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Thermodynamics of the Eu(μ)-Mg-SO₄-H₂O and Eu(μ)-Na-SO₄-H₂O 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 MgSO₄ and Na₂SO₄ 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 Eu₂(SO₄)₃–MgSO₄–H₂O (19 samples) and Eu₂(SO₄)₃–Na₂SO₄–H₂O (26 samples). Eu₂(SO₄)₃·8H₂O(cr) was found to control the solubility of Eu(III) in all investigated MgSO₄ solutions, as well as in dilute Na₂SO₄ systems. The transformation of Eu₂(SO₄)₃.8H₂O(cr) into the double salt Na₂Eu₂(SO₄)₄·2H₂O(cr) was observed at mNa₂SO₄ > 0.01 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 MgSO₄ and Na₂SO₄ 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 ($K_{s,0}^{(0)}$) and binary and ternary specific interaction parameters ($\beta_{II}^{(0)}$, $\beta_{II}^{(1)}$, $C_{II}^{(0)}$, θ_{IIK} , Ψ_{IIK}).

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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,^{1,2} europium is often considered as an analogue of trivalent actinides relevant in the context of nuclear waste disposal, *e.g.* Am(m) and Pu(m).^{2–4} Besides sharing the same valency, these trivalent lanthanides and actinides have similar structural properties (ionic radii and f-orbital electrons) leading to comparable

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chemical behavior, including the formation of a queous complexes of similar stability. $^{5\mathchar`-8}$

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^{-1} , mostly due to its Na, Ca, Mg, Cl and SO_4 content.^{9–11} 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^{-1} .¹² 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.¹³⁻¹⁵ 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.^{6,16-22}Only a few studies considering the solubility of Eu₂(SO₄)₃·8H₂O(cr) in the Eu₂(SO₄)₃-H₂O binary system are reported in the literature.^{3,23,24} Amongst these studies, the work published by Rard³ stands out, involving isopiestic and solubility experiments in the absence of any background electrolyte. However, data on the solubility of Eu(III) in dilute to concentrated Eu₂(SO₄)₃-Na₂SO₄-H₂O and Eu₂(SO₄)₃-MgSO₄-H₂O systems are not available so far. The study by Keyes and James²⁵ focused on the solubility of Sm(III) in the ternary system $Sm_2(SO_4)_3$ -Na₂SO₄-H₂O. The authors observed the formation of a hydrated salt in the form of $Na_2Sm_2(SO_4)_4 \cdot 2H_2O(cr)$ at Na₂SO₄ concentrations above ~0.01 M. Some studies with different lanthanides reporting the formation of double salts with Na^+ are also available in the literature, $^{25-28}$ but they were performed at low Na₂SO₄ concentrations and did not conclude any solubility information for a supposed $Na_2Ln_2(SO_4)_4 \cdot 2H_2O(cr)$ phase. The lack of relevant experimental solubility data makes it difficult to understand the behavior of Eu(III) and feed the thermodynamic models used to reproduce and estimate such properties. This fact is emphasized by the work of Das et al.,²⁶ showing that in contrast to La, Ce, Pr, Nd and Sm, the experimental data for Eu(III)-SO₄²⁻ 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 Eu₂(SO₄)₃-Na₂SO₄-H₂O and Eu₂(SO₄)₃-MgSO₄-H₂O, including the solubility of europium in dilute to concentrated aqueous solutions of sodium and magnesium sulfate (0–2.85 mol 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 Eu₂(SO₄)₃·8H₂O(cr) and Na₂Eu₂(SO₄)₄·2H₂O(cr) salts in Na₂SO₄ aqueous solutions and Eu₂(SO₄)₃·8H₂O(cr) in MgSO₄ aqueous solutions. Part II of this work is presented in a separate contribution, involving spectroscopic characterization of Eu(III) in the aqueous phase and including thermodynamic and (SIT, Pitzer) activity models specifically accounting for Eu(III)-SO₄ 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 Ω cm, 22 ± 2 °C, pore size 0.22 µm). Magnesium sulfate heptahydrate (MgSO₄·7H₂O, p.a., 99.5 wt%) and europium(III) sulfate octahydrate (Eu₂(SO₄)₃·8H₂O(cr), p.a., 99.9 wt%) were purchased from ThermoFisher Scientific. Anhydrous sodium sulfate (Na₂SO₄, p.a., >99 wt%) was obtained from Merck.

2.2. pH measurements

Proton concentration, pH_m ($pH_m = -log[H^+]$ in molal units, mol 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 pH_m values were obtained from the measured pH_{exp} values considering pH_m = pH_{exp} + A_m , where A_m is a correction factor entailing the activity coefficient of H⁺ and the liquid junction potential of the electrode at a given background electrolyte concentration. A_m values for Na₂SO₄ systems were taken as reported by Duckworth *et al.*²⁹ (work performed between 0.1 and 1.75 mol kg⁻¹ of Na₂SO₄). A_m values for the MgSO₄ aqueous systems are not available in the literature, and experimental values, pH_{exp}, are reported instead.

