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Liquid/liquid extraction kinetics of Eu(III) and Am(III) by extractants designed for the industrial reprocessing of nuclear wastes

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Abstract

Results about the kinetics of extraction of Eu(III) and Am(III) by extractants designed for the industrial reprocessing of nuclear wastes are reported. They were obtained using the rotating membrane cell technique (RMC). Extraction and stripping kinetic rate constants were determined for various compositions of the aqueous and organic phases. The transfer was studied at liquid/liquid interfaces between an aqueous nitric acid solution and an organic solvent containing the diglycolamide extractant molecule TODGA, or a mixture of the bipyridine molecule CyMe₄BTBP with TODGA (the latter being used as a phase-transfer catalyst), dissolved in an aliphatic diluent. In some experiments, an aqueous ligand (a sulfonated bis triazinyl pyridine, SO₃-Ph-BTP,

or a PyTri-diol) was added to the aqueous phase as a stripping agent. The diffusion coefficients of Eu(III) and Am(III), that are key in the analysis of the kinetic data, were measured using the RMC and the closed capillary technique. Whenever possible, mechanisms are proposed to interpret the experimental results.

1 Introduction

The reprocessing of nuclear wastes resulting from spent nuclear fuel is a worldwide topic of utmost importance in the nuclear industry and for the society itself. Various processes, generally based on liquid/liquid (L/L) extraction stages, have been proposed with the aim of reducing the volume, heat and radiotoxicity of highly radioactive waste (plutonium and americium in particular) for their disposal in a geologic repository.¹ These processes involve the separation of the most problematic radioactive elements in the wastes.

Various strategies have been developed worldwide for the reprocessing of used fuel. An overview of the main solvent extraction processes² (besides Europe) is presented in Table 1. References are indicated in the table that give more details on the policies of the countries in this domain.

Table 1: Main reprocessing strategies worldwide.

Country	Name	Extractant	Aim ^a	TRL ^b	Refs.
USA	TRUEX	CMPO + TBP	Extract U, Pu, Am, Cm	5-7	3
	TALSPEAK	D2EHPA	An/Ln separation	4-5	
China	TRPO	Cyanex 923	Extract Pu, Am, Cm	4-5	4-6
	Cyanex 301	Cyanex 301	Extract An	4-5	
Japan	SETFICS	CMPO + TBP	Extract An & Ln	3-4	4
	ARTIST	D2EHBA, TODGA	Recover U, TU ^c	2-3	
India	TRUEX	CMPO + TBP	Extract U, Pu, Am, Cm	5-7	4

^aLn=lanthanide, An=actinide; ^bTechnology Readiness Level; ^cTransuranic elements

In Europe this topic has been tackled with determination through the financing of successive European EURATOM projects since the early 90's: NEWPART, PARTNEW, EUROPART, ACSEPT, SACSESS,⁷ and now GENIORS (GEN IV Integrated Oxide fuels Re-

cycling Strategies).⁸ The European approach was centered around the use of selective extractants and molecular diluents that would generate a minimal amount of secondary waste. A feature of this strategy is the use of chemicals that only comprise the C, H, O, and N atoms (often referred to as the CHON principle⁹), which makes them suitable for subsequent incineration.

Reference aqueous separation process routes have emerged from these in-depth studies. They are depicted in Figure 1.^{10,11}

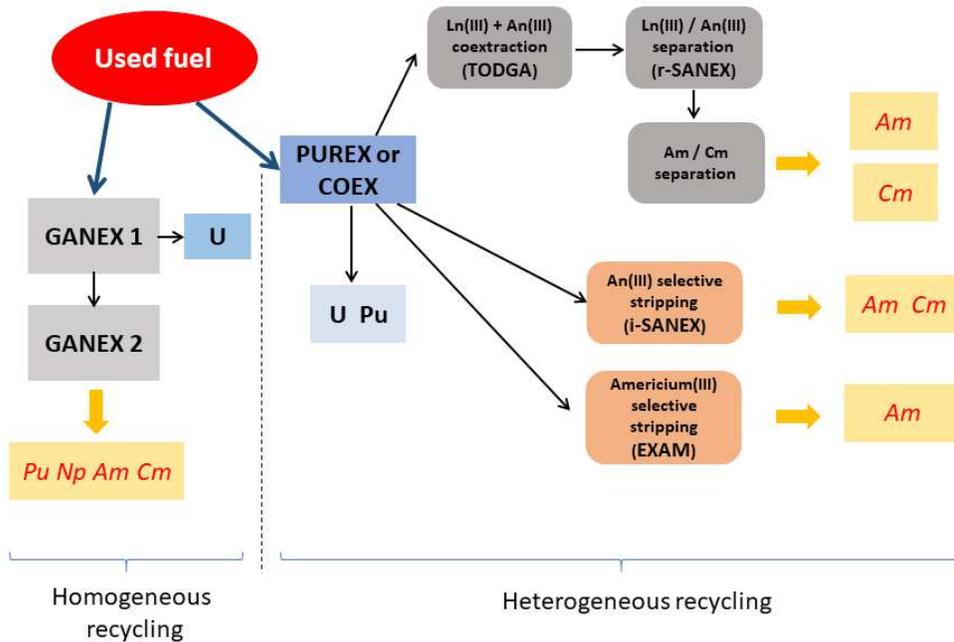


Figure 1: Main routes of the European partitioning process strategy envisaged for the recycling of actinides (An) from used fuel (Ln = lanthanides). EXAM = Extraction of americium.

One route uses the GANEX (Grouped Actinide EXtraction) process^{12,13} in which uranium is separated from the waste in a first step, and then transuranic actinide elements are isolated (Np, Pu, Am, and Cm) from all fission products.

In the other route, the PUREX (Plutonium, Uranium, Reduction, EXtraction) process,¹⁴ first implemented in the Manhattan project, is employed for the separation of uranium and

plutonium from other fission products by using tributyl phosphate (TBP) as the extractant. The COEX process is a modified version of PUREX. Then the DIAMEX (DIAMide EXtraction) process developed at CEA (Commissariat à l’Energie Atomique) in France may be used. It consists of the co-extraction of trivalent minor actinides (MA’s, mainly composed of americium(III) and curium(III)) and lanthanides (Ln’s) from a PUREX raffinate by employing a malondiamide extractant. Although they constitute less than 0.1% of the initial spent fuel mass, the minor actinides (especially neptunium, americium, and curium) will be the main contributors to the radiotoxicity (and heat generation) after a three century storage of high-level radioactive liquid waste (obtained after the PUREX stage). In the mixture so obtained, MA’s and Ln’s may be separated by using the r-SANEX (Regular Separation of ActiNides by EXtraction) process.¹⁵ The separation of Ln’s from MA’s is required for the subsequent transmutation of MA’s into short-lived or stable nuclides (which is the aim of the so-called “Partition and Transmutation” strategies¹⁶).

It has been proposed to replace the malondiamide molecule used in the DIAMEX and SANEX processes by a diglycolamide, which results from the insertion of an ether group between the two amide groups of the malondiamide. The reference molecule is the N,N,N’,N’-Tetra-n-Octyl-DiGlycolAmide (TODGA) that was first synthesized by a group from the Japan atomic Energy Agency.¹⁷

Variations of the original SANEX step have been proposed in order to reduce the number of stages following the PUREX process, namely the innovative SANEX (i-SANEX)¹⁸ and the 1-cycle SANEX processes.^{10,19} In i-SANEX it is expected to employ a solvent phase comprising 0.2 M TODGA with 5 vol.% 1-octanol (corresponding to a concentration of 0.32 M) in an inert diluent. The purpose of adding 1-octanol is to prevent the onset of a third phase.^{20,21} In 1-cycle SANEX the solvent comprises a mixture of CyMe₄BTBP and 5 mM TODGA in an aliphatic diluent,¹⁰ in which CyMe₄BTBP designates the 6,6’-bis(5,5,8,8-tetramethyl-5,6,7,8-tetrahydrobenzo[1,2,4]-triazin-3-yl)-[2,2’]-bipyridine molecule.

Modeling and simulation are essential tools to design flowsheets efficiently and to test

the flowsheet against maloperation scenarios, or process upsets. An experimental study of maloperations is not feasible in practice because it would require a huge programme of active pilot plant trials that would be long, hazardous, and very costly. Kinetic data are essential for an accurate simulation of a process flowsheet, but interfacial mass transfer is still one of the most poorly modeled aspects of solvent extraction. This is especially true when considering the case of short residence time centrifugal contactors (with aqueous/organic contact times of a few seconds typically), which is the most likely scenario for future MA separation plants.

In the literature, kinetic data are extremely scarce concerning the transfer of trivalent lanthanide and actinide ions from nitric acid solutions (as commonly used in nuclear re-processing) by the reference extractants TODGA and CyMe₄-BTBP in molecular solvents. To our best knowledge, two papers have been published in the case of TODGA, about the extraction of Am(III)²⁴ (using a Lewis cell) and that of Ce(III)²⁵ (using a rotating diffusion cell). Besides, one finds a publication in which TODGA was dissolved in an ionic liquid for the extraction of Eu(III).²⁶ No reference was found in the case of CyMe₄-BTBP.

The research presented in this communication was done within the European project GENIORS (2017-2021). For the development of separation processes, the extraction and stripping kinetics of the lanthanide Eu(III) ion and the actinide Am(III) ion were investigated at the interface between nitric acid solutions and organic phases comprising TODGA, or a mixture of CyMe₄-BTBP and TODGA, in the diluent TPH (‘tétrapropylène hydrogéné’, which is a mixture of isomers of dodecane and a widespread diluent in the nuclear industry). The rotating membrane cell (RMC) technique²⁷ was employed for the study of the kinetics. The aqueous phase sometimes contained an hydrophilic aqueous ligand, SO₃-Ph-BTP (a sulfonated bis triazinyl pyridine)²⁸ or PTD (2,6-bis[1-(propan-1-ol)-1,2,3-triazol-4-yl]pyridine),^{29,30} to mimic the conditions of a SANEX process. The PTD molecule is more interesting than SO₃-Ph-BTP because, contrary to the latter, it satisfies the CHON principle.

