Na}_2\text{SO}_4 > 0.01 \text{ mol kg}^{-1}$ . The latter phase is characterized by significantly lower solubility. Based on these experimental solubility measurements, thermodynamic and activity models were proposed based on the Pitzer equations considering the full dissociation of the Eu(III) species in MgSO<sub>4</sub> and Na<sub>2</sub>SO<sub>4</sub> aqueous solutions, *i.e.* deliberately excluding Eu(III)–sulfate complex formation. A combination of the geochemical calculation code PhreeSCALE and the parameter estimation code PEST was used to determine the values of solubility products ( $K_{s,0}^\circ$ ) and binary and ternary specific interaction parameters ( $\beta_{ij}^{(0)}$ ,  $\beta_{ij}^{(1)}$ ,  $C_{ij}^\beta$ ,  $\theta_{ik}$ ,  $\Psi_{ijk}$ ).

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## 1. Introduction

Europium belongs to the rare-earth elements (REEs), a group of metals comprising the lanthanide series (Ln), which plays an important role in human technology mainly in the field of energy. In addition to its technology-oriented properties, including the well-known Eu luminescence,<sup>1,2</sup> europium is often considered as an analogue of trivalent actinides relevant in the context of nuclear waste disposal, *e.g.* Am(III) and Pu(III).<sup>2–4</sup> Besides sharing the same valency, these trivalent lanthanides and actinides have similar structural properties (ionic radii and f-orbital electrons) leading to comparable

chemical behavior, including the formation of aqueous complexes of similar stability.<sup>5–8</sup>

The emplacement of nuclear waste in underground repositories is one of the options favored by the international community for the disposal of nuclear waste. In the event of water intrusion, aqueous systems involving radionuclides may form. In this context, accurate knowledge of the aqueous properties and solubility upper limit provides important inputs for the prediction of radionuclide behavior under repository conditions. The composition of these aqueous systems will be largely defined by the waste components, technical barriers and geological formation. For instance, the characteristic pore water of the Callovian–Oxfordian clay rock, selected in the French concept, has an average salinity of 4.3 g L<sup>−1</sup>, mostly due to its Na, Ca, Mg, Cl and SO<sub>4</sub> content.<sup>9–11</sup> In the Opalinus clay of the Mont-Terri underground laboratory in Switzerland, the salinity is slightly higher, with an average value of 20.7 g L<sup>−1</sup>.<sup>12</sup> Sulfate is an abundant component in natural groundwater and one of the most relevant anions (besides chloride) in brines, possibly inherited from ancient seawater or forms during salt rock formation, where sulfate concentrations of up to 0.2 M can be expected.<sup>13–15</sup> Thermodynamic

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models can help understand the chemical behavior of dilute to concentrated sulfate-bearing aqueous solutions, including the interactions and/or aqueous complexation reactions that involve such species, as well as the solubility of the corresponding salts.<sup>6,16–22</sup> Only a few studies considering the solubility of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  in the  $\text{Eu}_2(\text{SO}_4)_3\text{--H}_2\text{O}$  binary system are reported in the literature.<sup>3,23,24</sup> Amongst these studies, the work published by Rard<sup>3</sup> stands out, involving isopiestic and solubility experiments in the absence of any background electrolyte. However, data on the solubility of  $\text{Eu}(\text{III})$  in dilute to concentrated  $\text{Eu}_2(\text{SO}_4)_3\text{--Na}_2\text{SO}_4\text{--H}_2\text{O}$  and  $\text{Eu}_2(\text{SO}_4)_3\text{--MgSO}_4\text{--H}_2\text{O}$  systems are not available so far. The study by Keyes and James<sup>25</sup> focused on the solubility of  $\text{Sm}(\text{III})$  in the ternary system  $\text{Sm}_2(\text{SO}_4)_3\text{--Na}_2\text{SO}_4\text{--H}_2\text{O}$ . The authors observed the formation of a hydrated salt in the form of  $\text{Na}_2\text{Sm}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  at  $\text{Na}_2\text{SO}_4$  concentrations above  $\sim 0.01$  M. Some studies with different lanthanides reporting the formation of double salts with  $\text{Na}^+$  are also available in the literature,<sup>25–28</sup> but they were performed at low  $\text{Na}_2\text{SO}_4$  concentrations and did not conclude any solubility information for a supposed  $\text{Na}_2\text{Ln}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  phase. The lack of relevant experimental solubility data makes it difficult to understand the behavior of  $\text{Eu}(\text{III})$  and feed the thermodynamic models used to reproduce and estimate such properties. This fact is emphasized by the work of Das *et al.*,<sup>26</sup> showing that in contrast to La, Ce, Pr, Nd and Sm, the experimental data for  $\text{Eu}(\text{III})\text{--SO}_4^{2-}$  are very scarce.

In the context of a collaborative project between BRGM, KIT and Andra, the present work aimed to acquire experimental data in order to assess the chemical behavior of the two ternary systems  $\text{Eu}_2(\text{SO}_4)_3\text{--Na}_2\text{SO}_4\text{--H}_2\text{O}$  and  $\text{Eu}_2(\text{SO}_4)_3\text{--MgSO}_4\text{--H}_2\text{O}$ , including the solubility of europium in dilute to concentrated aqueous solutions of sodium and magnesium sulfate ( $0\text{--}2.85$  mol  $\text{kg}^{-1}$  of water) at room temperature. In addition, assuming full dissociation of the dissolved electrolytes, a set of Pitzer-specific interaction parameters and solubility products are proposed for describing the solubility of the  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  and  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  salts in  $\text{Na}_2\text{SO}_4$  aqueous solutions and  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  in  $\text{MgSO}_4$  aqueous solutions. Part II of this work is presented in a separate contribution, involving spectroscopic characterization of  $\text{Eu}(\text{III})$  in the aqueous phase and including thermodynamic and (SIT, Pitzer) activity models specifically accounting for  $\text{Eu}(\text{III})\text{--SO}_4$  aqueous complexes.

## 2. Experimental

### 2.1. Chemicals

All solutions were prepared with ultra-pure water and purified with a Milli-Q academic apparatus (Merck Millipore,  $18.2$  M $\Omega$  cm,  $22 \pm 2$  °C, pore size  $0.22$   $\mu\text{m}$ ). Magnesium sulfate heptahydrate ( $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$ , p.a., 99.5 wt%) and europium(III) sulfate octahydrate ( $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$ , p.a., 99.9 wt%) were purchased from ThermoFisher Scientific. Anhydrous sodium sulfate ( $\text{Na}_2\text{SO}_4$ , p.a., >99 wt%) was obtained from Merck.

### 2.2. pH measurements

Proton concentration,  $\text{pH}_m$  ( $\text{pH}_m = -\log[\text{H}^+]$  in molal units, mol  $\text{kg}^{-1}$ ), was measured using a combination of pH electrodes (ROSS Orion, with 3 M KCl as the filler solution) calibrated against pH standards (pH 2–7, Merck). The  $\text{pH}_m$  values were obtained from the measured  $\text{pH}_{\text{exp}}$  values considering  $\text{pH}_m = \text{pH}_{\text{exp}} + A_m$ , where  $A_m$  is a correction factor entailing the activity coefficient of  $\text{H}^+$  and the liquid junction potential of the electrode at a given background electrolyte concentration.  $A_m$  values for  $\text{Na}_2\text{SO}_4$  systems were taken as reported by Duckworth *et al.*<sup>29</sup> (work performed between 0.1 and 1.75 mol  $\text{kg}^{-1}$  of  $\text{Na}_2\text{SO}_4$ ).  $A_m$  values for the  $\text{MgSO}_4$  aqueous systems are not available in the literature, and experimental values,  $\text{pH}_{\text{exp}}$ , are reported instead.