2.3. Solubility experiments

Solubility experiments were performed using well-characterized $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$ 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 Na_2SO_4 in the range 0.000-1.784 mol kg⁻¹ (20 samples) or of MgSO₄ in the range 0.006–2.854 mol kg⁻¹ samples). Above the Na₂SO₄ concentration of (19 0.015 mol kg⁻¹, the $Na_2Eu_2(SO_4)_4$ ·2H₂O(cr) double salt formed and was recovered for a subsequent series of experiments. The subsequent series of batch solubility experiments used the $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ double salt and was prepared with a concentration of Na₂SO₄ ranging between 0.015 and 0.122 mol 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 Eu(III) concentration. Eu(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). Aliguots of the original samples were diluted with 2% HNO3 before ICP-OES measurements. The accuracy of ICP-OES measurements was $\pm 2-5\%$.

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 Na_2SO_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 MgSO₄ were subjected to 8 washing and centrifugation steps using a solution of absolute ethanol at four dilution ratios, namely: two steps the absolute temperatur 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 β 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 N₂ 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⁻¹ 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 $50 \times (NA = 0.75)$ or $100 \times (NA = 0.9)$ objectives. Typical laser power was 20 mW (measured on the sample). The measurements as single spots were performed with 1800 L mm⁻¹ and 600 L mm⁻¹ gratings, providing spectral resolution better than 1 cm⁻¹ and 3 cm⁻¹, respectively. Due to strong Eu³⁺ lumine-scence, 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⁻¹ with 64 scans and a spectral resolution of 2 cm⁻¹. 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% HNO₃ and the solution was used for quantitative chemical analysis.

3. Thermodynamic modelling

The free energy change of a reaction, ΔG , is related to the standard free energy change, ΔG° , and the ion activity product, Q, according to the following classical equation:

$$\Delta G = \Delta G^{\circ} + RT \ln Q \tag{1}$$

where *R* is the ideal gas constant (8.314 J K⁻¹ mol⁻¹) 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:

$$Q = \prod_{i} a_{i}^{\nu_{i}} \tag{2}$$

 ν_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 γ_i are the molality (mol kg⁻¹) and the activity coefficient (unitless) of i, respectively. m_0 is a reference molality, set to 1 mol kg⁻¹, 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:

$$\Delta G = RT \, \ln \frac{Q}{K} \tag{3}$$

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):

$$SR = \frac{Q}{K_{s,0}^{\circ}}$$
(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:

$$K_{\rm s,0}^{\circ} = Q \tag{5}$$

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 $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$ at infinite dilution, $K_{s,0}^{\circ} \{Eu_2(SO_4)_3 \cdot 8H_2O(cr)\}$, is defined as follows:

$$Eu_{2}(SO_{4})_{3} \cdot 8H_{2}O(cr) \leftrightarrow 2Eu_{(aq)}^{3+} + 3SO_{4(aq)}^{2-} + 8H_{2}O_{(l)}$$
(6)

$$K_{s,0}^{\circ}\{\mathrm{Eu}_{2}(\mathrm{SO}_{4})_{3}\cdot 8\mathrm{H}_{2}\mathrm{O}(\mathrm{cr})\} = a_{\mathrm{Eu}^{3+}}^{2}a_{\mathrm{SO}_{4}^{2-}}^{3}a_{\mathrm{H}_{2}\mathrm{O}}^{8} \tag{7}$$

$$K_{\rm s,0}^{\circ}\{{\rm Eu}_{2}({\rm SO}_{4})_{3}\cdot 8{\rm H}_{2}{\rm O}({\rm cr})\} = m_{{\rm Eu}^{3+}}^{2}m_{{\rm SO}_{4}^{2-}}^{3}\gamma_{{\rm SO}_{4}^{2-}}^{2}a_{{\rm H}_{2}{\rm O}}^{8} \quad (8)$$

$$\log_{10} K_{\rm s,0}^{\circ} \{ \mathrm{Eu}_{2}(\mathrm{SO}_{4})_{3} \cdot 8\mathrm{H}_{2}\mathrm{O}(\mathrm{cr}) \}$$

= $\log_{10} \left(m_{\mathrm{Eu}^{3+}}^{2} m_{\mathrm{SO}_{4}^{2-}}^{3} \right) + \log_{10} \left(a_{\mathrm{H}_{2}\mathrm{O}}^{8} \right) + \log_{10} \left(\gamma_{\mathrm{Eu}^{3+}}^{2} \gamma_{\mathrm{SO}_{4}^{2-}}^{3} \right)$ (9)