The first aim of this work is to provide kinetic data for these important systems. The results are analyzed from a fundamental point of view, and the practical consequences for in-

dustrial processes are examined. An additional underlying aim of this study is to interrogate the capability of the technique to provide reliable kinetic data.

This work is organized as follows. The RMC technique is described in the next section. Then the experimental features of this work are exposed, and the results for the transport coefficients (viscosities of the solutions and diffusion coefficients of the solutes) are given. After that, kinetic experiments are reported in conditions of extraction and of stripping independently on the same system. Expectedly, the kinetic rate constants obtained in the two cases should have similar values. These experiments provided a test of the technique and an estimation of the experimental uncertainty with the RMC. After this preliminary study, the extraction and stripping kinetics of Eu(III) and Am(III) are investigated in various conditions. The effect of varying the concentration of the extractants, of the aqueous ligands, of nitric acid, and of temperature, is examined. This work ends with a conclusion summarizing the results and exposing some prospects for the future.

2 Experimental

2.1 Overview of experimental techniques

The RMC possesses the advantage of controlling the hydrodynamics in the two phases. Few techniques possess this property.³¹ The first such technique was devised by a Russian group.³² It used a strip of chromatographic paper to stabilize the aqueous phase, that was immersed into the organic phase. Then Albery et al. proposed the rotating diffusion cell (RDC).³³

The control of transport in the two phases is a fundamental requirement without which no reliable result can be obtained. In this respect, the Lewis cell should not be used, except in the case of very slow kinetics. Otherwise it may yield kinetic rate constants that are in error by several orders of magnitude.³⁴ It was recognized by Danesi (who used this cell extensively) that the plateau region was not necessarily an indication of a kinetic regime

because of a “slip effect” on the propellers when the rotation speed was increased beyond some value.³⁵

Microfluidic devices that ensure direct contact of the phases have been proposed in the past two decades.³⁶ However, it is not easy to reliably control the flow and the transport of the species in cells in which the interface is free. It still seems difficult to estimate the contribution from diffusional transport in these cells without making some approximations, in the case of parallel flows³⁷ (in which the geometries are not yet simple enough to allow an estimation of the diffusional contribution) as well as moving droplets.^{38,39} As a consequence, these studies generally deliver values for *global* (apparent) kinetic rate constants combining the effect of the interfacial reaction and diffusional transport in the layers adjacent to the interface.

2.2 The RMC technique

2.2.1 Description of the RMC

The cell consists of a thin membrane that is glued on the base of a cylinder made of perspex (see Figure 2).

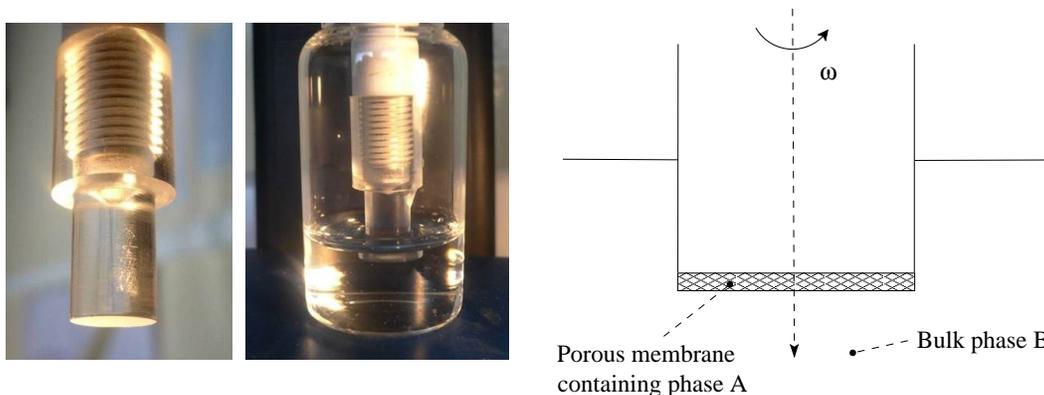


Figure 2: The RMC technique. Left: View of the cell with the membrane glued at the bottom. Center: Cell rotating in the outer phase. Right: Sketch of the technique.

Two types of membranes, one hydrophilic and one hydrophobic, were used in this work to contain the aqueous and organic phases, respectively (see section 2.7). Besides their

chemical compatibility with the phases, these membranes were selected because they were found to give consistent results for the diffusion coefficients of Eu(III) measured with the closed capillary technique and the RMC (see Supplementary Information).

The phase contained in the membrane, denoted by A, was spiked with the radioactive tracer to be extracted. The cell was mounted on a rotating-electrode spindle that can be rotated at a definite speed. Initially, it was set into rotation at a known speed and it was immersed into the outer phase B. The rotating electrode was set on a rack that could be lowered or raised easily.

More practical details about the technique are given in the Supporting Information addendum.

2.2.2 Modeling

It was assumed that the transfer of solute occurs strictly at the interface between the two phases. This assumption was made because of the very low solubility of extractant in aqueous phases.⁴⁰ This contrasts with a model that was first proposed a few decades ago by the renowned Carl Hanson^{41,42} and later extended by Rod,⁴³ in which the complexation reaction may occur in a thin layer in the aqueous phase, adjacent to the interface. This model was not used in the present work.

With the assumption that the complexation reaction is strictly interfacial, the ratio of matter extracted as a function of time can be written as,²⁷

$$P(t) = 1 - \exp(-t/\tau) \quad \text{or} \quad -\ln[1 - P(t)] = t/\tau \quad (1)$$

where τ is the mean-passage time of the solute in the overall A-to-B transfer process. It is given by,

$$\tau = \tau_A + \tau_i + \tau_B \quad (2)$$

in which τ_A is the mean diffusion time in the membrane (phase A), τ_i is the characteristic

time for the interfacial reaction (from A to B) and τ_B is the mean residence time of solute in the diffusion layer, that results from the competition between the back transfer from B to A at the interface and removal from the diffusion layer by dilution in B.

The expressions of these times are,

$$\tau_A = L^2\theta/(3D_A) \quad \tau_i = L/k_{A \rightarrow B} \quad \tau_B = \sigma L \delta_B/(K_{B/A}D_B) \quad (3)$$

in which L is the membrane thickness, θ is its tortuosity, D_A and D_B are the solute diffusion coefficients in the bulk of phases A and B, respectively, σ is the membrane porosity, $K_{B/A} = C_B^{eq}/C_A^{eq}$, and δ_B is the diffusion layer thickness in phase B given by the Levich equation,⁴⁴

$$\delta_B = \frac{1.612}{Sc^{1/3}} \sqrt{\frac{\nu_B}{\omega}} \quad (4)$$

with Sc the Schmidt number ($Sc = \nu_B/D_B$, which is commonly of the order of 10^3), ν_B the kinematic viscosity of phase B, and ω the rotation speed of the membrane (in rad s^{-1} , $\omega = 2\pi N/60$ where N is the rotation speed in rpm).

In practice, the value of the diffusion time in the membrane, τ_A , is of the order of a few seconds (typically 5-10 s) when the membrane contains the aqueous phase. It is much larger in the case of an organic phase (50-200 s) because then the diffusion coefficient of the solute is much smaller.

In these relations $k_{A \rightarrow B}$ and $k_{B \rightarrow A}$ are the forward (from A to B) and reverse (B to A) kinetic rate constants, respectively, and $K_{B/A}$ satisfies the relation,³¹

$$K_{B/A} = k_{A \rightarrow B}/k_{B \rightarrow A} \quad (5)$$

When A is aqueous and B is organic, then $K_{B/A}$ is the usual organic-to-aqueous distribution ratio, K . In that case (extraction), $k_{A \rightarrow B}$ is the extraction rate constant, k_{ex} , and $k_{B \rightarrow A}$ is the stripping (back-extraction) rate constant, k_{st} . In the reverse case of stripping (A=org,

B=aq) then $K_{B/A} = 1/K$, and $k_{A \rightarrow B} = k_{st}$, $k_{B \rightarrow A} = k_{ex}$.

By virtue of eq 5, the time τ_B in eq 3 may be alternatively expressed as,

$$\tau_B = \frac{\sigma k_{B \rightarrow A}}{D_B/\delta_B} \tau_i \quad (6)$$

which expression highlights the competition between back-transfer of solute into the membrane at a rate $\sigma k_{B \rightarrow A}$ and its removal into B by convective diffusion at a rate D_B/δ_B .

In the case of infinitely fast interfacial kinetics ($k_{A \rightarrow B}$ and $k_{B \rightarrow A} \rightarrow \infty$ with $K_{B/A}$ unchanged), the process becomes diffusion controlled and eq 2 reduces to

$$\tau_\infty = \tau_A + \tau_B \quad (7)$$

Replacing τ by τ_∞ in eq 1 yields the diffusive limit, P_∞ , of the process.

According to eq 1 a fit of the experimental results for $-\ln[1 - P(t)]$ vs. t yields the value of the time τ . Then, the latter may lead to the determination of $k_{A \rightarrow B}$ value (using eqs 2-5), which is the only unknown ($k_{B \rightarrow A}$ being given by eq 5), provided all bulk transport parameters (diffusion coefficients, viscosities,...) and the characteristics of the membranes have been determined experimentally.

This technique has been shown to give constant values for the rate constant of transfer of acetic acid when the rotation speed was varied.⁴⁵ This result showed that a good control of the hydrodynamics is achieved with this technique, which avoids the need of prior calibration.

2.2.3 Use of the RMC in this study

From a practical point of view, the stripping configuration offers the silver lining of using very small amounts of extractant, since in this case the organic solution is placed in the membrane, which is of very small volume (a few μL). However, the question arises as to whether the extraction configuration might be preferable when the distribution ratio K is large, which would make the time τ_B very small in eq 3 and therefore provide more accuracy

in the determination of the kinetic rate constants. This point is illustrated in section 3 below.