### 2.3. Solubility experiments

Solubility experiments were performed using well-characterized  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  as the starting material. Batch samples were equilibrated at  $T = (22 \pm 2)$  °C in an independent series of binary aqueous solutions with different concentrations of  $\text{Na}_2\text{SO}_4$  in the range  $0.000\text{--}1.784$  mol  $\text{kg}^{-1}$  (20 samples) or of  $\text{MgSO}_4$  in the range  $0.006\text{--}2.854$  mol  $\text{kg}^{-1}$  (19 samples). Above the  $\text{Na}_2\text{SO}_4$  concentration of  $0.015$  mol  $\text{kg}^{-1}$ , the  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  double salt formed and was recovered for a subsequent series of experiments. The subsequent series of batch solubility experiments used the  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  double salt and was prepared with a concentration of  $\text{Na}_2\text{SO}_4$  ranging between  $0.015$  and  $0.122$  mol  $\text{kg}^{-1}$  (6 samples). All samples were subjected to constant agitation until thermodynamic equilibrium was achieved, which was assumed after repeated measurements with a constant total  $\text{Eu}(\text{III})$  concentration.  $\text{Eu}(\text{III})$  concentration was measured by ICP-OES (inductively coupled plasma optical emission spectroscopy, PerkinElmer Optima 8300 DV) after phase separation by ultrafiltration ( $10$  kDa  $\approx 2$  nm, Pall Life Science). Aliquots of the original samples were diluted with 2%  $\text{HNO}_3$  before ICP-OES measurements. The accuracy of ICP-OES measurements was  $\pm 2\text{--}5\%$ .

### 2.4. Solid-phase characterization

Part of the solid phase of the samples was collected and subjected to several washing and centrifugation steps, with the aim of washing out the background electrolyte. The solid samples of the system containing  $\text{Na}_2\text{SO}_4$  were subjected to 5 successive washing steps with absolute ethanol followed by centrifugation at  $10,000$  g (5 minutes). The samples of solutions containing  $\text{MgSO}_4$  were subjected to 8 washing and centrifugation steps using a solution of absolute ethanol at four dilution ratios, namely: two steps the absolute temperature with a ratio of 1 : 5, two steps at 1 : 3, two steps at 1 : 2 and two steps with absolute ethanol only. Once the recovered solid was completely dry, it was subjected to all analyses for solid characterization, namely X-ray diffraction (XRD), Raman and infrared (IR) spectrometry, quantitative chemical analysis and thermogravimetric analysis (TG-DTA) before and after solubility experiments.



XRD measurements were performed on an Empyrean diffractometer (Malvern-Panalytical) equipped with a Cu tube in the range 5–90° 2theta with 0.0131° 2theta steps. The CuK $\beta$  radiation was filtered with a Bragg–Brentano HD device mounted on the primary beam path. Divergent slits of 1/8°, anti-scattering slits of 1/2° and soller slits of 0.04 Rad were used. The detection was performed with a multi-strip PIXcel 3D detector covering a 2theta of 3.348° simultaneously with 255 single channels. The samples mixed with ethanol were placed on Si sample holders. The data were evaluated with HighscorePlus v.5 with integrated ICDD2004 and COD2021 databases.

Thermogravimetry (TG) and differential thermal analysis (DTA) were performed simultaneously under a N<sub>2</sub> atmosphere using a Netzsch instrument (model STA409C Jupiter) with a DSC (differential scanning calorimetry) measuring head. Measurements were performed with 10–50 mg of the solid phase, and the heating rate was 10 K min<sup>-1</sup> up to 1200 °C.

Raman spectrometry was performed on an Alpha 300R instrument (WITec, Ulm, Germany) equipped with a Zeiss microscope. A 532 nm laser with tunable power in 0.1 mW steps (TruePower) was used as an excitation source focused on 50× (NA = 0.75) or 100× (NA = 0.9) objectives. Typical laser power was 20 mW (measured on the sample). The measurements as single spots were performed with 1800 L mm<sup>-1</sup> and 600 L mm<sup>-1</sup> gratings, providing spectral resolution better than 1 cm<sup>-1</sup> and 3 cm<sup>-1</sup>, respectively. Due to strong Eu<sup>3+</sup> luminescence, the spectra were recorded with very short acquisition times varying between 0.01 and 1 s, and 10 scans. Raman signals were recorded using a back illuminated CCD with 96% quantum efficiency (QE). The spectra were evaluated with Project 5.3+ software (WITec).

IR spectrometry was performed in ATR mode on a Tensor II spectrometer (Bruker Optics, Ettlingen, Germany) equipped with a deuterated triglycine sulfate (DTGS) detector and a Golden Gate ATR cell with a diamond crystal (Specac Ltd, Orpington, UK). Spectra were acquired in the range 400–4000 cm<sup>-1</sup> with 64 scans and a spectral resolution of 2 cm<sup>-1</sup>. OPUS 8 software was used for the acquisition and evaluation of the IR spectra.

Quantitative chemical analysis was performed by ICP-OES with a PerkinElmer Optima 8300 DV equipment to determine the Eu, S and Na content in the solid. The solid phase was dissolved in 7–10 mL of 2% HNO<sub>3</sub> and the solution was used for quantitative chemical analysis.

### 3. Thermodynamic modelling

The free energy change of a reaction,  $\Delta G$ , is related to the standard free energy change,  $\Delta G^\circ$ , and the ion activity product,  $Q$ , according to the following classical equation:

$$\Delta G = \Delta G^\circ + RT \ln Q \quad (1)$$

where  $R$  is the ideal gas constant (8.314 J K<sup>-1</sup> mol<sup>-1</sup>) and  $T$  is the absolute temperature (in K). The ionic activity product,  $Q$ ,

is the numerical value of the mass action expression for the system:

$$Q = \prod_i a_i^{\nu_i} \quad (2)$$

$\nu_i$  is the algebraic stoichiometric coefficient of species  $i$  in the dissolution/precipitation reaction and  $a_i$  is its chemical activity (unitless), with  $a_i = \frac{m_i}{m_0} \cdot \gamma_i$ , where  $m_i$  and  $\gamma_i$  are the molality (mol kg<sup>-1</sup>) and the activity coefficient (unitless) of  $i$ , respectively.  $m_0$  is a reference molality, set to 1 mol kg<sup>-1</sup>, and is thus generally disregarded, like in the following.

By defining the equilibrium constant  $K$  of the reaction according to  $\Delta G^\circ = -RT \ln K$ , eqn (1) becomes:

$$\Delta G = RT \ln \frac{Q}{K} \quad (3)$$

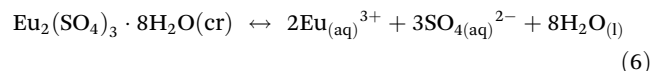
For a dissolution/precipitation reaction,  $K$  represents the solubility product at infinite dilution and can be denoted as  $K_{s,0}^\circ$ . Generally, when considering a dissolution/precipitation reaction, equilibrium is not reached instantaneously because of kinetic limitations. Out of equilibrium,  $\Delta G \neq 0$  J and the saturation ratio, SR, can be defined by eqn (4):

$$SR = \frac{Q}{K_{s,0}^\circ} \quad (4)$$

SR is commonly used to measure the deviation from equilibrium for a given mineral–solution system. SR > 1 represents supersaturation and thus the tendency towards precipitation. In contrast, SR < 1 represents undersaturation, indicating that, if not present, the solid cannot form or, if in contact with the solution, the solid has a tendency to dissolve. SR = 1 means that  $\Delta G = 0$  and, hence, that equilibrium is reached. In this case, eqn (4) reduces to the mass action law:

$$K_{s,0}^\circ = Q \quad (5)$$

The saturation ratio is thus of great importance in determining the equilibrium constant of minerals when the solubility is actually characterized. At the thermodynamic equilibrium, the expression of the solubility product of Eu<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>·8H<sub>2</sub>O(cr) at infinite dilution,  $K_{s,0}^\circ\{\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})\}$ , is defined as follows:



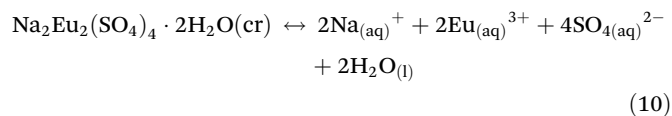
$$K_{s,0}^\circ\{\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})\} = a_{\text{Eu}^{3+}}^2 a_{\text{SO}_4^{2-}}^3 a_{\text{H}_2\text{O}}^8 \quad (7)$$

$$K_{s,0}^\circ\{\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})\} = m_{\text{Eu}^{3+}}^2 m_{\text{SO}_4^{2-}}^3 \gamma_{\text{Eu}^{3+}}^2 \gamma_{\text{SO}_4^{2-}}^3 a_{\text{H}_2\text{O}}^8 \quad (8)$$

$$\begin{aligned} & \log_{10} K_{s,0}^\circ\{\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})\} \\ &= \log_{10} \left( m_{\text{Eu}^{3+}}^2 m_{\text{SO}_4^{2-}}^3 \right) + \log_{10} \left( a_{\text{H}_2\text{O}}^8 \right) + \log_{10} \left( \gamma_{\text{Eu}^{3+}}^2 \gamma_{\text{SO}_4^{2-}}^3 \right) \end{aligned} \quad (9)$$