Analogously, the solubility product of the double salt $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ at infinite dilution, $K_{s,0}^{\circ}\{Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)\}$, can be defined as follows:

$$\begin{split} \mathrm{Na_2Eu_2(SO_4)_4} \cdot \mathrm{2H_2O(cr)} \, &\leftrightarrow \, \mathrm{2Na_{(aq)}}^+ + \mathrm{2Eu_{(aq)}}^{3+} + \mathrm{4SO_{4(aq)}}^{2-} \\ &+ \mathrm{2H_2O_{(l)}} \end{split}$$

(10)

$$K_{s,0}^{\circ}\{\operatorname{Na}_{2}\operatorname{Eu}_{2}(\operatorname{SO}_{4})_{4}\cdot 2\operatorname{H}_{2}\operatorname{O}(\operatorname{cr})\} = a_{\operatorname{Na}^{+}}^{2}a_{\operatorname{Eu}^{3+}}^{2}a_{\operatorname{SO}_{4}}^{4}a_{\operatorname{H}_{2}\operatorname{O}}^{2} \qquad (11)$$

$$K_{\rm so}^{\circ} \{ Na_2 Eu_2(SO_4)_4 \cdot 2H_2O(cr) \} = m_{\rm Na^+}^2 m_{\rm Eu^{3+}}^2 m_{\rm SO_4^{2-}}^2 \gamma_{\rm Na^+}^2 \gamma_{\rm Eu^{3+}}^2 \gamma_{\rm SO_4^{2-}}^2 a_{\rm H_2O}^2$$
(12)

$$\log_{10} K_{s,0} \{ \operatorname{Na}_{2}\operatorname{Eu}_{2}(\operatorname{SO}_{4})_{4} \cdot 2\operatorname{H}_{2}\operatorname{O}(\operatorname{cr}) \}$$

= $\log_{10} \left(m_{\operatorname{Na}^{+}}^{2} m_{\operatorname{Eu}^{3+}}^{2} m_{\operatorname{SO}_{4}^{2-}}^{4} \right) + \log_{10} \left(a_{\operatorname{H}_{2}\operatorname{O}}^{2} \right)$ (13)
+ $\log_{10} \left(\gamma_{\operatorname{Na}^{+}}^{2} \gamma_{\operatorname{Eu}^{3+}}^{2} \gamma_{\operatorname{SO}_{4}^{2-}}^{4} \right)$

The activity coefficient γ_i of an aqueous species i is given by the derivative of the excess Gibbs free energy G^{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,^{30,31} 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.^{19,32,33} Hence, full dissociation has been previously considered for other 3 : 2 type electrolytes like $Al_2(SO_4)_3$,^{34,35} $Cr_2(SO_4)_3$, $Fe_2(SO_4)_3$ and $La_2(SO_4)_3$.^{36–38} In the present work, this assumption indicates that only Eu³⁺, Na⁺, Mg²⁺ and SO₄²⁻ aqueous species are considered in the chemical systems studied (in addition to H₂O, H⁺ and 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 $Eu(SO_4)^+$, $Eu(SO_4)_2^-$ and $Eu(SO_4)_3^{3-}$ aqueous complexes.^{4,39–48} 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 overparametrization of the investigated system. These aspects are further addressed in Part II of this work (F. dos Santos *et al.*;⁴⁹ this journal).

4. Parametrization procedure

To reproduce the solubility behavior of the $Eu_2(SO_4)_3$ -MgSO₄-H₂O and $Eu_2(SO_4)_3$ -Na₂SO₄-H₂O systems, the parameters to be determined are:

- The solubility products at infinite dilution, $\log_{10} K_{s,0}^{\circ} \{ \operatorname{Eu}_2(\operatorname{SO}_4)_3 \cdot 8H_2\operatorname{O}(\operatorname{cr}) \}$ and $\log_{10} K_{s,0}^{\circ} \{ \operatorname{Na}_2\operatorname{Eu}_2(\operatorname{SO}_4)_4 \cdot 2H_2\operatorname{O}(\operatorname{cr}) \};$ - The binary $(\beta_{ij}^{(0)}, \beta_{ij}^{(1)}, C_{ij}^{\phi}, \theta_{ik})$ and ternary (Ψ_{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 MgSO₄-H₂O and Na₂SO₄-H₂O, which were determined previously by Lach *et al.*⁵⁰ 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 Eu₂(SO₄)₃-MgSO₄-H₂O system, which is more constrained than the Eu₂(SO₄)₃-Na₂SO₄-H₂O 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 Eu₂(SO₄)₃-Na₂SO₄-H₂O system were determined.