By and large, the strategy adopted in this work was to mainly perform stripping experiments and, whenever possible, to carry out a few extraction experiments in order to confirm the result.

2.3 Chemicals and methods

The chemical structures of the molecules used in this study are shown in Figure 3.

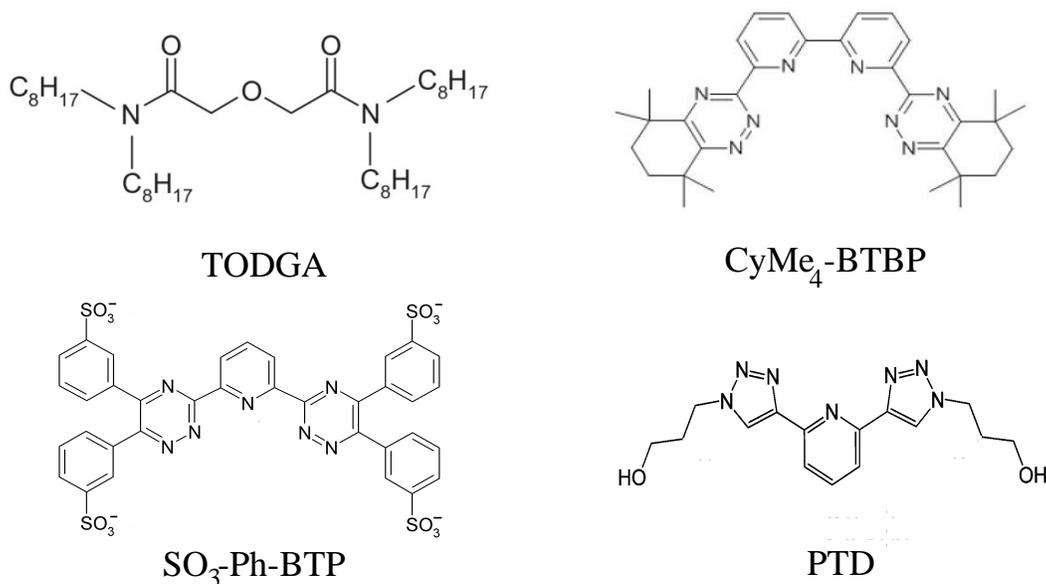


Figure 3: Chemical structures of the molecules used in this study (SO₃-Ph-BTP in tetravalent ionic form, counterion: Na⁺).

The extractants TODGA and PTD, were synthesized by authors of this work (at Twente and Parma, respectively).^{29,46} The extractant CyMe₄-BTBP and the complexing agent SO₃-Ph-BTP were synthesized by Karlsruhe Institute for Technology (KIT). The SO₃-Ph-BTP was provided and used as the tetra-sodium salt. The TPH diluent was provided by CEA (purchased from NOVASEP). These compounds were used as received. The ¹⁵²Eu(III) and

$^{241}\text{Am}(\text{III})$ radioactive sources were purchased from ORANO-LEA (France). Aqueous solutions were prepared with ultrapure water (Millipore, 18.2 M Ω cm). All other chemicals were purchased from AnalaR (Normapur), Sigma-Aldrich and Fluka, and were used without further purification.

A small amount of 1-octanol (5 vol.%, which corresponds to 0.32 M) was added to the TPH diluent in order to prevent the formation of a third phase. Hereafter, for convenience, this solvent will be designated by the abbreviation TPH-O.

The organic phases were pre-equilibrated with the aqueous phase prior to a kinetic experiment, in order to saturate the organic phase with water and nitric acid. The aqueous phases were not pre-equilibrated with the organic phase (the equilibrium concentration of 1-octanol in the aqueous phases was neglected). In contrast, in the kinetic experiments with 1-octanol as the diluent, the two phases were pre-equilibrated with each other because 1-octanol is slightly soluble in the aqueous phase ($\sim 4 \times 10^{-3}$ M in water at 25°C).

The distribution ratios were determined by equilibrating 500 μL of the two phases in a 5 mL vial made of Teflon. The phases were highly stirred during 1 to 4 hours. After centrifugation, aliquots of the two phases (350 μL) were separated and their activities were measured using a gamma counter, Packard Cobra II Auto Gamma (calibrated weekly, according to the standard procedure using a ^{137}Cs calibration source of 9435 Bq).

In this work, the RMC was rotated at a speed of 600 rpm, and the diffusion and extraction experiments were carried out at a temperature of $22 \pm 1^\circ\text{C}$, unless otherwise indicated.

2.4 Solute diffusion coefficients

The bulk transport parameters and the characteristics of the membranes are key parameters that must be determined experimentally.

No data for the diffusion coefficient of $\text{Eu}(\text{III})$ and $\text{Am}(\text{III})$ were found in the literature in the case of our systems. The diffusion coefficients of $\text{Eu}(\text{III})$ were determined using the closed capillary technique suitable for a γ -emitter,^{47,48} and the RMC. Typical results are shown in

Figures S1 and S2. The diffusion coefficients of $^{241}\text{Am(III)}$ could not be determined with the capillary technique because it is a γ -emitter of low energy, which would impose the use of a too large amount of tracer for a reliable measurement to be done.

In the case of the RMC, identical solutions A and B (see Figure 2) were employed, except for the diffusing solute which was present in A only. The porosity and the tortuosity of a membrane are required for the determination of the D 's. They were assessed as described below in section 2.7.

The measurement of a diffusion coefficient with the RMC typically lasts 10 seconds for an aqueous solution, and 1 minute for an organic solution, as compared to several days with the closed capillary technique. Nevertheless, the RMC requires a larger amount of solution (of the order of 2 or 3 mL) than the capillary technique (less than 100 μL). Similar results for the D values were obtained with the two techniques in the case of Eu(III) (see Supporting Information).

The Stokes hydrodynamic radii values, R , of Eu(III) and Am(III) ions may be derived from the D values by using the Stokes-Einstein equation in the form,

$$R = k_B T / (6\pi\eta D) \quad (8)$$

with k_B the Boltzmann constant, T the temperature, and η the dynamic viscosity of water or of the organic diluent (the medium in the vicinity of the complex) at temperature T . The hydrodynamic radius R provides an estimation of the size of the complex in solution.

The results for the diffusion coefficients are gathered in the Supporting Information.

2.5 Diffusion coefficients measured using NMR

Initially, it was planned to measure the self-diffusion coefficients of (inactive) Eu(III) by using NMR (nuclear magnetic resonance).⁴⁹ However, this was not possible because the europium nucleus is paramagnetic. It was not possible to make this measurement for the

diamagnetic lanthanum nucleus either, because its resonance frequency is very close to that of deuterium (used for the lock in our NMR spectrometer). Then the lock filter blocked the signal produced by the lanthanum nucleus.

The diffusion coefficients of the solvent species in solutions of TODGA + 1-octanol in TPH were investigated by using NMR. Most experiments were carried out in presence of lanthanide (Ln) ions, Eu(III) and La(III), at a fixed concentration of 0.02 M (in the form of Ln-nitrate). La(III) was used in addition to Eu(III) in order to confirm the results obtained with the latter. For comparison, some experiments were conducted in the absence of Ln ions, at 0.06 M and 0.2 M TODGA. All solutions were pre-equilibrated with a 0.5 M HNO₃ solution (prior to the incorporation of Ln(III)).

The technical details of these experiments and the results are presented in the Supporting Information.

2.6 Solution viscosities

The values of the kinematic viscosities of the solutions, ν , are required in the analysis of the kinetic data, and those of the dynamic viscosity, η , are needed for the calculation of the Stokes radii.

The values of η at 22°C were measured using an Anton Paar AMVn Automated Micro Viscometer. The data at 22°C, and those estimated at 35°C, are collected in the Supporting Information.

2.7 Membranes

The physical characteristics of the membranes, thickness L , porosity σ , and tortuosity θ , are needed in the treatment of the experimental data.

Two types of membranes were purchased from Merck Millipore: the hydrophilic Omnipore PTFE membrane (JHWP04700, manufacturer's data: pore size 0.45 μm , porosity of 80%) was employed to contain aqueous solutions, and the hydrophobic Durapore PVDF

membrane (HVHP04700, manufacturer’s data: pore size 0.45 μm , porosity of 75%) was employed for organic solutions. The porosity values were also measured by impregnating membranes (glued on a plastic cylinder) with TPH and by measuring the corresponding mass of diluent. The membrane thicknesses, L , were measured by using a digital micrometer.

The tortuosities of the two types of membrane were obtained as follows. First the diffusion coefficient of Eu(III) ion was measured by using the closed capillary technique (with a capillary of 3 cm total length), in the case of a 3 M HNO_3 aqueous solution for the hydrophilic membrane, and in the case of a 0.2 M TODGA organic solution in TPH-O solvent pre-equilibrated with a 3 M HNO_3 solution. In a second step diffusion experiments were carried out with the RMC for the same systems, with identical solutions A and B ($A \equiv B$, aqueous or organic). The tortuosity value of a membrane was then adjusted so as to recover the experimental data obtained with the RMC, and by using the D value for Eu(III) measured using the closed capillary technique. The adjustment was performed by employing the formula for P_∞ , eq 1 together with eq 7 and taking $K_{B/A} = 1$.

This methodology relies on the consideration that the pores of the membranes are of macroscopic size ($\sim 0.45 \mu\text{m}$ according to the manufacturer). This feature allows one to consider the diffusion process of an ion in the pores as being of the same nature as diffusion in the bulk solution (no effect of confinement).

The values of L , σ , and θ are collected in Table 2. These values were used for the subsequent determination of the diffusion coefficients of the solutes in other solutions and of the kinetic rate constants with the RMC technique.

Table 2: Physical characteristics of the membranes.