Analogously, the solubility product of the double salt  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  at infinite dilution,  $K_{s,0}^\circ\{\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})\}$ , can be defined as follows:



$$K_{s,0}^\circ\{\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})\} = a_{\text{Na}^+}^2 a_{\text{Eu}^{3+}}^2 a_{\text{SO}_4^{2-}}^4 a_{\text{H}_2\text{O}}^2 \quad (11)$$

$$K_{s,0}^\circ\{\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})\} = m_{\text{Na}^+}^2 m_{\text{Eu}^{3+}}^2 m_{\text{SO}_4^{2-}}^4 \gamma_{\text{Na}^+}^2 \gamma_{\text{Eu}^{3+}}^2 \gamma_{\text{SO}_4^{2-}}^4 a_{\text{H}_2\text{O}}^2 \quad (12)$$

$$\begin{aligned} \log_{10} K_{s,0}^\circ\{\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})\} \\ = \log_{10} \left( m_{\text{Na}^+}^2 m_{\text{Eu}^{3+}}^2 m_{\text{SO}_4^{2-}}^4 \right) + \log_{10} \left( a_{\text{H}_2\text{O}}^2 \right) \\ + \log_{10} \left( \gamma_{\text{Na}^+}^2 \gamma_{\text{Eu}^{3+}}^2 \gamma_{\text{SO}_4^{2-}}^4 \right) \end{aligned} \quad (13)$$

The activity coefficient  $\gamma_i$  of an aqueous species  $i$  is given by the derivative of the excess Gibbs free energy  $G^{\text{ex}}$  of the solution with respect to the number of moles of species  $i$ . In the present work, the excess Gibbs free energy is calculated according to the Pitzer equations,<sup>30,31</sup> which involve semi-empirical parameters that represent the specific interactions between solutes (see the ESI†).

The total dissociation of the dissolved electrolytes is assumed analogously to several previous studies under high saline conditions involving Pitzer equations.<sup>19,32,33</sup> Hence, full dissociation has been previously considered for other 3 : 2 type electrolytes like  $\text{Al}_2(\text{SO}_4)_3$ ,<sup>34,35</sup>  $\text{Cr}_2(\text{SO}_4)_3$ ,  $\text{Fe}_2(\text{SO}_4)_3$  and  $\text{La}_2(\text{SO}_4)_3$ .<sup>36–38</sup> In the present work, this assumption indicates that only  $\text{Eu}^{3+}$ ,  $\text{Na}^+$ ,  $\text{Mg}^{2+}$  and  $\text{SO}_4^{2-}$  aqueous species are considered in the chemical systems studied (in addition to  $\text{H}_2\text{O}$ ,  $\text{H}^+$  and  $\text{OH}^-$  water species). Neither aqueous complexes nor ion pairs are explicitly accounted for in the following: they are represented by specific interactions, instead.

It should however be noted that previous studies have reported the formation of the  $\text{Eu}(\text{SO}_4)^+$ ,  $\text{Eu}(\text{SO}_4)_2^-$  and  $\text{Eu}(\text{SO}_4)_3^{3-}$  aqueous complexes.<sup>4,39–48</sup> Although the consideration of these aqueous complexes allows a more realistic description of the aqueous phase, the number of interaction parameters required for accurate model calculations using the Pitzer formalism increases significantly, with the consequent risk of over-parametrization of the investigated system. These aspects are further addressed in Part II of this work (F. dos Santos *et al.*;<sup>49</sup> this journal).

## 4. Parametrization procedure

To reproduce the solubility behavior of the  $\text{Eu}_2(\text{SO}_4)_3\text{-MgSO}_4\text{-H}_2\text{O}$  and  $\text{Eu}_2(\text{SO}_4)_3\text{-Na}_2\text{SO}_4\text{-H}_2\text{O}$  systems, the parameters to be determined are:

- The solubility products at infinite dilution,  $\log_{10} K_{s,0}^\circ\{\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})\}$  and  $\log_{10} K_{s,0}^\circ\{\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})\}$ ;

- The binary ( $\beta_{ij}^{(0)}$ ,  $\beta_{ij}^{(1)}$ ,  $C_{ij}^\phi$ ,  $\theta_{ik}$ ) and ternary ( $Y_{ijk}$ ) specific interaction parameters of the Pitzer model (see the ESI†).

Estimation of these parameters not only requires the experimental solubility data acquired in the present work for the two aforementioned salts, but also the interaction parameters for the binary systems  $\text{MgSO}_4\text{-H}_2\text{O}$  and  $\text{Na}_2\text{SO}_4\text{-H}_2\text{O}$ , which were determined previously by Lach *et al.*<sup>50</sup> In the present case, the parameterization procedure includes two successive steps in order to develop a consistent model for the two systems investigated. First, the specific interaction parameters and the solubility product  $\log_{10} K_{s,0}$  were determined for the  $\text{Eu}_2(\text{SO}_4)_3\text{-MgSO}_4\text{-H}_2\text{O}$  system, which is more constrained than the  $\text{Eu}_2(\text{SO}_4)_3\text{-Na}_2\text{SO}_4\text{-H}_2\text{O}$  system due to the formation of the double-salt in the latter case. In the second step, the parameters obtained in the first step were kept unchanged and the parameters specific to the  $\text{Eu}_2(\text{SO}_4)_3\text{-Na}_2\text{SO}_4\text{-H}_2\text{O}$  system were determined.

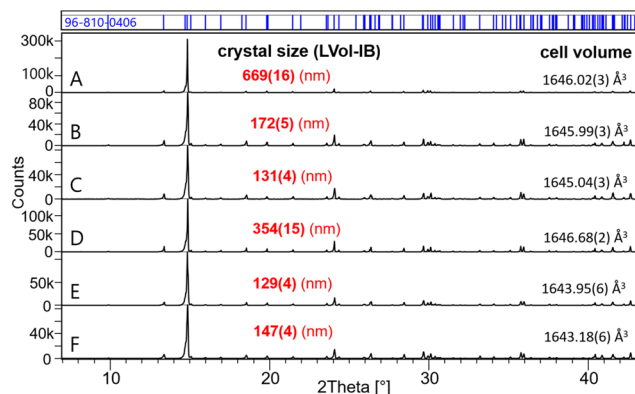
Optimizations were performed by coupling the geochemical calculation code PhreeSCALE<sup>50</sup> with the parameter estimation software PEST<sup>51</sup> according to the following principle.<sup>16,50,52</sup> PEST calls PhreeSCALE to calculate the target properties of the chemical systems of interest (in this work, the saturation ratio, SR), using an initial set of guess values for the parameters to be estimated (binary and ternary interaction parameters and solubility products). The calculation results are compared to the corresponding experimental values by calculating the objective function that characterizes the deviation with respect to the experimental data. If the convergence criterion is not satisfactory, a new series of calculations is started with a new set of parameters. The optimization is carried out by successive iterations of this procedure until the convergence criterion is satisfactory or the objective function does not evolve any further. In total, 45 solubility data points were used to estimate 9 parameters.

## 5. Results and discussion

### 5.1. Solid phase characterization

Fig. 1 shows the powder XRD patterns of the samples A–F equilibrated in  $\text{MgSO}_4$  solutions (see Table 1 for a description of the sample conditions). In all these samples,  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  was found to be the only solid phase controlling the solubility of  $\text{Eu}(\text{III})$ . A Rietveld refinement was performed using the structure data of Xu *et al.*<sup>53</sup> The size of the coherent scattering domains was calculated by the double Voigt approach based on the integral breadth using Lorentzian and Gaussian type component convolutions.<sup>54</sup> From sample A to sample C, a gradual decrease in the crystal size from 670 to 130 nm is observed, correlating with a small decrease in the volume of the unit cell. Sample D shows once again a greater crystal size of around 350 nm. Samples E and F further show significantly smaller crystal sizes of 130 and 147 nm, respectively. A similar correlation is observed considering the volume of the unit cell. The overall large crystal size of all investigated solids reflects





**Fig. 1** XRD patterns of samples A–F (in  $\text{MgSO}_4$  aqueous solutions). All patterns match the COD 96-810-0406 for  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$ . The calculated crystal size and the cell volume are also shown.

the high crystallinity of the solid phases controlling the solubility.