Optimizations were performed by coupling the geochemical calculation code PhreeSCALE⁵⁰ with the parameter estimation software PEST⁵¹ according to the following principle.^{16,50,52} 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 MgSO₄ solutions (see Table 1 for a description of the sample conditions). In all these samples, $Eu_2(SO_4)_3 \cdot 8H_2O_3$ (cr) was found to be the only solid phase controlling the solubility of Eu(III). A Rietveld refinement was performed using the structure data of Xu et al.53 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.⁵⁴ 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 MgSO₄ aqueous solutions). All patterns match the COD 96-810-0406 for $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$. 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 Na₂SO₄ solutions $(0.00-1.56 \text{ mol kg}^{-1})$. Samples G, H and I consist entirely of Eu₂(SO₄)₃·8H₂O(cr), whereas the double-salt Na₂Eu₂(SO₄)₄·2H₂O(cr) is the only solid phase identified in samples J–N. Samples M and N show the additional presence of thenardite Na₂SO₄(vı) and a minor fraction of Na₂SO₄(uı). The latter is known as a product of the thermal treatment of the former. With increasing Na₂SO₄ concentration from 0.1 mol kg⁻¹ (sample J) to 0.35 mol kg⁻¹ (sample L), the formed Na₂Eu₂(SO₄)₄·2H₂O(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 Na₂SO₄ (samples M and N) show again smaller crystal sizes of 125 and

145 nm, respectively. As in the case of MgSO₄ systems, these results confirm the high crystallinity of the solid phases controlling the solubility of Eu(m) in Na₂SO₄ systems. It should be noted that for solid phases above 100 nm, the impact of the particle size on the $\Delta_f G^\circ$ of the solid phase (and by extension on its solubility product) is considered negligible.⁵⁵

The IR spectra of samples A-F are shown in Fig. 3. All spectra show typical features of $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$. The band assignment was made according to Denisenko et al.56-58 and Ram.⁵⁹ The internal vibrations of the SO₄ tetrahedra are observed as: (1) broad bands in the range 1050-1200 cm⁻¹ with the main peaks at 1076 and 1126 cm⁻¹ assigned to $\nu_3(SO_4)$, (2) a sharp band at 1000 cm⁻¹ with shoulders at 984 and 940 cm⁻¹ ($\nu_1(SO_4)$), and (3) small bands at 638 and 650 cm⁻¹, assigned to $\nu_4(SO_4)$. The observed frequencies match very well those reported by Buyer et al.⁶⁰ for $Y_2(SO_4)_3 \cdot 8H_2O_1$, although their assignment of the bands at 688 and 743 cm⁻¹ to $\nu_4(SO_4)$ raises some doubts. We favor the assignment of the similar bands at 683 and 745 cm⁻¹ in our spectra to H₂O vibrations as reported by Ram,⁵⁹ given the broadness of these bands. The vibrations due to H₂O are manifested by: (1) a very sharp band at 1637 cm⁻¹ (H–O–H bending, ν_2), (2) several broad bands with low intensity at 683, 745 and 803 cm⁻¹ assigned to H₂O vibrations (rocking, twisting), and (3) broad bands in the OH stretching region $(3000-3600 \text{ cm}^{-1})$ centered at 3225, 3300, 3340, and 3452 cm⁻¹. The latter confirm the presence of four distinct H₂O coordinating Eu along with four oxygens from SO₄ tetrahedra. Once again, there is a striking similarity to the frequencies of the OH stretching vibrations reported by Buyer et al.60

The trivalent Eu³⁺ 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 Na_2SO_4 and $MgSO_4$ 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, nH_2O)

		$m\mathrm{Na}_2\mathrm{SO}_4\ (\mathrm{mol}\ \mathrm{kg}^{-1})$	XRD		Raman/IR		ICP-OES		
Samples	(mol kg^{-1})		Single salt ^a	Double salt ^b	Single salt ^a	Double salt ^b	S : Eu	Na : Eu	nH_2O
A	0.006		X ^c		$\mathbf{X}^{c,d}$		1.62 ± 0.08		7.98 ± 0.03
В	0.052		\mathbf{X}^{c}		$\mathbf{X}^{c,d}$		1.60 ± 0.08		
С	0.804		\mathbf{X}^{c}		$\mathbf{X}^{c,d}$		1.66 ± 0.08		
D	1.310		\mathbf{X}^{c}		$\mathbf{X}^{c,d}$		1.60 ± 0.08		
Е	2.490		\mathbf{X}^{c}		$\mathbf{X}^{c,d}$		1.61 ± 0.08		
F	2.695		\mathbf{X}^{c}		$\mathbf{X}^{c,d}$		1.81 ± 0.08		
G		0.000	\mathbf{X}^{c}		$X^{c,e}$		1.55 ± 0.08		8.03 ± 0.07
Н		0.008	\mathbf{X}^{c}		$X^{c,e}$		1.58 ± 0.08		
I		0.010	\mathbf{X}^{c}		$X^{c,e}$		1.64 ± 0.08		
J		0.100		\mathbf{X}^{c}		$X^{c,e}$		1.10 ± 0.06	2.08 ± 0.13
K		0.122		\mathbf{X}^{c}		$\mathbf{X}^{c,e}$		1.07 ± 0.05	
L		0.035		\mathbf{X}^{c}		$\mathbf{X}^{c,e}$		1.07 ± 0.05	
М		0.643		X ^c		$X^{c,e}$		$6.59^{f} \pm 0.33$	
N		1.560		X ^c		X ^{c,e}		$7.03^f \pm 0.35$	