Type of membrane	Membrane thickness $L / \mu\text{m}$	Porosity σ	Tortuosity θ
Hydrophilic JHWP04700	58	0.80	2.51
Hydrophobic HVHP04700	102	0.75	1.94

The membranes were glued on the base of the plastic cylinder with liquid Kapton (a

polyimide purchased from Aldrich: Pyre-ML RC-5019, CAS no. 25038-81-7). Moreover, the lower part of the plastic cylinder was covered with Kapton in order to isolate its surface from the solutions (especially organic). Kapton was chosen because it is known to have very weak interactions with most compounds.

3 Kinetic rate constants from extraction and stripping experiments

As mentioned in section 2.2.3, the kinetic rate constants were determined by carrying out preferentially stripping experiments. Additionally, a few extraction experiments were performed to confirm the results, when the extractant was available in sufficient amount (a kinetic extraction experiment consumes about 500 times more extractant than a stripping experiment).

So the question arises of whether the two configurations yield consistent values for the kinetic rate constants. In this section, this problem is illustrated in the case of a system exhibiting a high value of the distribution ratio K . This system was comprised of 0.2 M TODGA in TPH-O solvent (pre-equilibrated with a 0.5 M HNO_3 solution) as the organic phase, and a 0.5 M HNO_3 solution as the aqueous. For this system one has $K = 329$. This high K value makes this system an extreme case to study.

Figure 4 and Table 3 show the results obtained for the rate constants.

Table 3: Characteristic times, contrast parameter, and extraction and stripping kinetic rate constants obtained from extraction and stripping experiments, for the solutions of Figure 4.

	$K_{B/A}$	τ_A (/s)	τ_B (/s)	τ_i (/s)	C	k_{ex} (cm s ⁻¹)	k_{st} (cm s ⁻¹)
Extraction	329	5.27	0.0135	2.54	0.32	2.28×10^{-3}	6.95×10^{-6}
Stripping	0.00304	54.3	768	1617	0.66	2.07×10^{-3}	6.31×10^{-6}

Figure 4 shows that extraction (left) exhibits much faster kinetics than the stripping experiment (right). The minimum duration of an extraction experiment was ~ 5 s.

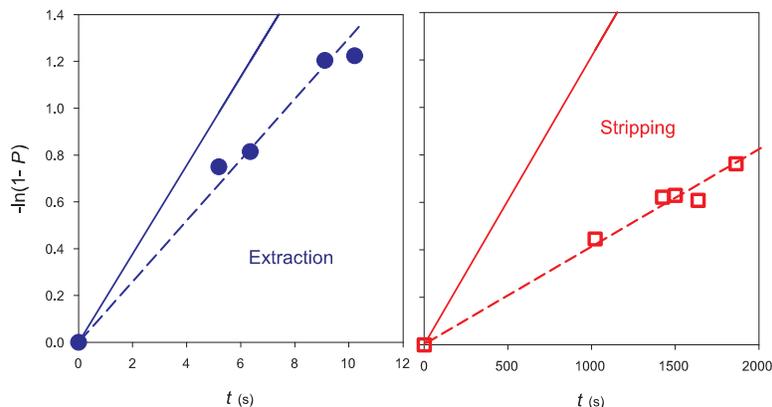


Figure 4: Proportion of extracted $^{152}\text{Eu}(\text{III})$, P , as a function of time in the case of extraction (left) and stripping (right) experiments. Aqueous phase = 0.5 M HNO_3 . Organic phase = 0.2 M TODGA in TPH-O. Dashed lines = fit of experimental data using eq 1; solid lines = diffusive limit (eqs 1 and 7).

The characteristic times of transfer, τ , had values of ~ 7.8 s and 2440 s in the extraction and stripping cases, respectively. The difference is due to the much higher values of τ_A , τ_B , and τ_i in the stripping case (see Table 3). These high values originate from the lower diffusion coefficient in the organic phase, and from the low values of $K_{aq/org}$, and k_{st} , respectively.

The experimental points are closer to the diffusive limit (DL) in the extraction configuration. The distance between the two reflects the effect of the interfacial reaction which slows down the transfer of the ion across the interface. The larger relative distance in the case of stripping entails that the accuracy of the rate constant determination should be higher than in the extraction experiment.

This distance may be quantified by defining a contrast parameter as the relative difference between the slopes, $C = -(\tau^{-1} - \tau_\infty^{-1})/\tau_\infty^{-1}$, which by virtue of eqs 2 and 7 may be rewritten as

$$C = \frac{\tau_i}{\tau} \quad (9)$$

The values of C are given in Table 3 in the present case.

It can be seen in Table 3 that the rate constants k_{ex} and k_{st} obtained in the two exper-

imental configurations are comparable, with a discrepancy of $\sim 10\%$. In other experiments of the same type, with other systems, the discrepancy could sometimes reach $\sim 20\%$. These values give an order of magnitude of the experimental uncertainty.

This may seem a rather large uncertainty. However, the extraction and stripping experiments were carried out using two different types of membrane. Moreover, experimental kinetic rate constants may be in error by several orders of magnitude if the hydrodynamics and the diffusive transport are not well controlled. Therefore an uncertainty of 10% or 20% is actually small in comparison of what it can be when an unsuitable technique is used. It also reflects the intrinsic difficulty of determining L/L interfacial kinetic rate constants.

In view of this uncertainty value, the results for the rate constant will hereafter be given with two significant figures only.

4 Kinetics with TODGA in TPH-O

4.1 Effect of TODGA concentration

The influence of the TODGA concentration on the transfer kinetics of $^{152}\text{Eu(III)}$ was investigated. Figure 5 shows the results for the distribution ratio K and the kinetic rate constants. The numerical values are given in the Supporting Information.

One notes in Figure 5 that, when the TODGA concentration was increased from 0.1 M to 0.4 M, the distribution ratio K was highly enhanced, showing a strong complexation by TODGA. Since the distribution ratio is high for every TODGA concentration, *extraction* experiments were conducted exceptionally in this case, because stripping experiments would have been accompanied by a larger uncertainty in the values of the rate constants (due to a high influence of the transport process in the outer aqueous phase).

The stripping rate constant (k_{st}) is rather small (of the order of a few 10^{-6} or 10^{-5} cm s^{-1}). It was considerably reduced by an increase in the concentration of TODGA. At the same time, the extraction rate (k_{ex}) is rather high (a few 10^{-3} cm s^{-1}) and it dropped

notably with [TODGA]. Since the extraction rate of Am(III) is equally fast (see Table 4), this implies that the co-extraction stage of Eu(III) and Am(III) by TODGA (top of Figure 1) should be fast in the industrial process.

A log-log plot of K (see Figure S5), k_{ex} , and k_{st} (not shown for the rate constants) indicates approximately linear variations with slopes of ~ 1.5 , -0.4 , and -1.9 , respectively. This dependency suggests a number of TODGA molecules between 1 and 2 per Eu(III) ion. This result is lower than slopes of ~ 3.2 and ~ 3.7 found previously in the literature in the case of 1 M HNO_3 and lower TODGA concentration in *n*-dodecane (ranging from 0.01 M to 0.1 M) without 1-octanol.^{50,51} This discrepancy may be due to the different experimental conditions (nitric acid and TODGA concentrations, use of TPH instead of dodecane, and presence of 1-octanol in our measurements), and to the fact that the slope of the log-log plot of K in Figure S5 is expected to be larger for TODGA concentrations below 0.1 M in our system.

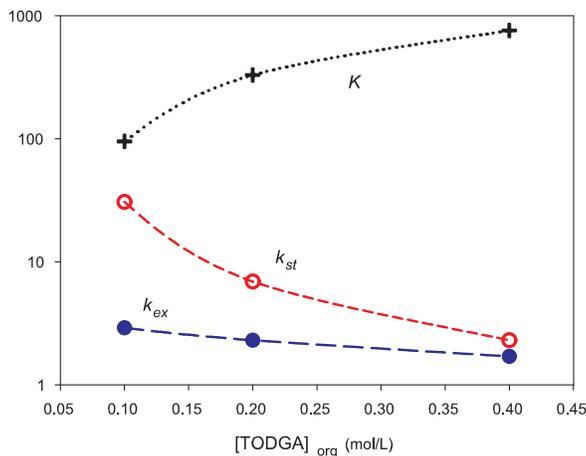


Figure 5: Results for $^{152}\text{Eu(III)}$ as a function of [TODGA] in organic phase (log plot on vertical axis): (+) = Distribution ratio K , (\bullet) = k_{ex} in units of $10^{-3} \text{ cm s}^{-1}$, (\circ) = k_{st} in units of $10^{-6} \text{ cm s}^{-1}$. Organic phase: TODGA in TPH-O. Aqueous phase: 0.5 M HNO_3 .

In order to get more insight into these results, interfacial tension measurements were performed by using Lecomte du Noüy ring method. The result is shown in Figure 6.

This plot shows that TODGA is surface active in TPH in the absence of 1-octanol

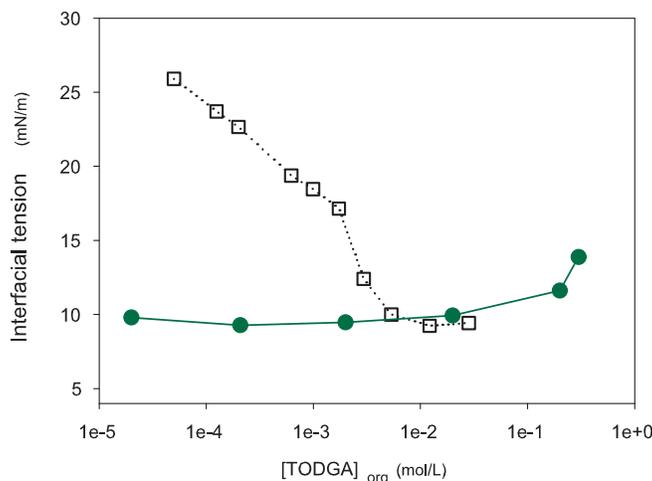


Figure 6: Interfacial tension at the interface: 3 M HNO₃ aqueous solution/organic phase, vs. TODGA concentration at 22°C. (□)= Organic phase comprised of TODGA in TPH; (●) = Organic phase comprised of TODGA in TPH-O.