Fig. 2 shows the powder XRD patterns of samples G–N equilibrated in  $\text{Na}_2\text{SO}_4$  solutions (0.00–1.56  $\text{mol kg}^{-1}$ ). Samples G, H and I consist entirely of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$ , whereas the double-salt  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  is the only solid phase identified in samples J–N. Samples M and N show the additional presence of thenardite  $\text{Na}_2\text{SO}_4(\text{vi})$  and a minor fraction of  $\text{Na}_2\text{SO}_4(\text{III})$ . The latter is known as a product of the thermal treatment of the former. With increasing  $\text{Na}_2\text{SO}_4$  concentration from 0.1  $\text{mol kg}^{-1}$  (sample J) to 0.35  $\text{mol kg}^{-1}$  (sample L), the formed  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  shows an increasing crystallinity with the size of the coherent scattering domains varying from 170 nm (sample J), 180 nm (sample K) to 390 nm (sample L). The samples with excess  $\text{Na}_2\text{SO}_4$  (samples M and N) show again smaller crystal sizes of 125 and

145 nm, respectively. As in the case of  $\text{MgSO}_4$  systems, these results confirm the high crystallinity of the solid phases controlling the solubility of  $\text{Eu}(\text{III})$  in  $\text{Na}_2\text{SO}_4$  systems. It should be noted that for solid phases above 100 nm, the impact of the particle size on the  $\Delta_r G^\circ$  of the solid phase (and by extension on its solubility product) is considered negligible.<sup>55</sup>

The IR spectra of samples A–F are shown in Fig. 3. All spectra show typical features of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$ . The band assignment was made according to Denisenko *et al.*<sup>56–58</sup> and Ram.<sup>59</sup> The internal vibrations of the  $\text{SO}_4$  tetrahedra are observed as: (1) broad bands in the range 1050–1200  $\text{cm}^{-1}$  with the main peaks at 1076 and 1126  $\text{cm}^{-1}$  assigned to  $\nu_3(\text{SO}_4)$ , (2) a sharp band at 1000  $\text{cm}^{-1}$  with shoulders at 984 and 940  $\text{cm}^{-1}$  ( $\nu_1(\text{SO}_4)$ ), and (3) small bands at 638 and 650  $\text{cm}^{-1}$ , assigned to  $\nu_4(\text{SO}_4)$ . The observed frequencies match very well those reported by Buyer *et al.*<sup>60</sup> for  $\text{Y}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}$ , although their assignment of the bands at 688 and 743  $\text{cm}^{-1}$  to  $\nu_4(\text{SO}_4)$  raises some doubts. We favor the assignment of the similar bands at 683 and 745  $\text{cm}^{-1}$  in our spectra to  $\text{H}_2\text{O}$  vibrations as reported by Ram,<sup>59</sup> given the broadness of these bands. The vibrations due to  $\text{H}_2\text{O}$  are manifested by: (1) a very sharp band at 1637  $\text{cm}^{-1}$  (H–O–H bending,  $\nu_2$ ), (2) several broad bands with low intensity at 683, 745 and 803  $\text{cm}^{-1}$  assigned to  $\text{H}_2\text{O}$  vibrations (rocking, twisting), and (3) broad bands in the OH stretching region (3000–3600  $\text{cm}^{-1}$ ) centered at 3225, 3300, 3340, and 3452  $\text{cm}^{-1}$ . The latter confirm the presence of four distinct  $\text{H}_2\text{O}$  coordinating Eu along with four oxygens from  $\text{SO}_4$  tetrahedra. Once again, there is a striking similarity to the frequencies of the OH stretching vibrations reported by Buyer *et al.*<sup>60</sup>

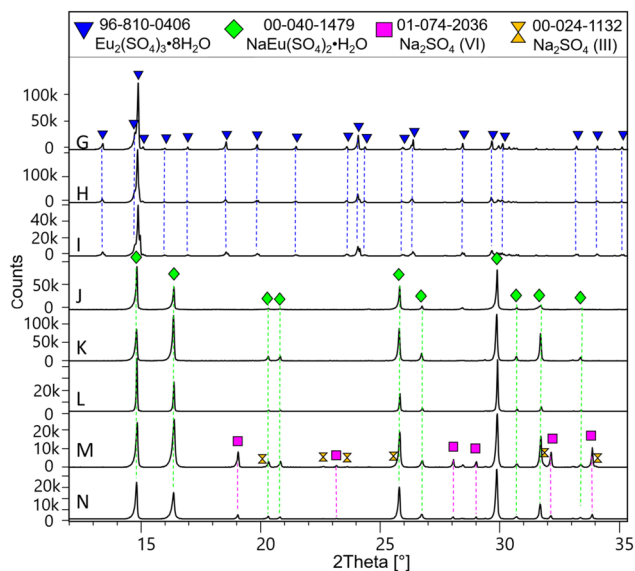
The trivalent  $\text{Eu}^{3+}$  ion shows very strong “red” luminescence, which obscures the Raman spectra when excited with a 532 nm laser. Due to the strong dependence of the number and intensity of the emission lines on the site symmetry and

**Table 1** Main characteristics of the selected solid samples equilibrated in  $\text{Na}_2\text{SO}_4$  and  $\text{MgSO}_4$  systems according to various characterization techniques: XRD (phase identification), Raman and IR spectrometry (phase identification), ICP-OES (S : Eu or Na : Eu ratio) and TG-DTA (number of hydration water molecules,  $n\text{H}_2\text{O}$ )

Samples	$m\text{MgSO}_4$ ( $\text{mol kg}^{-1}$ )	$m\text{Na}_2\text{SO}_4$ ( $\text{mol kg}^{-1}$ )	XRD		Raman/IR		ICP-OES		TG-DTA $n\text{H}_2\text{O}$
			Single salt <sup>a</sup>	Double salt <sup>b</sup>	Single salt <sup>a</sup>	Double salt <sup>b</sup>	S : Eu	Na : Eu	
A	0.006		X <sup>c</sup>		X <sup>c,d</sup>		1.62 ± 0.08		7.98 ± 0.03
B	0.052		X <sup>c</sup>		X <sup>c,d</sup>		1.60 ± 0.08		
C	0.804		X <sup>c</sup>		X <sup>c,d</sup>		1.66 ± 0.08		
D	1.310		X <sup>c</sup>		X <sup>c,d</sup>		1.60 ± 0.08		
E	2.490		X <sup>c</sup>		X <sup>c,d</sup>		1.61 ± 0.08		
F	2.695		X <sup>c</sup>		X <sup>c,d</sup>		1.81 ± 0.08		
G		0.000	X <sup>c</sup>		X <sup>c,e</sup>		1.55 ± 0.08		8.03 ± 0.07
H		0.008	X <sup>c</sup>		X <sup>c,e</sup>		1.58 ± 0.08		
I		0.010	X <sup>c</sup>		X <sup>c,e</sup>		1.64 ± 0.08		
J		0.100		X <sup>c</sup>		X <sup>c,e</sup>		1.10 ± 0.06	2.08 ± 0.13
K		0.122		X <sup>c</sup>		X <sup>c,e</sup>		1.07 ± 0.05	
L		0.035		X <sup>c</sup>		X <sup>c,e</sup>		1.07 ± 0.05	
M		0.643		X <sup>c</sup>		X <sup>c,e</sup>		6.59 <sup>f</sup> ± 0.33	
N		1.560		X <sup>c</sup>		X <sup>c,e</sup>		7.03 <sup>f</sup> ± 0.35	

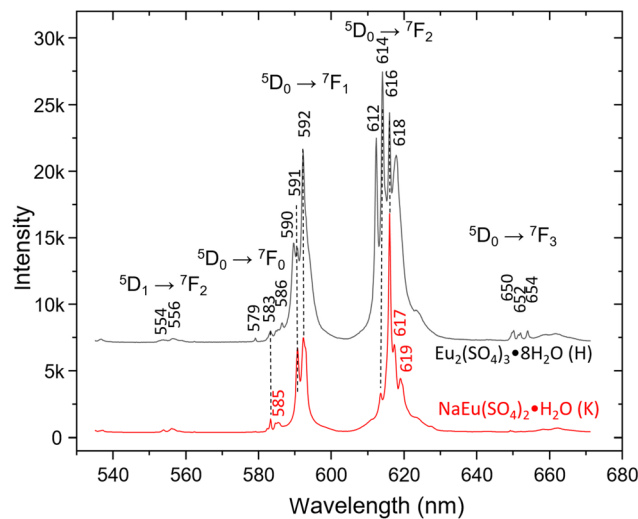
<sup>a</sup>  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$ . <sup>b</sup>  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$ . <sup>c</sup> Identified phase. <sup>d</sup> By IR spectrometry. <sup>e</sup> By Raman spectrometry. <sup>f</sup> Presence of  $\text{Na}_2\text{SO}_4$  solid.





