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Introduction

- Long-term performance assessment of deep geological nuclear waste repositories
→ prediction of chemical behavior of An and long lived FP in aqueous solutions needed.
- Waste disposal in rock-salt formations in USA; option under consideration in Germany
→ high [Na⁺], [Mg²⁺] and [Cl⁻] expected in water intrusion scenarios.
- Nitrate can be found in high concentrations (≥ 1 M) as part of certain waste forms
→ waste originated from reprocessing facilities.

- Previous complexation studies with nitrate focused on acidic conditions; no MgCl₂ systems considered.

Objectives of this work

- Assessment of NO₃⁻ effect on Ln(III)/An(III) solubility under repository relevant conditions.
- Development of chemical, thermodynamic and activity models for the system Ln(III)/An(III) in NaCl–NaNO₃ and MgCl₂–Mg(NO₃)₂ solutions.

Experimental

Solubility experiments

- Batch experiments in Ar atmosphere (22 ± 2°C)
- Undersaturation approach in 0.1–5.0 M NaCl–NaNO₃ and 0.25–4.5 M MgCl₂–Mg(NO₃)₂ mixtures → up to 7 M NO₃⁻
- pH range: 7.5 ≤ pH_m ≤ 13.0

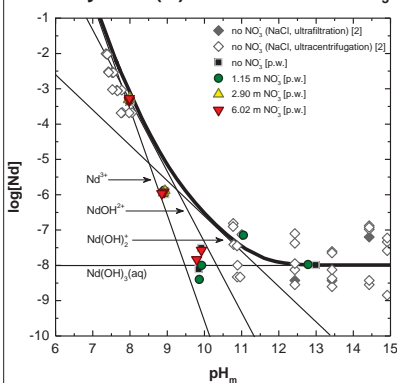
- 6–12 mg Nd(OH)₃(am) solid phase used in each experiment
- Equilibration time: t ≤ 500 days
- pH measurements: pH_m = -log m_{H+} = pH_{exp} + A_m [1]; A_m for Cl⁻–NO₃⁻ mixtures determined in this study
- [Nd(III)] measured by ICP–MS after 10 kD (2–3 nm) ultrafiltration
- Solid phase characterization: XRD, SEM–EDX

Cm(III)–TRLFS

- Sample preparation in Ar atmosphere (22 ± 2°C)
- TRLFS studies in 5.0 M NaCl–NaNO₃, 0.25 and 3.5 M MgCl₂–Mg(NO₃)₂ mixtures → up to 7 M NO₃⁻
- pH range: 1 ≤ pH_m ≤ 9
- [Cm(III)] ~ 1 × 10⁻⁷ M per sample

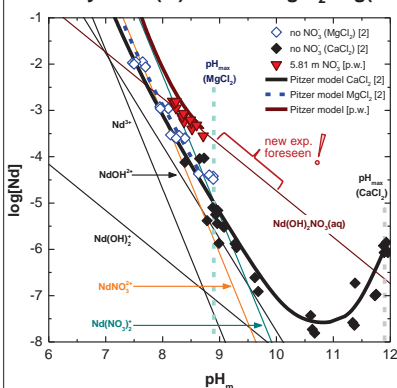
Results and discussion

Solubility of Nd(III) in 5.0 M NaCl–NaNO₃



- Very good agreement with nitrate-free solubility data reported in [2].
- No effect of NO₃⁻ on Nd(OH)₃(am) solubility in NaCl–NaNO₃ systems (even in 5 M NaNO₃).

Solubility of Nd(III) in 3.5 M MgCl₂–Mg(NO₃)₂



- Significant effect of [NO₃⁻] on Nd(OH)₃(am) solubility.
- Slope of solubility curve increases at pH_m ≥ 8.44 → change in number of OH⁻ involved in solubility reaction.
- Additional experiments in CaCl₂–Ca(NO₃)₂ (pH_{max} ~12) planned to confirm this trend.
- Handouts with experimental data at other ionic strength can be shown upon request.