(represented by squares in Figure 6), and that it is *not* surface active in the presence of 5 vol.% 1-octanol (full circles). These findings suggest that 1-octanol may replace TODGA at the interface when it is introduced in TPH. This interpretation is reinforced by the experimental observation that TODGA is not surface active when it is dissolved in pure 1-octanol (not shown).

Consequently, the slowdown of the kinetics observed in Figure 5 is counterintuitive, because one may expect the extraction process to be faster when more extractant is present in the organic phase, as has been for example observed in the past in a system involving dithizone as the extractant, which exhibited no surface activity.⁵² Indeed, one may expect that more TODGA in the organic phase would increase the probability of a TODGA molecule to reach the interface, hence also the encounter between an Eu(III) ion and a TODGA molecule there, which would accelerate the transfer. The fact that it is not so, and that both the extraction and back-extraction processes are affected, may be related to the aggregation properties of TODGA in the organic phase,⁵³ a phenomenon which does not occur in the case of dithizone. Nevertheless the precise way in which aggregation may cause the slowdown of the kinetics is unclear at present. This topic would require further experimental

investigations.

4.2 Effect of HNO₃ concentration

The effect of nitric acid concentration needs be investigated. ¹⁵²Eu(III) and ²⁴¹Am(III) were extracted from 0.5 M and 3 M HNO₃ solutions into a 0.2 M TODGA solution in TPH-O.

The results for the distribution ratio and the kinetic rate constants are collected in Table 4.

Table 4: Distribution ratios and kinetic rate constants for ¹⁵²Eu(III) and ²⁴¹Am(III) at the interface between 0.2 M TODGA in TPH-O and HNO₃ solutions of different concentrations.

Cation	Aqueous phase	K	k_{ex} (cm s ⁻¹)	
			aq → org	org → aq
¹⁵² Eu(III)	0.5 M HNO ₃	329	2.4×10^{-3}	6.9×10^{-6}
	3 M HNO ₃	1044	3.1×10^{-3}	3.0×10^{-6}
²⁴¹ Am(III)	0.5 M HNO ₃	58	2.8×10^{-3}	4.8×10^{-6}
	3 M HNO ₃	2977	3.9×10^{-3}	1.3×10^{-6}

As seen in this table, the distribution ratios K for Eu(III) and Am(III) were found to be strongly enhanced when [HNO₃] was varied from 0.5 M to 3 M (by factors of ~ 3.2 and 51, respectively). This behavior is in keeping with previous work.⁵⁰ It may be interpreted along the Le Châtelier principle because the extracted complex contains nitrate ions,⁵⁰ and maybe also nitric acid molecules as in the case of other extractants.⁵⁴

In contrast, the extraction rate constants, k_{ex} , for Eu(III) and Am(III) were increased but they did not vary as much as K did. They were of comparable high magnitudes for the two species, with k_{ex} values of a few 10^{-3} cm s⁻¹. They increased by about 30% when [HNO₃] was changed from 0.5 M to 3 M. This acceleration may be attributed to a higher rate of encounter for a Eu(III) or Am(III) ion with a nitrate ion and a nitric acid molecule (which combine to form the extracted complex⁵⁰) when the acid concentration is increased.

On the other hand, the stripping of these cations from TODGA solution to aqueous nitric acid was slow with k_{st} values of the order of a few 10^{-6} cm s⁻¹. Moreover, the stripping rate constants dropped by $\sim 60\%$ and 70% in the case of Eu(III) and Am(III), respectively, when [HNO₃] was increased from 0.5 M to 3 M. This slowdown may be caused by a more difficult

release of nitrate or nitric acid from the complex, at the interface, when nitric acid is more concentrated in the aqueous phase.

These results suggest that, in the industrial process, the co-extraction of lanthanides and actinides should be performed from concentrated nitric acid solutions (≥ 3 M) that will favor *both* the partitioning (higher K , which is well known) and the extraction kinetics (higher k_{ex} and lower k_{st}) of the elements.

5 Kinetics of stripping in the presence of aqueous ligands

The $\text{SO}_3\text{-Ph-BTP}$ ²⁸ and PTD ²⁹ molecules are promising aqueous ligands envisaged for the SANEX and GANEX An/Ln separation processes.⁵⁵ The effect of these ligands on the kinetics of stripping of Eu(III) and Am(III) was investigated.

5.1 Addition of $\text{SO}_3\text{-Ph-BTP}$

5.1.1 Effect of $\text{SO}_3\text{-Ph-BTP}$ concentration

Figure 7 shows the results for the inverse of the distribution ratios, $1/K = C_{aq}^{eq}/C_{org}^{eq}$ of Eu(III) and Am(III) between an organic 0.2 M TODGA solution in TPH-O and an aqueous 0.5 M HNO_3 solution for concentrations of $\text{SO}_3\text{-Ph-BTP}$ in the aqueous phase up to 40 mM. Figures 8 and S6 give the corresponding results for the stripping and extraction rate constants, respectively. The values of these quantities are given in the Supporting Information.

As expected, an increase of $[\text{SO}_3\text{-Ph-BTP}]_{aq}$ strongly increased the aqueous-to-organic distribution ratio $1/K$ of Eu(III) and Am(III) (see Figure 7) because the ligand tends to capture the solute in the aqueous phase. It is seen in Table S8 that the addition of only 10 mM $\text{SO}_3\text{-Ph-BTP}$ produces a rise of the value of $1/K$ by a factor of ~ 3 and ~ 400 in the case of Eu(III) and Am(III), respectively. Then doubling the concentration of the aqueous ligand

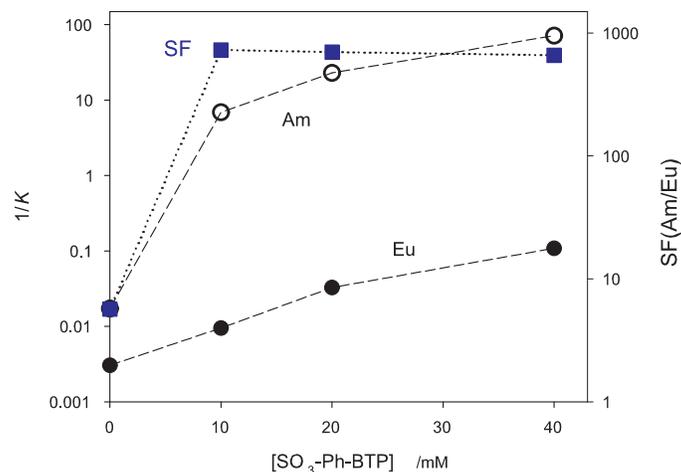


Figure 7: Aq/org distribution ratios ($1/K$, left scale), and separation factor ($SF(Am/Eu)$, right scale), of Eu(III) and Am(III) for an organic 0.2 M TODGA solution in TPH-O and an aqueous 0.5 M HNO_3 solution as a function of SO_3 -Ph-BTP concentration in aqueous phase up to 40 mM. (\bullet)= Eu(III); (\circ)= Am(III).

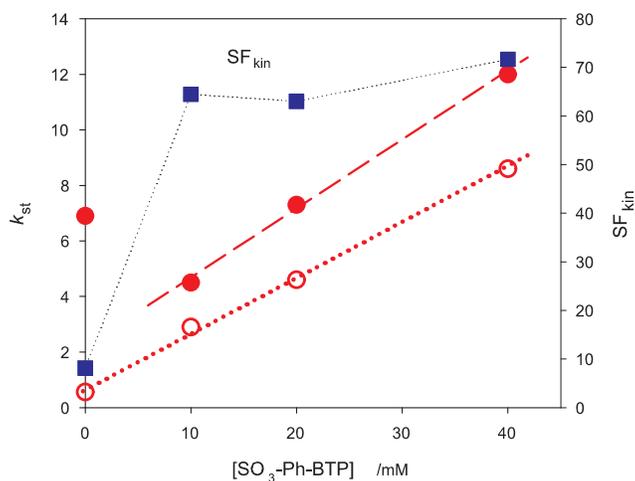


Figure 8: Stripping rate constants (left scale), and kinetic separation factor (SF_{kin} , right scale), for Eu(III) (in $10^{-6} \text{ cm s}^{-1}$) and Am(III) (in $10^{-4} \text{ cm s}^{-1}$) for the system considered in Figure 7. (\bullet)= Eu(III); (\circ)= Am(III)

causes $1/K$ to be increased by a factor of approximately 3 in both cases. This is reflected in the fact that the variation of $1/K$ is nearly linear in log-log scale (not shown) with a slope of ~ 1.7 for the two ions. This slope seems a little small in absolute value in view of reported association constants for the possible ligand-metal complexes,⁵⁶ which would suggest a slope between 2 and 3. Deviations from ideality involving the highly charged $\text{SO}_3\text{-Ph-BTP}$ ion (see Figure 3) in the aqueous phase might be partly responsible for the difference.

The Am/Eu separation factors, $\text{SF}(\text{Am}/\text{Eu}) = (1/K(\text{Am})) / (1/K(\text{Eu})) = K(\text{Eu}) / K(\text{Am})$ are high, between 727 at 10 mM $\text{SO}_3\text{-Ph-BTP}$ and 660 at 40 mM. This result shows the selectivity of the aqueous ligand $\text{SO}_3\text{-Ph-BTP}$ towards the actinide Am(III) through stripping.