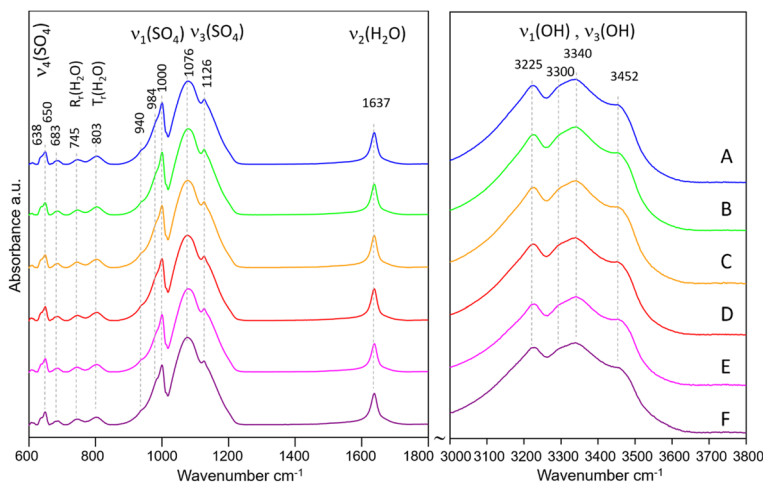
**Fig. 2** XRD patterns of samples G–N (in  $\text{Na}_2\text{SO}_4$  aqueous solutions). The chemical formula  $\text{NaEu}(\text{SO}_4)_2 \cdot \text{H}_2\text{O}$  stands for the  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  solid phase.

environment of the Eu ion in different structures,<sup>1</sup> it could be used for identification purposes. Fig. 4 shows the emission spectra of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  (sample H) and  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  (sample K) excited by the 532 nm laser recorded with 600  $\text{L mm}^{-1}$  grating in the range 530–680 nm. The hypersensitive  $^5\text{D}_0 \rightarrow ^7\text{F}_2$  transition between 610 and 630 nm dominates both spectra, followed by  $^5\text{D}_0 \rightarrow ^7\text{F}_1$  transition lines. The spectrum of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  shows strong resemblance to that published by Xu *et al.*<sup>53</sup> with the highest intensity of the 614 nm line. On the other side, the spectrum of  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  resembles the one published by Wu and Liu<sup>61</sup> and matches perfectly that reported by Buyer *et al.*<sup>60</sup> for  $\text{Na}_2\text{Y}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  doped with  $\text{Eu}^{3+}$  at 300 K.



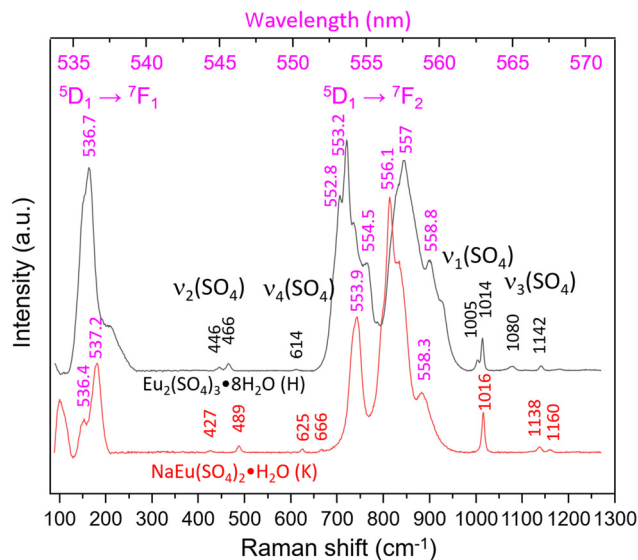
**Fig. 4** Emission spectra of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  (sample H) and  $\text{NaEu}(\text{SO}_4)_2 \cdot \text{H}_2\text{O}(\text{cr})$  (sample K) excited by a 532 nm laser. The chemical formula  $\text{NaEu}(\text{SO}_4)_2 \cdot \text{H}_2\text{O}$  stands for the  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  solid phase.

Fig. 5 shows the Raman spectra of both aforementioned samples taken with 1800  $\text{L mm}^{-1}$  grating in the range 100–1300  $\text{cm}^{-1}$  (530–570 nm). Once again the spectra are dominated by luminescence, but higher excited states most probably  $^5\text{D}_1 \rightarrow ^7\text{F}_1$  and  $^5\text{D}_1 \rightarrow ^7\text{F}_2$ , to our knowledge, have not been reported so far. The Raman bands typical of the internal vibrations of  $\text{SO}_4^{2-}$  are visible and allow the differentiation between the  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  and  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  systems. The former shows bands at 446 and 466  $\text{cm}^{-1}$  ( $\nu_2$ ), 614  $\text{cm}^{-1}$  ( $\nu_4$ ), 1005  $\text{cm}^{-1}$ , 1014  $\text{cm}^{-1}$  ( $\nu_1$ ) and 1080  $\text{cm}^{-1}$ , 1142  $\text{cm}^{-1}$ , 1180  $\text{cm}^{-1}$  ( $\nu_3$ ) in accordance with Ram.<sup>59</sup> In the spectrum of  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$ , the corresponding modes give rise to bands at 427, 489  $\text{cm}^{-1}$  ( $\nu_2$ ), 625, 666  $\text{cm}^{-1}$  ( $\nu_4$ ), 1016  $\text{cm}^{-1}$  ( $\nu_1$ ) and 1138, 1160  $\text{cm}^{-1}$  ( $\nu_3$ ). The number and positions of these bands show strong simi-



**Fig. 3** IR spectra of samples A–F (in  $\text{MgSO}_4$  solutions).





**Fig. 5** Raman spectra of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  (sample H) and  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  (sample K) excited by a 532 nm laser. The most intense “bands” are the emission lines from higher excited states. The chemical formula  $\text{NaEu}(\text{SO}_4)_2 \cdot \text{H}_2\text{O}(\text{K})$  stands for the  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  solid phase.

larity to those reported by Buyer *et al.*<sup>60</sup> for  $\text{Na}_2\text{Y}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  doped with  $\text{Eu}^{3+}$  with maximal shifts of 2–4  $\text{cm}^{-1}$ .

Quantitative evaluation of the TG-DTA data obtained for the initial solid of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  indicates a content of  $(8.01 \pm 0.03)$  mol of  $\text{H}_2\text{O}$  per mol of solid. After completing the solubility experiments, the water content quantified by TG-DTA for the solid samples equilibrated in  $\text{MgSO}_4$  aqueous solutions was  $(7.98 \pm 0.03)$  mol of  $\text{H}_2\text{O}$  per mol of solid. As for the samples G, H, I of the  $\text{Na}_2\text{SO}_4$  aqueous system (in the region with no phase change, *i.e.*  $\text{Na}_2\text{SO}_4 \leq 0.01$  mol  $\text{kg}^{-1}$ ), a water content of  $(8.03 \pm 0.07)$  mol of  $\text{H}_2\text{O}$  per mol of solid was identified. This confirms that in these samples the  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  phase actually controlled equilibrium – as was already shown by XRD and Raman spectroscopy measurements. In the samples where the presence of  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  was identified by XRD and Raman spectroscopy, the measured water content was  $(2.08 \pm 0.13)$  mol of  $\text{H}_2\text{O}$  per mol of solid, consistent with the formation of the double salt in the samples J to N, with concentrations of  $\text{Na}_2\text{SO}_4 > 0.01$  mol  $\text{kg}^{-1}$ .