^a Eu₂(SO₄)₃·8H₂O(cr). ^b Na₂Eu₂(SO₄)₄·2H₂O(cr). ^c Identified phase. ^d By IR spectrometry. ^e By Raman spectrometry. ^f Presence of Na₂SO₄ solid.

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Eu₂(SO₄)₃•8H₂O (H)

NaEu(SO₄)₂•H₂O (K)

660

680

640



environment of the Eu ion in different structures,¹ it could be used for identification purposes. Fig. 4 shows the emission spectra of $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$ (sample H) and $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ (sample K) excited by the 532 nm laser recorded with 600 L mm⁻¹ grating in the range 530–680 nm. The hypersensitive ${}^{5}D_{0} \rightarrow {}^{7}F_{2}$ transition between 610 and 630 nm dominates both spectra, followed by ${}^{5}D_{0} \rightarrow {}^{7}F_{2}$ transition lines. The spectrum of Eu₂(SO₄)₃·8H₂O(cr) shows strong resemblance to that published by Xu et al.53 with the highest intensity of the 614 nm line. On the other side, the spectrum of $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ resembles the one published by Wu and Liu⁶¹ and matches perfectly that reported by Buyer et al.⁶⁰ for $Na_2Y_2(SO_4)_4 \cdot 2H_2O(cr)$ doped with Eu^{3+} at 300 K.

Fig. 5 shows the Raman spectra of both aforementioned samples taken with 1800 L mm⁻¹ grating in the range 100–1300 cm⁻¹ (530–570 nm). Once again the spectra are dominated by luminescence, but higher excited states most probably ${}^{5}D_{1} \rightarrow {}^{7}F_{1}$ and ${}^{5}D_{1} \rightarrow {}^{7}F_{2}{}^{1}$, to our knowledge, have not been reported so far. The Raman bands typical of the internal vibrations of SO₄²⁻ are visible and allow the differentiation between the Eu₂(SO₄)₃·8H₂O(cr) and Na₂Eu₂(SO₄)₄·2H₂O(cr) systems. The former shows bands at 446 and 466 cm⁻¹ (ν_2), 614 cm⁻¹ (ν_4), 1005 cm⁻¹, 1014 cm⁻¹ (ν_1) and 1080 cm⁻¹, 1142 cm⁻¹, 1180 cm⁻¹ (ν_3) in accordance with Ram.⁵⁹ In the spectrum of Na₂Eu₂(SO₄)₄·2H₂O(cr), the corresponding modes give rise to bands at 427, 489 cm⁻¹ (ν_2), 625, 666 cm⁻¹ (ν_4), 1016 cm⁻¹ (ν_1) and 1138, 1160 cm⁻¹ (ν_3). The number and positions of these bands show strong simi-

614⁰ ²D⁰

⁵D₀

592

590

⁷F₀

579 583 586

580

600

Fig. 4 Emission spectra of $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$ (sample H) and $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ (sample K) excited by a 532 nm laser. The chemi-

cal formula NaEu(SO₄)₂·H₂O stands for the Na₂Eu₂(SO₄)₄·2H₂O(cr) solid

620

Wavelength (nm)

⁷F₂

560

554 556

30k

25k

20k

Intensity 12k

10k

5k

0

phase.

540

^{'7}F2



Fig. 3 IR spectra of samples A-F (in MgSO₄ solutions)



Fig. 5 Raman spectra of $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$ (sample H) and $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ (sample K) excited by a 532 nm laser. The most intense "bands" are the emission lines from higher excited states. The chemical formula $NaEu(SO_4)_2 \cdot H_2O$ stands for the $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ solid phase.

larity to those reported by Buyer *et al.*⁶⁰ for $Na_2Y_2(SO_4)_4$ ·2H₂O (cr) doped with Eu³⁺ with maximal shifts of 2–4 cm⁻¹.