It is observed in Figure 8 that the stripping rates for both $^{152}\text{Eu}(\text{III})$ and $^{241}\text{Am}(\text{III})$ exhibit a linear increase by addition of $\text{SO}_3\text{-Ph-BTP}$. In the case of Eu(III), k_{st} initially drops between 0 mM and 10 mM $\text{SO}_3\text{-Ph-BTP}$ before increasing linearly. The k_{st} value for Am(III) increases continuously over the whole range. With k_{st} values of $^{241}\text{Am}(\text{III})$ in the range of 10^{-4} to 10^{-3} cm s^{-1} , the stripping rate for $^{241}\text{Am}(\text{III})$ was rather fast, and greatly faster than for $^{152}\text{Eu}(\text{III})$. The ratio of the stripping rate constants, $k_{st}(\text{Am})/k_{st}(\text{Eu})$, varies from 63 for 10 mM $\text{SO}_3\text{-Ph-BTP}$ to 118 for 40 mM $\text{SO}_3\text{-Ph-BTP}$.

The behavior of the stripping rate constant vs. $[\text{SO}_3\text{-Ph-BTP}]$ makes a case for a mechanism in which the aqueous ligand captures the ion at the interface with a velocity that is proportional to its concentration. This may be expressed by writing $k_{st} = k_{st}^{(0)} + \lambda L$, for $L \geq 10$ mM, with $L = [\text{SO}_3\text{-Ph-BTP}]$, in which $k_{st}^{(0)}$ and λ are constants.

A look at Table S8 shows that the extraction kinetic rate constants have moderate values and are considerably lower for both $^{152}\text{Eu}(\text{III})$ and $^{241}\text{Am}(\text{III})$ when $[\text{SO}_3\text{-Ph-BTP}]_{aq}$ is added, especially in the case of Am(III). In the presence of 10 mM aqueous ligand, the values of k_{ex} for the two ions are reduced by a factor of ~ 5 and 80, respectively, as compared to a ligand-free solution. For each value of $[\text{SO}_3\text{-Ph-BTP}]_{aq}$, the extraction rate constant k_{ex} for $^{152}\text{Eu}(\text{III})$ is observed to be about 10 times that for $^{241}\text{Am}(\text{III})$.

It may be added that the ligand $\text{SO}_3\text{-Ph-BTP}$ does not accelerate the extraction kinetics

of Eu(III) and Am(III). Such an acceleration had been observed in the literature, e.g. by the acetate anion on the extraction of divalent cations like Co(II).⁵⁷⁻⁵⁹ An interpretation of this phenomenon was that the ligand may replace water molecules bound to the metal aquo-ion to form a kinetically more labile complex.^{57,58} The reverse situation is observed in the present case, with a ligand that binds very hard to the ion and slows down its kinetics of extraction.

From the important point of view of applications for the industrial processes SANEX and GANEX, it is interesting to look at the kinetic selectivity of the ligand in a stripping stage. This may be done by computing a stripping kinetic Am/Eu separation factor, defined as,

$$SF_{kin} = k_{st}(\text{Am})/k_{st}(\text{Eu}) \quad (10)$$

which quantity allows one to compare the respective fluxes of Am and Eu when an organic phase loaded with these elements is put in contact with a fresh aqueous solution.

The values of SF_{kin} are plotted in Figure 8. It is seen that this kinetic separation factor is strongly enhanced by addition of ligand, as is the separation factor at equilibrium, SF.

Therefore the separation of Am(III) from Eu(III) by stripping is greatly enhanced by SO₃-Ph-BTP, both thermodynamically *and* kinetically. This outcome will likely be quite favorable for An/Ln separation using short-time phase contacting extractors in SANEX and GANEX processes.

5.1.2 Effect of HNO₃ concentration (in the presence of SO₃-Ph-BTP)

In the extraction processes envisaged for the future, actinides may have to be re-extracted from aqueous solutions containing a hydrophilic ligand (e.g., SO₃-Ph-BTP or PTD) into an organic TODGA-based solvent by increasing the TODGA concentration and/or that of nitric acid. The kinetics of re-extraction of Am(III) by a 0.2 M TODGA solution was studied by increasing [HNO₃]_{aq} from 0.5 M to 2 M in the presence of 20 mM SO₃-Ph-BTP.

In Figure 9 are plotted the results for the re-extraction of $^{241}\text{Am}(\text{III})$ in these conditions (the corresponding data are given in the Supporting Information). It is worth noting that the 3 plots exhibit a linear behavior when plotted with log-log axes. These results may be compared with those displayed in Table 4, which gives data for the effect of HNO_3 in the absence of $\text{SO}_3\text{-Ph-BTP}$.

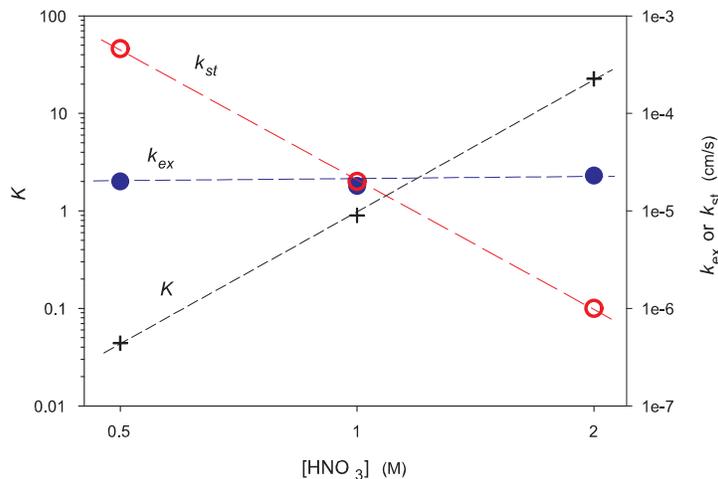


Figure 9: Re-extraction and stripping rates (right scale), and K values (left scale), for $^{241}\text{Am}(\text{III})$ between aqueous 20 mM $\text{SO}_3\text{-Ph-BTP}$ solutions of different HNO_3 concentrations and organic 0.2 M TODGA in TPH-O. (+) = K ; (●) = k_{ex} ; (○) = k_{st} .

First, it is seen in Figure 9 that the partition coefficient, $K = k_{ex}/k_{st}$, is dramatically enhanced (by ~ 500 times) when $[\text{HNO}_3]_{aq}$ is varied from 0.5 M to 2 M. This outcome is similar to that found in section 4.2 (Table 4), but the magnitude of the enhancement is much larger than in the absence of $\text{SO}_3\text{-Ph-BTP}$. The interpretation of the effect of adding nitric acid is identical to that of section 4.2, invoking the Le Châtelier principle.

Then, it is noticed in Figure 9 that the extraction rate constant remains approximately constant vs. $[\text{HNO}_3]$ (it was found to increase slightly in section 4.2). In contrast, the stripping rate constant decreases by more than two orders of magnitude when $[\text{HNO}_3]_{aq}$ is increased from 0.5 M to 2 M. This drop is therefore much more pronounced in the presence of $\text{SO}_3\text{-Ph-BTP}$ than when it is absent (see Table 4, suggesting a decline by a factor of about 3 in this concentration range). The cause for this drop might be similar to that put forward

in section 4.2, i.e. a more difficult release of nitrate and nitric acid from the complex at the interface.

In the industrial process, the kinetics of this extraction step will therefore be somewhat faster when using a more concentrated nitric acid solution, because the back-extraction rate constant k_{st} will then be much smaller (k_{ex} being approximately constant).

The $\text{SO}_3\text{-Ph-BTP}$ ligand (in which $\text{SO}_3\text{-Ph-BTP}$ denotes the tetravalent anionic form of the ligand) may become protonated in the presence of nitric acid (probably on a N donor atom of the central pyridine group⁶⁰), with a first protonation constant, $\log_{10} K_1 \simeq 0.5$.⁶¹ The resulting $\text{H}\cdot\text{SO}_3\text{-Ph-BTP}$ molecule, may not complex the Am(III) ions.⁶¹ As the concentration of nitric acid in the aqueous phase is increased, more ligand becomes (mono-)protonated, which decreases the amount of unprotonated ligand available to complex and strip Am(III). This process provides a qualitative interpretation of the variation of K . However, a simple calculation, using the value of K_1 and assuming thermodynamic ideality in the aqueous phase, shows that the drawdown of unprotonated ligand alone cannot explain the huge increase of K and the big drop of k_{st} observed when the acid concentration is varied in this range. We note that a similar difficulty in interpreting the equilibrium data was encountered in the case of the uranyl cation in the presence of this same ligand.⁶² Further investigations would be needed to better understand the partitioning and kinetic behavior of this system.

5.2 Addition of PTD

As mentioned in the Introduction section, a drawback of the $\text{SO}_3\text{-Ph-BTP}$ ligand is that it does not comply with the CHON principle suitable for complete incineration. On the contrary, the PTD ligand satisfies this condition. This molecule has proved to be a good alternative for stripping actinides from organic solutions.^{29,30} Its effect on the kinetics was studied in this work.

5.2.1 Effect of PTD concentration

The results for the stripping and extraction rate constants, for 0.5 M HNO₃ and 0.2 M TODGA, are shown in Figures 10 (for Eu(III)) and 11 (for Am(III)), and collected in the Supporting Information. The values for the distribution ratio are given in the latter.

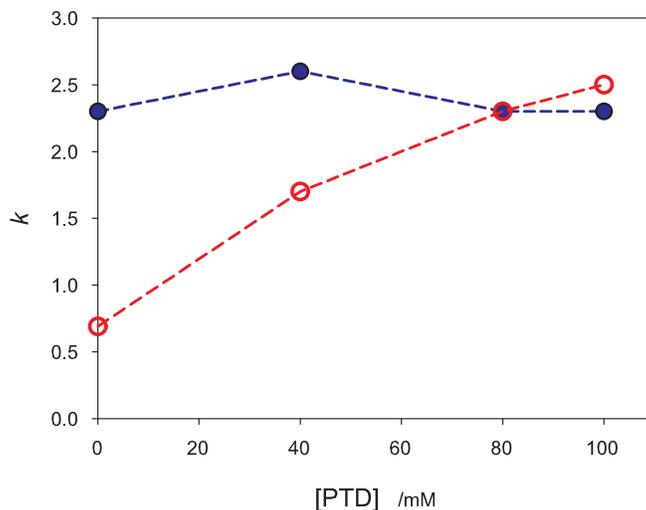


Figure 10: Extraction (●) and stripping (○) kinetic rate constants for ¹⁵²Eu(III) as a function of PTD concentration in aqueous phase in 0.5 M HNO₃ solution, and organic phase: 0.2 M TODGA in TPH-O. Plots in units of 10⁻³ cm s⁻¹ for k_{ex} , and of 10⁻⁵ cm s⁻¹ for k_{st} . Dashed lines are guides for the eye.