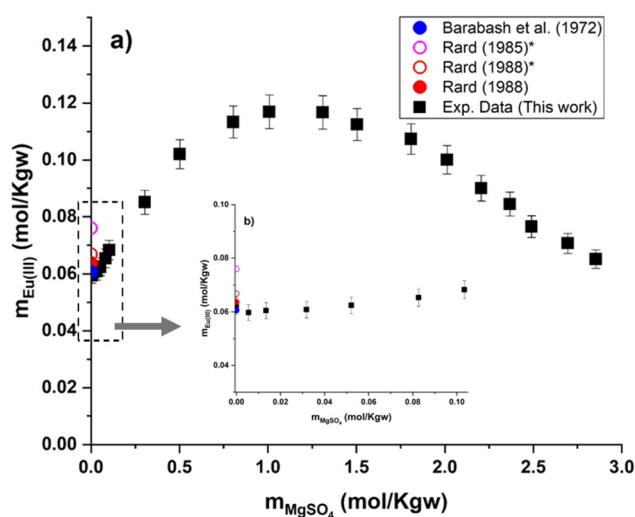
Furthermore, in the  $\text{Eu}_2(\text{SO}_4)_3$ – $\text{Na}_2\text{SO}_4$ – $\text{H}_2\text{O}$  system, the S : Eu ( $\sim 1.5$ ) and Na : Eu ( $\sim 1.0$ ) ratios determined by ICP-OES and quantitative chemical analyses on solid samples are in excellent agreement with the XRD data. They support the predominance of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  (samples G to I) and  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  (samples J to N) at room temperature (Table 1). In samples M and N, with higher concentrations of  $\text{Na}_2\text{SO}_4$ , namely 0.643 and 1.560 mol  $\text{kg}^{-1}$ , the Na : Eu ratios are well above 1, suggesting the presence of sodium sulfate in the solid phase. This is confirmed in Fig. 2, where the XRD pattern shows peaks of thenardite  $\text{Na}_2\text{SO}_4(\text{VI})$  and  $\text{Na}_2\text{SO}_4(\text{III})$ . In the  $\text{Eu}_2(\text{SO}_4)_3$ – $\text{MgSO}_4$ – $\text{H}_2\text{O}$  system, the S : Eu ratio values

found confirm the presence of the  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  phase only (samples A to F), in agreement with the other analyses carried out on the corresponding solid phases.

The main results of the solid phase characterization obtained by XRD, Raman and IR spectrometry, quantitative chemical analysis and TG-DTA are summarized in Table 1.

## 5.2. Solubility measurements

Experimental results of  $\text{Eu}(\text{III})$  solubility at different aqueous concentrations of  $\text{MgSO}_4$  and  $\text{Na}_2\text{SO}_4$  are shown in Fig. 6 and 7. Experimental solubility data, as well as the pH and uncertainty values of each sample are summarized in the ESI.† As mentioned earlier, only experimental data on solubility in pure water have been published. Taking these data into account, our  $\text{Eu}(\text{III})$  solubility point differs by 2.7% from the data published by Barabash *et al.*<sup>23</sup> and by 2.3% from the data published by Rard<sup>3</sup> – showing good consistency among the sources of solubility data in pure water. Comparing the experimental results in water with two other interpolated points (0.076 and 0.0668 mol  $\text{kg}^{-1}$ ) discussed previously by Rard<sup>3,62</sup> taking into account the work of Spedding and Jaffe<sup>63</sup> and Jackson and Rienäcker,<sup>24</sup> we obtain differences of 22.0% and 7.2%, respectively. Spedding and Jaffe<sup>63</sup> performed several measurements of the solubility of the sulfates of various lanthanides (Ce, Pr, Nd, Sm, Gd, Ho, Er, Yb, except Eu) at 25 °C. Rard<sup>60</sup> then performed an interpolation of these solubility points to estimate the solubility of  $\text{Eu}_2(\text{SO}_4)_3$  in water. Jackson and Rienäcker<sup>24</sup> reported  $\text{Eu}(\text{III})$  solubility values in water at temperatures of 20 and 40 °C. Rard<sup>3</sup> then interpolated these values to 25 °C. Such discrepancies provide evidence that these interpolations can lead to considerable overestimation, as discussed by Rard himself. Thus, these interpolation values will no longer be taken into account in our work.



**Fig. 6** (a) Solubility in the system  $\text{Eu}_2(\text{SO}_4)_3$ – $\text{MgSO}_4$ – $\text{H}_2\text{O}$  as a function of  $\text{MgSO}_4$  concentration at room temperature on a linear scale. (b) Zoom of graph (a) in the low  $\text{MgSO}_4$  concentration region. \*Interpolated points proposed by Rard.<sup>3,62</sup>



$\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  is the only solid phase that controls the  $\text{Eu}(\text{III})$  solubility in all the  $\text{MgSO}_4$  aqueous solutions investigated. The europium concentration in the studied  $\text{MgSO}_4$  concentration range (up to  $2.854 \text{ mol kg}^{-1}$ ) increases by a factor of 2 with increasing ionic strength, up to a maximum of  $0.117 \text{ mol kg}^{-1}$  (for an ionic strength of  $I \approx 4.8 \text{ eq. per kg}$ ), and then decreases again (Fig. 6). This behavior is probably due to ionic interactions and aqueous complexation of  $\text{Eu}(\text{III})$  with sulfates in concentrated aqueous  $\text{MgSO}_4$  systems. As illustrated

by the limited dispersion of the experimental data points, the measurements are very repeatable. However, it is not possible to compare our experimental data since they are the first of this type, to our knowledge.

Fig. 7 shows the experimental results of  $\text{Eu}(\text{III})$  solubility in  $\text{Na}_2\text{SO}_4$  aqueous solutions. Two well-defined regions can be distinguished: (1) a flat region that corresponds to the phase  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  prevailing at concentrations of  $m\text{Na}_2\text{SO}_4 \leq 0.01 \text{ mol kg}^{-1}$  and (2) a second region ( $m\text{Na}_2\text{SO}_4 > 0.01 \text{ mol kg}^{-1}$ ) that corresponds to the stability of  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$ . As the concentration of  $\text{Na}_2\text{SO}_4$  increases in this region, the solubility of the double salt decreases down to a minimum (at  $m\text{Na}_2\text{SO}_4 \approx 1 \text{ mol kg}^{-1}$ ) and then slightly increases beyond. No experimental data on the solubility of  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  in  $\text{Na}_2\text{SO}_4$  aqueous solutions have been published that could be used for direct comparison purposes, but experiments with other lanthanides (La, Ce, Nd, Sm, and Gd)<sup>26</sup> have been reported. The solubility results reported by Keyes and James<sup>25</sup> for the  $\text{Sm}_2(\text{SO}_4)_3$ – $\text{Na}_2\text{SO}_4$ – $\text{H}_2\text{O}$  system are also shown in Fig. 7. These authors mentioned the presence of the double salt  $\text{Na}_2\text{Sm}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  while the solubility trend of  $\text{Sm}(\text{III})$ , which is also discussed in the work of Das *et al.*,<sup>26</sup> is clearly similar to our finding. This shows that our experiments are consistent with the expected behavior of the ternary system  $\text{Eu}_2(\text{SO}_4)_3$ – $\text{Na}_2\text{SO}_4$ – $\text{H}_2\text{O}$ .

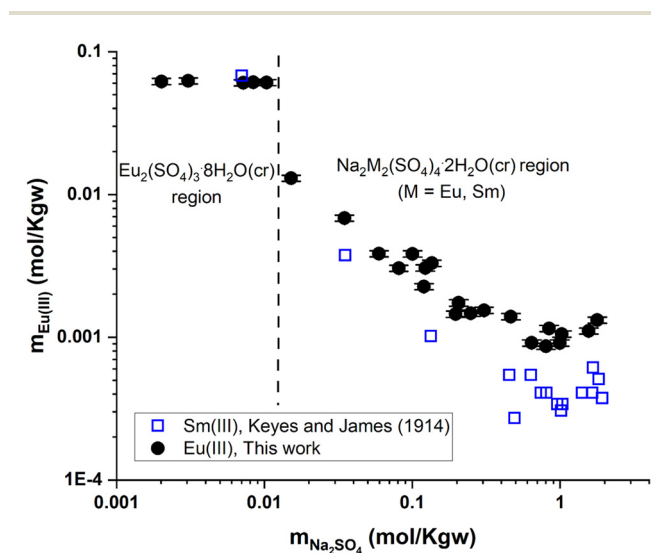


Fig. 7 Solubility in the system  $\text{Eu}_2(\text{SO}_4)_3$ – $\text{Na}_2\text{SO}_4$ – $\text{H}_2\text{O}$  (this work) and  $\text{Sm}_2(\text{SO}_4)_3$ – $\text{Na}_2\text{SO}_4$ – $\text{H}_2\text{O}$ <sup>25</sup> as a function of  $\text{Na}_2\text{SO}_4$  concentration at room temperature on a logarithmic scale.