Quantitative evaluation of the TG-DTA data obtained for the initial solid of Eu₂(SO₄)₃·8H₂O(cr) indicates a content of (8.01 \pm 0.03) mol of H₂O per mol of solid. After completing the solubility experiments, the water content quantified by TG-DTA for the solid samples equilibrated in MgSO₄ aqueous solutions was (7.98 \pm 0.03) mol of H₂O per mol of solid. As for the samples G, H, I of the Na₂SO₄ aqueous system (in the region with no phase change, *i.e.* $Na_2SO_4 \leq 0.01 \text{ mol kg}^{-1}$), a water content of (8.03 ± 0.07) mol of H₂O per mol of solid was identified. This confirms that in these samples the $Eu_2(SO_4)_3 \cdot 8H_2O$ (cr) phase actually controlled equilibrium - as was already shown by XRD and Raman spectroscopy measurements. In the samples where the presence of $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ was identified by XRD and Raman spectroscopy, the measured water content was (2.08 ± 0.13) mol of H₂O per mol of solid, consistent with the formation of the double salt in the samples J to N, with concentrations of $Na_2SO_4 > 0.01$ mol kg⁻¹.

Furthermore, in the Eu₂(SO₄)₃–Na₂SO₄–H₂O system, the S : Eu (~1.5) and Na : Eu (~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 Eu₂(SO₄)₃·8H₂O(cr) (samples G to I) and Na₂Eu₂(SO₄)₄·2H₂O(cr) (samples J to N) at room temperature (Table 1). In samples M and N, with higher concentrations of Na₂SO₄, namely 0.643 and 1.560 mol 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 Na₂SO₄(vI) and Na₂SO₄(III). In the Eu₂(SO₄)₃-MgSO₄-H₂O system, the S : Eu ratio values

found confirm the presence of the $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$ 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 Eu(III) solubility at different aqueous concentrations of MgSO₄ and Na₂SO₄ 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 Eu(m) solubility point differs by 2.7% from the data published by Barabash et al.²³ and by 2.3% from the data published by Rard³ - 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 kg⁻¹) discussed previously by Rard^{3,62} taking into account the work of Spedding and Jaffe⁶³ and Jackson and Rienäcker,24 we obtain differences of 22.0% and 7.2%, respectively. Spedding and Jaffe⁶³ 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. Rard⁶⁰ then performed an interpolation of these solubility points to estimate the solubility of $Eu_2(SO_4)_3$ in water. Jackson and Rienäcker²⁴ reported Eu(III) solubility values in water at temperatures of 20 and 40 °C. Rard³ 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 $Eu_2(SO_4)_3$ –MgSO₄–H₂O as a function of MgSO₄ concentration at room temperature on a linear scale. (b) Zoom of graph (a) in the low MgSO₄ concentration region. *Interpolated points proposed by Rard.^{3,62}

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 $\operatorname{Eu}_2(\operatorname{SO}_4)_3$ ·8H₂O(cr) is the only solid phase that controls the Eu(m) solubility in all the MgSO₄ aqueous solutions investigated. The europium concentration in the studied MgSO₄ concentration range (up to 2.854 mol kg⁻¹) increases by a factor of 2 with increasing ionic strength, up to a maximum of 0.117 mol kg⁻¹ (for an ionic strength of $I \approx 4.8$ eq. per kg), and then decreases again (Fig. 6). This behavior is probably due to ionic interactions and aqueous complexation of Eu(m) with sulfates in concentrated aqueous MgSO₄ systems. As illustrated

0.1 Na2M2(SO4)42H2O(cr) region $Eu_2(SO_4)_3$ $8H_2O(cr)$ n_{Eu(III)} (mol/Kgw) (M = Eu, Sm)0.01 reaion 0.001 Sm(III), Keyes and James (1914) Eu(III), This work 1E-4 0.001 0.01 0.1 1 m_{Na₂SO₄} (mol/Kgw)

Fig. 7 Solubility in the system $Eu_2(SO_4)_3-Na_2SO_4-H_2O$ (this work) and $Sm_2(SO_4)_3-Na_2SO_4-H_2O^{25}$ as a function of Na_2SO_4 concentration at room temperature on a logarithmic scale.

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 Eu(III) solubility in Na₂SO₄ aqueous solutions. Two well-defined regions can be distinguished: (1) a flat region that corresponds to the phase $Eu_2(SO_4)_3 \cdot 8H_2O(cr)$ prevailing at concentrations of mNa₂SO₄ \leq 0.01 mol kg⁻¹ and (2) a second region (mNa₂SO₄ > 0.01 mol kg⁻¹) that corresponds to the stability of Na₂Eu₂(SO₄)₄. 2H₂O(cr). As the concentration of Na₂SO₄ increases in this region, the solubility of the double salt decreases down to a minimum (at mNa₂SO₄ \approx 1 mol kg⁻¹) and then slightly increases beyond. No experimental data on the solubility of $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ in Na_2SO_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)²⁶ have been reported. The solubility results reported by Keyes and James²⁵ for the Sm₂(SO₄)₃-Na₂SO₄-H₂O system are also shown in Fig. 7. These authors mentioned the presence of the double salt $Na_2Sm_2(SO_4)_4 \cdot 2H_2O(cr)$ while the solubility trend of Sm(III), which is also discussed in the work of Das et al.,²⁶ is clearly similar to our finding. This shows that our experiments are consistent with the expected behavior of the ternary system Eu₂(SO₄)₃-Na₂SO₄-H₂O.