Chemical and thermodynamic model for the system

Nd³⁺/Cm³⁺–H⁺–Mg²⁺–OH⁻–Cl⁻–NO₃⁻

(preliminary Pitzer model available upon request)

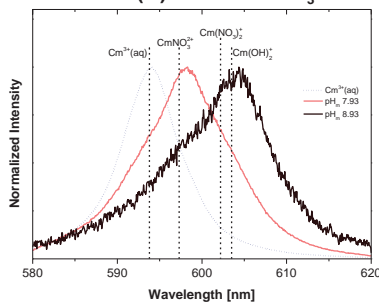
- Solid phase controlling solubility: Nd(OH)₃(am) (XRD, SEM–EDX).
- Slope –1 in presence of NO₃⁻ and pH_m ≥ 8.44 → 1 Nd(III) : 2 OH⁻ (solubility).
- Binary Cm(III)–NO₃ species relevant for pH_m ≤ 8.14 (TRLFS).
- Formation of Cm(OH)₂NO₃(aq) indicated by TRLFS at pH_m ≥ 8.44.



References

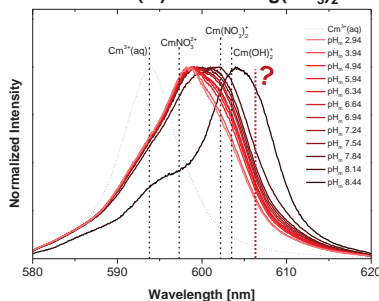
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TRLFS of Cm(III) in 5.0 M NaNO₃



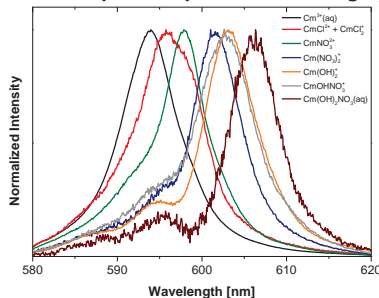
- CmNO₃²⁺ prevails in 5 M NaNO₃ and pH_m < 8.93.
- Cm(OH)₂⁺ dominates at pH_m ≥ 8.93.
- No clear evidence of relevant ternary Cm–OH–NO₃ species in 5 M NaNO₃.

TRLFS of Cm(III) in 3.5 M Mg(NO₃)₂



- CmNO₃²⁺ and Cm(NO₃)₂⁺ forming at pH_m ≤ 8.14, in good agreement with thermodynamic calculations based upon [3].
- New (ternary) species arising at pH_m ≥ 8.44.
- Three ligands complexing Cm(III) based upon red shift: 1 Cm(III) : 2 OH⁻ : 1 NO₃⁻.

Pure component spectra of 3.5 M MgCl₂–Mg(NO₃)₂



- Nitrate effect → genuine complexation reaction!
- Very complex Cm(III) speciation found in MgCl₂–Mg(NO₃)₂ mixtures → two ternary Cm–OH–NO₃ species forming.

Conclusion and outlook

- Nitrate significantly influences solubility of Nd(OH)₃(am) in concentrated and weakly alkaline MgCl₂–Mg(NO₃)₂ solutions at [Mg²⁺] ≥ 2.5 M and [NO₃⁻] ≥ 1 M.
- TRLFS data confirm that the effect of NO₃⁻ on solubility is resulting from complex formation reactions and not related to matrix effects (presence of NO₃⁻ instead of Cl⁻).
- A chemical model has been proposed including the formation of the ternary aqueous species Nd(OH)₂NO₃(aq) in equilibrium with solid Nd(OH)₃(am).
- Thermodynamic and activity models (Pitzer) for Nd³⁺/Cm³⁺–H⁺–Mg²⁺–OH⁻–Cl⁻–NO₃⁻ system are currently derived, based upon the proposed chemical model.
- Additional solubility experiments in CaCl₂–Ca(NO₃)₂ and use of advanced spectroscopic techniques (EXAFS/XANES) foreseen to confirm aqueous speciation.