### 5.3. Thermodynamic modelling

The solubility products at infinite dilution and the binary and ternary specific interaction parameters were optimized from the solubility data described above. The parameters determined for the Pitzer model are listed below in Table 2.

Table 2 Key parameters describing the solubility of  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}$  in  $\text{MgSO}_4$  and  $\text{Na}_2\text{SO}_4$  aqueous solutions at room temperature developed in this study

Species, i	Species, j	$\beta_{ij}^{(0)}$	$\beta_{ij}^{(1)}$	$\beta_{ij}^{(2)}$	$C_{ij}^{\phi}$	References
Pitzer binary parameters						
$\text{Eu}^{3+}$	$\text{SO}_4^{2-}$	1.775	6.877	—	–2.150	This work
$\text{La}^{3+}$	$\text{SO}_4^{2-}$	0.083	–0.202	–51.3	—	Pitzer and Silvester <sup>38</sup>
$\text{Ce}^{3+}$	$\text{SO}_4^{2-}$	0.448	6.197	4.869	0.020	Christov <sup>35</sup>
$\text{Al}^{3+}$	$\text{SO}_4^{2-}$	0.566	12.161	3.075	0.0005	Christov <sup>37</sup>
$\text{Cm}^{3+}$	$\text{SO}_4^{2-}$	1.792	15.04	—	0.600	Fanghänel and Kim <sup>64</sup>
Species, i	Species, j	Species, k	$\theta_{ik}$	$\Psi_{ijk}$		
Pitzer mixing parameters						
$\text{Eu}^{3+}$	$\text{SO}_4^{2-}$	$\text{Mg}^{2+}$	–0.182	1.347		
$\text{Eu}^{3+}$	$\text{SO}_4^{2-}$	$\text{Na}^+$	0.2223	0.3804		
Reactions					$\log_{10} K_{s,0}^{\circ}$	References
Solubility products of $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$ and $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$						
$\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr}) \leftrightarrow 2\text{Eu}_{(\text{aq})}^{3+} + 3\text{SO}_{4(\text{aq})}^{2-} + 8\text{H}_2\text{O}(\text{l})$					$-11.232 \pm 0.02^c$ $-11.911^a$	This work Das <i>et al.</i> <sup>26</sup>
$\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr}) \leftrightarrow 2\text{Na}_{(\text{aq})}^+ + 2\text{Eu}_{(\text{aq})}^{3+} + 4\text{SO}_{4(\text{aq})}^{2-} + 2\text{H}_2\text{O}(\text{l})$					$-9.510 \pm 0.100^b$ $-17.056 \pm 0.03^c$ $-17.518^a$	Jordan <i>et al.</i> <sup>2</sup> This work Das <i>et al.</i> <sup>26</sup>

<sup>a</sup> Values calculated from the Gibbs energies of formation proposed by the author together with the Gibbs energies of formation of each species from the Thermochimie database.<sup>65</sup> <sup>b</sup> Calculated from the Rard<sup>3</sup> solubility data and Davies equation<sup>66</sup> for the activity coefficient. <sup>c</sup> Uncertainty =  $2\sigma$ .



Despite the high charges of the aqueous species involved in the systems, only three binary interaction parameters ( $\beta_{ij}^{(0)}$ ,  $\beta_{ij}^{(1)}$  and  $C_{ij}^{\phi}$ ) were used to describe the binary solution properties of the  $\text{Eu}_2(\text{SO}_4)_3\text{-H}_2\text{O}$  system. Several published works used the fourth binary interaction parameter  $\beta_{ij}^{(2)}$  with 3:2 type electrolytes.<sup>34,35,38</sup> Pitzer and Silvester<sup>38</sup> studied the  $\text{La}_2(\text{SO}_4)_3\text{-H}_2\text{O}$  system, whose solubility is low (about  $0.024 \text{ mol kg}^{-1}$ ), and used enthalpy of dilution data to estimate a value for the  $\beta_{ij}^{(2)}$  (with  $i = \text{La}^{3+}$  and  $j = \text{SO}_4^{2-}$ ). Reardon<sup>34</sup> and Christov<sup>35</sup> studied the more soluble  $\text{Al}_2(\text{SO}_4)_3\text{-H}_2\text{O}$  system (up to  $1.118 \text{ mol kg}^{-1}$ ) and used osmotic coefficient data to determine specific interaction parameters, including  $\beta_{ij}^{(2)}$  (with  $i = \text{Al}^{3+}$  and  $j = \text{SO}_4^{2-}$ ). Although originally considered in our study, the use of the isopiestic experimental method to measure osmotic coefficients was finally disregarded due to the significantly low solubility observed in the two investigated  $\text{Eu(III)-SO}_4$  systems.<sup>64</sup>

As illustrated by the graphical representation in Fig. 8, the proposed model can reproduce the experimental data reliably. More specifically, the average absolute deviation

( $\text{AAD}_{\%} = \frac{\sum_{i=1}^{N_p} \text{AD}_{\%}^i}{N}$ , with  $N$  being the number of points) is less than 2% over the 19 data points measured in the present study for the  $\text{Eu}_2(\text{SO}_4)_3\text{-MgSO}_4\text{-H}_2\text{O}$  system. This value can be compared with the experimental dispersion of the experimental solubility data in pure water reported from the literature with respect to our measured value: 2.7% and 2.3% for Barabash *et al.*<sup>23</sup> and Rard,<sup>3</sup> respectively.

Regarding the  $\text{Eu}_2(\text{SO}_4)_3\text{-Na}_2\text{SO}_4\text{-H}_2\text{O}$  system, Fig. 9 also shows that the model can reasonably reproduce the experimental solubility data given their dispersion. The deviation between the model and the experimental data is up to 4.3% in the single salt region (*i.e.*,  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$ , flat region). This deviation generally lies within the reported experimental

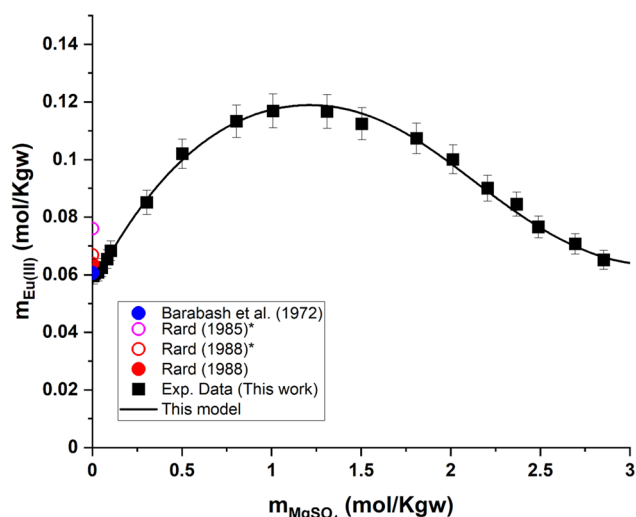


Fig. 8  $\text{Eu(III)}$  solubility in aqueous  $\text{MgSO}_4$  solution calculated using PhreeSCALE vs. experimental data at room temperature.<sup>3,23,62</sup> \*Interpolated points by Rard.<sup>3,62</sup>

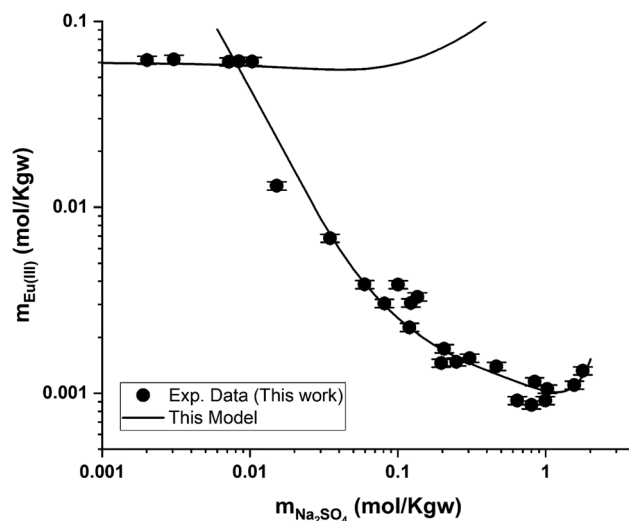


Fig. 9  $\text{Eu(III)}$  solubility in aqueous  $\text{Na}_2\text{SO}_4$  solution calculated using PhreeSCALE vs. experimental data at room temperature.

uncertainty. In the double salt region, the calculated solubility of  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$  is generally in good agreement with the experimental solubility data, although more significant deviations can be noted at intermediate  $\text{Na}_2\text{SO}_4$  concentration values ( $\leq 1.0 \text{ mol kg}^{-1}$ ).