It is seen in Table S10 that the aq/org distribution ratio, $1/K$, rises with the concentration of PTD, as expected because PTD forms a complex with the metal ion in aqueous phase.

Figure 10 shows that the extraction rate constant for Eu(III) is nearly constant (within experimental uncertainty) with respect to the concentration of PTD. The stripping rate constant for Eu(III) increases significantly with [PTD]. In Figure 11 it is seen that the behavior of the extraction rate constant in the case of americium is quite different from that of europium, with a k_{ex} for Am(III) that decreases sharply with the amount of added PTD. The extraction rate constant for Eu(III) is a few times larger than that for Am(III).

The rate constants for back-extraction, k_{st} , increase notably with [PTD] for the two ions, with a 40-fold increase from 0 to 100 mM of PTD in the case of Am(III). The stripping rate constant for Eu(III) is about 2 orders of magnitude smaller than that for Am(III) (but both

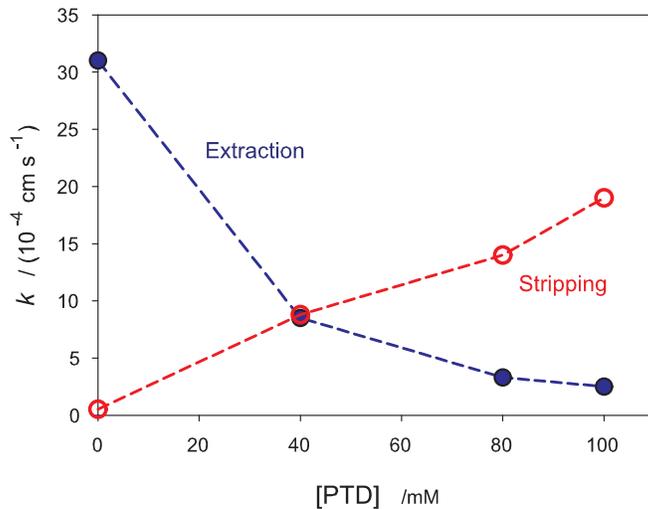


Figure 11: Extraction (●) and stripping (○) kinetic rate constants for $^{241}\text{Am}(\text{III})$ as a function of PTD concentration in aqueous phase in 0.5 M HNO_3 solution, and organic phase: 0.2 M TODGA in TPH-O. Plots in units of $10^{-4} \text{ cm s}^{-1}$. Dashed lines are guides for the eye.

increase with [PTD]).

The fact that the rate constant for back-extraction increases nearly linearly with [PTD] suggests, similarly to the case of $\text{SO}_3\text{-Ph-BTP}$, that PTD captures the ion at the interface with a probability that is proportional to [PTD], and takes it to the aqueous phase. One may write, $k_{st} = k_{st}^{(0)} + \lambda [\text{PTD}]$, in which $k_{st}^{(0)}$ is the rate constant in the absence of PTD and λ is a constant.

To interpret the steep variation of k_{ex} in the case of $\text{Am}(\text{III})$ it was examined whether the rate constant could be expressed as $k_{ex} = k_{ex}^{(0)}\alpha_0$, in which α_0 is the fraction of free Am^{3+} ion, and $k_{ex}^{(0)}$ is the extraction rate constant in the absence of PTD. Moreover, it was assumed that the complexed americium is predominantly in the form of one complex, $\text{Am}(\text{PTD})_n$ (with n a constant integer). A mass action law was used for this complex, and different values of n ($n= 1, 2$ and 3) were tested. A plot of $1/k_{ex}$ as a function of $[\text{PTD}]^2$ approximately gave a straight line on the whole PTD concentration range (not shown), thus suggesting a stoichiometry of 2 for the main complex ($\text{Am}(\text{PTD})_2$). Values of $n = 1$ or 3 did not give a straight line. The present result is in keeping with the speciation found in

ref. 30 (see Figure 5 of this reference) in which the predominant 1-2 complex was identified in the case of Cm(III), whose behavior is very close to that of Am(III).^{29,30} The fact that k_{ex} is constant when PTD is added to a Eu(III) solution might be related to the much lower complexation constant for this metal ion.³⁰

As in the case of SO₃-Ph-BTP, it is interesting to look at the Am/Eu equilibrium and kinetic separation factors, SF and SF_{kin} (eq 10), respectively. They are plotted in Figure 12 as a function of the PTD concentration. As previously observed for SO₃-Ph-BTP, both separation factors are strongly enhanced by addition of ligand.

Therefore, here too, the separation of Am(III) from Eu(III) by stripping is greatly enhanced by the aqueous ligand (PTD), both thermodynamically *and* kinetically. This should be quite favorable for An/Ln separation with the CHON-compliant stripping agent PTD, using short-time phase contacting extractors in SANEX and GANEX processes.

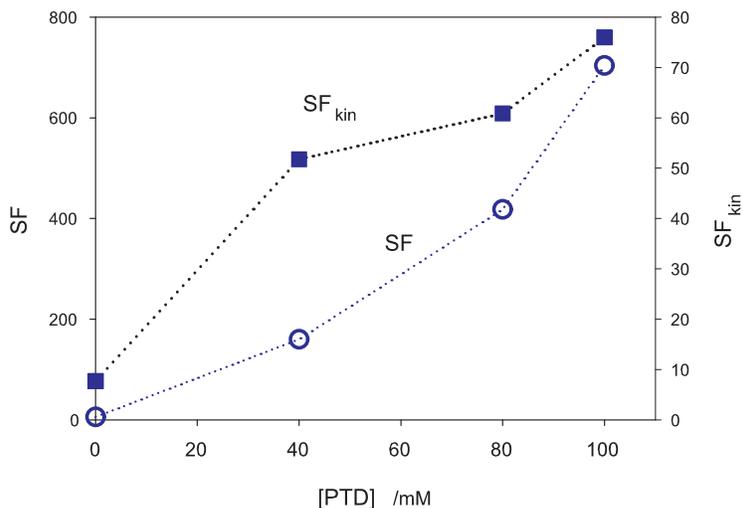


Figure 12: Am/Eu equilibrium (SF, left scale) and kinetic (SF_{kin}, right scale) separation factors in the case of PTD.

5.2.2 Effect of HNO₃ concentration (in the presence of PTD)

As done in section 5.1.2 in the case of SO₃-Ph-BTP, the influence of the HNO₃ concentration was studied in the case of Am(III), in conditions suitable for the stripping of actinide

ions,^{29,30,63} namely with [PTD] = 80 mM and 0.2 M TODGA, for 2 values of aqueous nitric acid concentration, [HNO₃] = 0.1 M and 0.5 M.

The results are collected in Table 5. They may be compared with those of sections 4.2 (absence of hydrophilic ligand) and 5.1.2 (use of SO₃-Ph-BTP ligand).

Table 5: Distribution ratio, K , and kinetic rate constants for ²⁴¹Am(III) at the interface between an organic phase of 0.2 M TODGA in TPH-O and aqueous solutions of various HNO₃ concentrations with [PTD] = 80 mM.

[HNO ₃]	K	k_{ex} (cm s ⁻¹)	k_{st} (cm s ⁻¹)
0.1 M	1.05×10^{-3}	1.9×10^{-7}	1.8×10^{-4}
0.5 M	0.237	4.1×10^{-4}	17×10^{-4}

In this table we see that K is strongly enhanced (by a factor of more than 200) when [HNO₃] is increased from 0.1 M to 0.5 M. This behavior is consistent with a previous study³⁰ in which decomplexation of the metal cation was observed when increasing the acid concentration, and it is similar to what was found in sections 4.2 and 5.1.2. As in these sections, the Le Châtelier principle may be invoked to explain the increase of the distribution ration with the acid concentration.

At the same time the extraction kinetics is dramatically enhanced by more than 3 orders of magnitude, and the stripping kinetics is enhanced by about 10 times. The huge enhancement of k_{ex} contrasts strongly with the mild rise observed in the absence of aqueous ligand, and the flat behavior found when using SO₃-Ph-BTP. The softer rise of k_{st} also contrasts with the drops found without ligand and in the presence of SO₃-Ph-BTP. The increase observed for k_{ex} may be explained along the same lines as those put forward in sections 4.2 and 5.1.2. The origin of the acceleration of the stripping step is unknown at this stage. The aqueous ligand PTD therefore behaves quite differently than the other ligand SO₃-Ph-BTP.

The results of this section suggest the use of concentrated nitric acid solutions in the process at the industrial scale, in order to accelerate the re-extraction of Am(III).

6 Selective extraction with CyMe₄-BTBP (+ TODGA) in 1-octanol

The combination of 10 mM CyMe₄-BTBP with 5 mM TODGA extractant, acting as a phase-transfer catalyst, in 1-octanol has been retained for actinide selective extraction in the one-cycle SANEX (1c-SANEX) process.^{64,65}

The kinetics of extraction of Eu(III) and Am(III) were investigated. The transfer was made from an aqueous 3 M HNO₃ solution into organic solutions of CyMe₄-BTBP + TODGA in 1-octanol. The phase-transfer catalyst, TODGA, was added in order to accelerate the extraction kinetics.

The results are shown in Table 6. The actinide-selective extractant CyMe₄-BTBP at 10 mM yielded a high separation factor, SF(Am/Eu)= $K(\text{Am})/K(\text{Eu})$, of about 105 when it was used alone, but a very slow extraction rate, with k_{ex} on the order of 10^{-7} – 10^{-6} cm s⁻¹. Very slow extraction kinetics were obtained for Eu(III) and Am(III). The rate constant for Am(III) ($k_{ex}(\text{Am}) = 7.2 \times 10^{-6}$ cm s⁻¹) was nevertheless 24 times larger than that for Eu(III).