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 $Eu_2(SO_4)_3 \cdot 8H_2O$ in MgSO₄ and Na₂SO₄ aqueous solutions at room temperature developed in this study

Species, i	Species, j	$eta_{ m ij}^{(0)}$	$eta_{\mathrm{ij}}^{(1)}$	$eta_{ m ij}^{(2)}$	$C^{\phi}_{ m ij}$	References
Pitzer binary pa	arameters					
Eu ³⁺	SO_4^{2-}	1.775	6.877	_	-2.150	This work
La ³⁺	SO_4^{2-}	0.083	-0.202	-51.3	_	Pitzer and Silvester ³⁸
Cr ³⁺	SO_4^{2-}	0.448	6.197	4.869	0.020	Christov ³⁵
Al^{3+}	SO_4^{2-}	0.566	12.161	3.075	0.0005	Christov ³⁷
Cm ³⁺	SO_4^{2-}	1.792	15.04	—	0.600	Fanghänel and Kim ⁶⁴
Species, i	Spo	ecies, j	Species	s, k	$ heta_{ m ik}$	$\Psi_{ m ijk}$
Pitzer mixing p	arameters					
Eu ³⁺	SO	2- 4	Mg ²⁺		-0.182	1.347
Eu ³⁺	SO	4^{2-}	Na ⁺		0.2223	0.3804
Reactions				\log_{10}	$K_{\rm s,0}^{\circ}$	References
Solubility prod	ucts of Eu ₂ (SO ₄) ₃ ·8H ₂ O	(cr) and Na ₂ Eu ₂ (SO	$_{4})_{4} \cdot 2H_{2}O(cr)$			
$Eu_2(SO_4)_3 \cdot 8H_2C$	$O(cr) \leftrightarrow 2Eu_{(aq)}^{3+} + 3SO_4$	$(aq)^{2-} + 8H_2O_{(1)}$	-11.	232 ± 0.02^{c}	This work	
				-11.	911 ^a	Das et al. ²⁶
				-9.5	10 ± 0.100^b	Jordan <i>et al</i> . ²
$Na_2Eu_2(SO_4)_4 \cdot 2$	$H_2O(cr) \leftrightarrow 2Na_{(aq)}^+ + 2H$	$Eu_{(aq)}^{3^+} + 4SO_{4(aq)}^{2^-}$	-17.	056 ± 0.03^{c}	This work	
				-17.	518 ^a	Das et al^{26}

^{*a*} 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.^{65 *b*} Calculated from the Rard³ solubility data and Davies equation⁶⁶ for the activity coefficient. ^{*c*} Uncertainty = 2σ .

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}^{ϕ} were used to describe the binary solution properties of the Eu₂(SO₄)₃-H₂O system. Several published works used the fourth binary interaction parameter $\beta_{ij}^{(2)}$ with 3:2 type electrolytes.^{34,35,38} Pitzer and Silvester³⁸ studied the La₂(SO₄)₃-H₂O system, whose solubility is low (about 0.024 mol kg⁻¹), and used enthalpy of dilution data to estimate a value for the $\beta_{ij}^{(2)}$ (with i = La³⁺ and j = SO₄²⁻). Reardon³⁴ and Christov³⁵ studied the more soluble Al₂(SO₄)₃-H₂O system (up to 1.118 mol kg⁻¹) and used osmotic coefficient data to determine specific interaction parameters, including $\beta_{ij}^{(2)}$ (with i = Al³⁺ and j = SO₄²⁻). 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 Eu(m)-SO₄ systems.⁶⁴

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

 $(AAD_{\%} = \frac{\sum_{i=1}^{N_{p}} AD_{\%}^{i}}{N}$, with *N* being the number of points) is less than 2% over the 19 data points measured in the present study for the Eu₂(SO₄)₃-MgSO₄-H₂O 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.*²³ and Rard,³ respectively.

Regarding the $Eu_2(SO_4)_3$ -Na₂SO₄-H₂O 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.*, $Eu_2(SO_4)_3$ ·8H₂O(cr), flat region). This deviation generally lies within the reported experimental



Fig. 8 Eu(III) solubility in aqueous MgSO₄ solution calculated using PhreeSCALE *vs.* experimental data at room temperature.^{3,23,62} *Interpolated points by Rard.^{3,62}



Fig. 9 Eu(III) solubility in aqueous Na_2SO_4 solution calculated using PhreeSCALE vs. experimental data at room temperature.