Table 2 summarizes the solubility product values proposed in the literature for the single and double salts considered in the present study. Das *et al.*<sup>26</sup> proposed values of the Gibbs energies of formation ( $\Delta G_f$ ) for both phases –  $\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})$  and  $\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})$ . The value of  $\Delta G_f\{\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})\}$  was reported on the basis of a regression of experimental data, whereas the value of  $\Delta G_f\{\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})\}$  was estimated empirically based on the trends observed along the lanthanide series. The authors did not report the values of  $\Delta G_f$  considered for the species involved in the dissolution/precipitation reactions ( $\text{Na}^+$ ,  $\text{Eu}^{3+}$ ,  $\text{SO}_4^{2-}$  and  $\text{H}_2\text{O}$ ). The corresponding values selected in the Thermochemie database<sup>65</sup> have been considered instead, leading to  $\log_{10} K_{s,0}$  values of  $-11.911$  and  $-17.518$  for the simple salt and double salt, respectively. These values can be compared moderately well with those obtained in our work:  $-11.232$  and  $-17.056$ , respectively (Table 2).

Jordan *et al.*<sup>2</sup> proposed a calculation of  $\log_{10} K_{s,0}^{\circ}\{\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})\}$  (“for information only”, according to the author) using the Rard<sup>3</sup> solubility value ( $0.06372 \text{ mol kg}^{-1}$ ) as a basis. Since Rard<sup>3</sup> did not use any background electrolyte, Jordan *et al.*<sup>2</sup> calculated the  $\log_{10} K_{s,0}^{\circ}$  value by using the Davies equation to estimate the activity coefficients  $\gamma_{\text{Eu}^{3+}}$  and  $\gamma_{\text{SO}_4^{2-}}$  (0.061 and 0.288, respectively), considering that the ionic strength of the saturated solution ( $I_m = 0.478 \text{ eq kg}^{-1}$ ) lies within the range of validity of the Davies equation (below seawater salinity,  $\sim 0.8 \text{ eq kg}^{-1}$ ). The authors obtained a value of  $-9.51 \pm 0.1$ . Using the same calculation as proposed by the author, but using the values of  $\gamma_{\text{Eu}^{3+}}$  and  $\gamma_{\text{SO}_4^{2-}}$  from our model instead (0.010 and 0.259, respect-



ively), we obtained a  $\log_{10} K_{s,0}^{\circ}$  value of  $-11.240$ , in good agreement with our value and that of Rard,<sup>3</sup> given the associated uncertainty. This is not surprising given the good agreement between the solubility values measured by Rard<sup>3</sup> and those in the present work.

The Pitzer interaction coefficients determined in this work can be compared with previous studies in the literature investigating 3 : 2 electrolytes<sup>34,35,38</sup> and assuming the complete dissociation of ions, as considered in this work. The parameters of interactions between La(III)-SO<sub>4</sub>, Cr(III)-SO<sub>4</sub> and Al(III)-SO<sub>4</sub> are all described taking into account the  $\beta_{ij}^{(2)}$  parameters and some without  $C_{ij}^{\phi}$  – as shown in Table 2. This comparison shows that the values of the Eu(III)-SO<sub>4</sub> interaction parameters of the present work are quite strong – values presented for  $\beta_{ij}^{(0)}$  by the authors are less than 1;  $C_{ij}^{\phi}$  positive and close to 0, but with differences between the proposed values of  $\beta_{ij}^{(1)}$  and  $\beta_{ij}^{(2)}$ . These strong values can be explained by: (1) the lack of solution data to better constrain the parameter optimization and the choice for null  $\beta_{ij}^{(2)}$ ; (2) the formation of Eu(III)-SO<sub>4</sub> aqueous complexes in the investigated solution as suggested by Pitzer and Silvester<sup>38</sup> for the La<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>-H<sub>2</sub>O system. This is thoroughly explored in Part II of this work (F. dos Santos;<sup>49</sup> this journal), where spectroscopic evidence is also reported.

## 6. Conclusion

A comprehensive solubility study was conducted at  $T = (22 \pm 2)$  °C for the systems Eu<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>-MgSO<sub>4</sub>-H<sub>2</sub>O and Eu<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>-Na<sub>2</sub>SO<sub>4</sub>-H<sub>2</sub>O, covering dilute to concentrated MgSO<sub>4</sub> (0.006–2.854 mol kg<sup>-1</sup>) and Na<sub>2</sub>SO<sub>4</sub> (0.000–1.784 mol kg<sup>-1</sup>) acidic solutions. Solid phase characterization (XRD, Raman, IR, TG-DTA and quantitative chemical analysis) conducted after attaining equilibrium conditions unequivocally confirms that Eu<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>·8H<sub>2</sub>O(cr) is the only phase controlling the solubility of Eu(III) in MgSO<sub>4</sub> systems. The same binary phase controls the solubility in Na<sub>2</sub>SO<sub>4</sub> systems with salt concentrations <0.01 mol kg<sup>-1</sup>, whereas the double salt Na<sub>2</sub>Eu<sub>2</sub>(SO<sub>4</sub>)<sub>4</sub>·2H<sub>2</sub>O(cr) was identified as the only Eu(III) solid phase above this Na<sub>2</sub>SO<sub>4</sub> concentration.

Based on the experimental solubility data acquired in this work, we derived a thermodynamic model including a set of ionic interaction parameters using the Pitzer formalism, which successfully describes the solubility of Eu(III) in sodium sulfate and magnesium sulfate systems up to high ionic strength conditions (11.5 eq·kg<sup>-1</sup>). The solubility product determined for the binary salt,  $\log_{10} K_{s,0}^{\circ}\{\text{Eu}_2(\text{SO}_4)_3 \cdot 8\text{H}_2\text{O}(\text{cr})\} = -11.232 \pm 0.02$ , is in excellent agreement with previous solubility studies available in the literature. For the first time, we report an experimentally determined solubility product for the double salt with Na,  $\log_{10} K_{s,0}^{\circ}\{\text{Na}_2\text{Eu}_2(\text{SO}_4)_4 \cdot 2\text{H}_2\text{O}(\text{cr})\} = -17.056 \pm 0.03$ . This model considers the full dissociation of the Eu(III) species in MgSO<sub>4</sub> and Na<sub>2</sub>SO<sub>4</sub> aqueous solutions, as is often favored in the description of high saline systems with Pitzer equations. On the basis of additional spectroscopic evidence, Part II of this contribution

will focus on the description of Eu(III)-SO<sub>4</sub> aqueous complexes using both SIT and Pitzer activity models and a re-interpretation of the data presented here with an alternate modeling approach.

This manuscript is the first of a series providing a quantitative thermodynamic description of high saline systems with special relevance in the context of nuclear waste disposal. The ternary and quaternary systems Eu(III)-Mg(NO<sub>3</sub>)<sub>2</sub>-H<sub>2</sub>O, Eu(III)-NaNO<sub>3</sub>-H<sub>2</sub>O, Eu(III)-MgSO<sub>4</sub>-Mg(NO<sub>3</sub>)<sub>2</sub>-H<sub>2</sub>O and Eu(III)-Na<sub>2</sub>SO<sub>4</sub>-NaNO<sub>3</sub>-H<sub>2</sub>O will be tackled in the following publications.

## Conflicts of interest

There are no conflicts to declare.

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