Table 6: Effect of the phase-transfer catalyst TODGA on the kinetics of extraction by CyMe₄-BTBP. Aqueous phase: 3 M HNO₃ solution spiked with ¹⁵²Eu or ²⁴¹Am. Organic phase: 5 mM or 10 mM CyMe₄-BTBP in 1-octanol, with or without 5 mM TODGA.

[CyMe ₄ -BTBP]	[TODGA]	$K(\text{Eu})$	$K(\text{Am})$	SF(Am/Eu)	$k_{ex}(\text{Eu})$ /(10 ⁻⁵ cm s ⁻¹)	$k_{ex}(\text{Am})$ /(10 ⁻⁵ cm s ⁻¹)
10 mM	0	0.281	29.3	105	0.0302	0.720
0	5 mM	0.394	0.142	0.360	10.5	29.4
Sum of previous entries		0.675	29.4		10.5	30.1
10 mM	5 mM	0.980	33.1	33.8	8.76	6.19
5 mM	5 mM	0.593	9.07	15.3	8.12	5.72

Fast extraction kinetics have been found in section 4.1 for TODGA alone in TPH-O. Here, even with a small amount of TODGA (5 mM), a rather high extraction rate constant was obtained for Am(III): $k_{ex}(\text{Am}) \sim 2.9 \times 10^{-4}$ cm s⁻¹, which is 2.8 times that for Eu(III): $k_{ex}(\text{Eu}) \sim 1.1 \times 10^{-4}$ cm s⁻¹ (see Table 6). However, the distribution ratios $K(\text{Eu})$ and

$K(\text{Am})$ were low.

A slight synergistic effect is observed in Table 6, where the value of the distribution ratio K for the mixture 10 mM $\text{CyMe}_4\text{-BTBP}$ + 5 mM TODGA is observed to be larger than the sum of the individual K values for the two extractants, for both Eu(III) ($0.98 > 0.28+0.39 = 0.67$) and Am(III) ($33.1 > 29.3+0.14 = 29.4$) extraction. Nonetheless, the separation factor is considerably reduced to $\text{SF}(\text{Am}/\text{Eu})\sim 34$ as compared to that with $\text{CyMe}_4\text{-BTBP}$ alone ($\text{SF} = 105$).

As regards the kinetics, it is clearly observed that the extraction kinetics of Eu(III) and Am(III) are greatly accelerated when the extractants are combined as compared to the case of $\text{CyMe}_4\text{-BTBP}$ alone: the rate constants $k_{ex}(\text{Eu})$ and $k_{ex}(\text{Am})$ are about 290 and 9 times higher, respectively (passing from $0.0302\times 10^{-5} \text{ cm s}^{-1}$ to $8.76\times 10^{-5} \text{ cm s}^{-1}$ in the case of Eu(III), and from $0.720\times 10^{-5} \text{ cm s}^{-1}$ to $6.19\times 10^{-5} \text{ cm s}^{-1}$ in that of Am(III)). However, the extraction rate constant k_{ex} is of the order of a few $10^{-5} \text{ cm s}^{-1}$, which is a moderate value.

It is seen on the last line of Table 6 that, when $[\text{CyMe}_4\text{-BTBP}]$ is lowered from 10 mM to 5 mM in the presence of 5 mM TODGA catalyst, the extraction kinetics of Eu(III) and Am(III) remain nearly unchanged (within the experimental uncertainty), but the distribution ratios are significantly lowered. This observation is consistent with the expectation that the kinetics are controlled by TODGA, not by $\text{CyMe}_4\text{-BTBP}$. However, it is also seen in Table 6 that the extraction rate constants for the mixture are lower than with TODGA alone ($8.8 < 10.5$ for $k_{ex}(\text{Eu})$, and above all $6.2 \ll 29.4$ for $k_{ex}(\text{Am})$).

As regards the 1c-SANEX process, it seems that the mixture 10 mM $\text{CyMe}_4\text{-BTBP}$ + 5 mM TODGA is a good formulation that preserves a rather high Am/Eu separation factor ($\text{SF}\sim 34$) and accelerates the extraction as compared to pure $\text{CyMe}_4\text{-BTBP}$. However, the extraction kinetics of Eu(III) and Am(III) are comparable ($k_{ex}(\text{Am})$ even seems to be a little smaller than $k_{ex}(\text{Eu})$). This point may reduce the interest of using a centrifugal extractor for the separation of Am(III) from Eu(III) in the case of $\text{CyMe}_4\text{-BTBP}$.

7 Effect of temperature

The influence of temperature on the kinetics of extraction was examined at temperatures of 22°C and 35°C (in a thermostated chamber). The results for the extraction of Eu(III) and Am(III) by 0.2 M TODGA in TPH-O, and by 10 mM CyMe₄-BTBP + 5 mM TODGA in 1-octanol, are collected in Tables 7 and 8, respectively.

Table 7: Influence of the temperature on the extraction kinetic rate constant for ¹⁵²Eu and ²⁴¹Am. Aqueous phase: 0.5 M HNO₃ + 20 mM SO₃-Ph-BTP. Organic phase: 0.2 M TODGA in TPH-O pre-equilibrated with the aqueous phase.

Ion	T	K	k_{ex} (cm s ⁻¹)	k_{st} (cm s ⁻¹)
¹⁵² Eu(III)	22°C	30.6	2.2×10^{-4}	7.3×10^{-6}
	35°C	11.6	2.5×10^{-4}	22×10^{-6}
²⁴¹ Am	22°C	0.0437	2.0×10^{-5}	4.6×10^{-4}
	35°C	0.0187	2.2×10^{-5}	12×10^{-4}

Table 8: Influence of the temperature on the extraction kinetic rate constant for ¹⁵²Eu and ²⁴¹Am. Aqueous phase: 3 M HNO₃. Organic phase: 10 mM CyMe₄-BTBP + 5 mM TODGA in 1-octanol (pre-equilibrated with the aqueous phase).

Ion	T	K	k_{ex} (cm s ⁻¹)	k_{st} (cm s ⁻¹)
¹⁵² Eu(III)	22°C	0.819	8.8×10^{-5}	9.0×10^{-5}
	35°C	0.674	6.1×10^{-5}	9.1×10^{-5}
²⁴¹ Am(III)	22°C	31.3	6.2×10^{-5}	2.0×10^{-6}
	35°C	22.4	5.6×10^{-5}	2.5×10^{-6}

An increase in the temperature from 22°C to 35°C lowers the distribution ratios K of Eu(III) and Am(III) by a factor of ~ 2 or 3 in the first case, and by about 30% in the second, indicating that the extraction reaction for both cations is exothermic. In other words, the activation energy of K is negative, as is often the case in extraction systems. This peculiarity has been shown to have potentially a strong influence on the apparent activation energy of the overall extraction process, and may lead to incorrect interpretations about the nature of the extraction regime (diffusion vs. kinetic control).³⁴

The values of the activation energies calculated using the K results at the two temperatures of 22°C and 35°C are: -24.5 kJ mol⁻¹ and -21.4 kJ mol⁻¹ for the data of Table 7 in

the case of Eu(III) and Am(III), respectively, and of -4.9 kJ mol^{-1} and -8.4 kJ mol^{-1} for the data of Table 8 for Eu(III) and Am(III), respectively.

The extraction kinetics of Eu(III) might be slightly accelerated by an increase of T from 22°C to 35°C in Table 7, and they seem to be moderately decelerated in Table 8. In contrast, they do not exhibit a clear trend in this temperature range in the case of Am(III). Although a small reduction is observed in the rate constants $k_{ex}(\text{Am})$, this variation is not significant in view of an uncertainty of the order of 10% or 20% (see section 3).

8 Conclusion

In this study several improvements have been brought to the RMC technique. Two types of Millipore membranes (hydrophilic vs. hydrophobic) have been used to contain the aqueous or the organic phase, respectively. The membranes were glued on the base of the plastic cylinder by using liquid Kapton. The lower part of the cylinder itself was covered with Kapton in order to isolate it from the solution in which it was immersed.

The results obtained in section 3 by performing extraction and stripping kinetic experiments were in agreement within experimental uncertainty. The values of the diffusion coefficients of the solutes Eu(III) and Am(III) in the phases, which are required in the treatment of the kinetic data, were measured using the RMC. The validity of the procedure was confirmed by also measuring separately a few diffusion coefficients by using the closed capillary technique. Measuring the D 's with the RMC makes this step much easier and shorter in time than with the capillary technique.

Extraction and stripping of Eu(III) and Am(III) were studied for various concentrations of nitric acid and TODGA, in mixtures of CyMe₄-BTBP with TODGA, and in the presence of the aqueous ligands SO₃-Ph-BTP and PTD. It was somewhat striking to find that TODGA is not surface active at the interface between nitric acid and TPH-O, which is in contrast with the case of TODGA in *n*-dodecane.

The kinetic data obtained in this work will be used as input parameters in simulation codes (such as, e.g., PAREX, developed at CEA^{22,23}) for a modeling of separation processes carried out in extractors (e.g., centrifugal) that operate with a short contact time between the phases.

The experimental results, obtained with TODGA and the two aqueous stripping ligands, show that faster transfer kinetics are associated with higher partitioning for Am(III) over Eu(III). This favorable outcome bodes well for future efficient actinide/lanthanide separation in the nuclear reprocessing industry.

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ASSOCIATED CONTENT

Supporting Information

This information is available free of charge via the Internet at <http://pubs.acs.org/>.

Details on the practical use of the RMC; Densities, and dynamic and kinematic viscosities of solutions; Diffusion coefficients; Distribution ratios and kinetic rate constants.

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9 Abstract Graphic For Table of Contents Only