uncertainty. In the double salt region, the calculated solubility of $Na_2Eu_2(SO_4)_4 \cdot 2H_2O(cr)$ is generally in good agreement with the experimental solubility data, although more significant deviations can be noted at intermediate Na_2SO_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.²⁶ proposed values of the Gibbs energies of formation (ΔG_f) for both phases Eu₂(SO₄)₃·8H₂O(cr) and Na₂Eu₂(SO₄)₄·2H₂O(cr). The value of $\Delta G_{\rm f} \{ Eu_2(SO_4)_3 \cdot 8H_2O(cr) \}$ was reported on the basis of a regression of experimental data, whereas the value of $\Delta G_{\rm f}$ {Na₂Eu₂(SO₄)₄·2H₂O(cr)} was estimated empirically based on the trends observed along the lanthanide series. The authors did not report the values of $\Delta G_{\rm f}$ considered for the species involved in the dissolution/precipitation reactions $(Na^+, Eu^{3+}, SO_4^{2-} and H_2O)$. The corresponding values selected in the Thermochimie database⁶⁵ 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).

 $al.^2$ proposed Iordan et a calculation of $\log_{10} K_{s,0}^{\circ} \{ Eu_2(SO_4)_3 \cdot 8H_2O(cr) \}$ ("for information only", according to the author) using the Rard³ solubility value $(0.06372 \text{ mol kg}^{-1})$ as a basis. Since Rard³ did not use any background electrolyte, Jordan *et al.*² calculated the $\log_{10} K_{s,0}^{\circ}$ value by using the Davies equation to estimate the activity coefficients γ_{Eu}^{3+} and $\gamma_{SO_4}^{2-}$ (0.061 and 0.288, respectively), considering that the ionic strength of the saturated solution $(I_{\rm m}$ = 0.478 eq kg⁻¹) lies within the range of validity of the Davies equation (below seawater salinity, $\sim 0.8 \text{ eq} \cdot \text{kg}^{-1}$). The authors obtained a value of -9.51 ± 0.1 . Using the same calculation as proposed by the author, but using the values of γ_{Eu}^{3+} and $\gamma_{SO_4}{}^{2-}$ from our model instead (0.010 and 0.259, respectively), we obtained a $\log_{10} K_{s,0}^{\circ}$ value of -11.240, in good agreement with our value and that of Rard,³ given the associated uncertainty. This is not surprising given the good agreement between the solubility values measured by Rard³ 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^{34,35,38} and assuming the complete dissociation of ions, as considered in this work. The parameters of interactions between La(III)-SO4, Cr(III)-SO4 and Al(III)-SO4 are all described taking into account the $\beta_{ij}^{(2)}$ parameters and some without C_{ii}^{ϕ} – as shown in Table 2. This comparison shows that the values of the Eu(m)-SO₄ interaction parameters of the present work are quite strong – values presented for $\beta_{ii}^{(0)}$ by the authors are less than 1; C_{ij}^{ϕ} positive and close to 0, but with differences between the proposed values of $\beta_{ii}^{(1)}$ and $\beta_{ii}^{(2)}$. 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)-SO₄ aqueous complexes in the investigated solution as suggested by Pitzer and Silvester³⁸ for the $La_2(SO_4)_3$ -H₂O system. This is thoroughly explored in Part II of this work (F. dos Santos;49 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 Eu₂(SO₄)₃–MgSO₄–H₂O and Eu₂(SO₄)₃–Na₂SO₄–H₂O, covering dilute to concentrated MgSO₄ (0.006–2.854 mol kg⁻¹) and Na₂SO₄ (0.000–1.784 mol kg⁻¹) acidic solutions. Solid phase characterization (XRD, Raman, IR, TG-DTA and quantitative chemical analysis) conducted after attaining equilibrium conditions unequivocally confirms that Eu₂(SO₄)₃·8H₂O(cr) is the only phase controlling the solubility of Eu(m) in MgSO₄ systems. The same binary phase controls the solubility in Na₂SO₄ systems with salt concentrations <0.01 mol kg⁻¹, whereas the double salt Na₂Eu₂(SO₄)₄·2H₂O(cr) was identified as the only Eu(m) solid phase above this Na₂SO₄ 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⁻¹). The solubility product determined for the binary salt, $\log_{10} K_{s,0}^{\circ} \{ Eu_2(SO_4)_3 \cdot$ $8H_2O(cr)$ = -11.232 ± 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, $\log_{10} K_{s,0}^{\circ} \{ Na_2 Eu_2(SO_4)_4 \cdot$ $2H_2O(cr)$ = -17.056 ± 0.03. This model considers the full dissociation of the Eu(III) species in MgSO4 and Na2SO4 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(m)-SO₄ 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 $Eu(m)-Mg(NO_3)_2-H_2O$, $Eu(m)-NaNO_3-H_2O$, $Eu(m)-MgSO_4-Mg(NO_3)_2-H_2O$ and $Eu(m)-Na_2SO_4-NaNO_3-H_2O$ will be tackled in the following publications.

Conflicts of interest

There are no conflicts to declare.

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