

Liquid-liquid extraction of rare earths elements from nitrate media in DMDOHEMA/ionic liquid system: performance and mechanism studies.

Cesar L. Usma,* S. Dourdain,* G. Arrachart and S. Pellet-Rostaing

Supplementary Materials

Table S1

Distribution coefficients and selectivity values for $[\text{HNO}_3] = 3.0 \text{ M}$

Organic solvent	[DMDOHEMA] (mol/L)	Distribution coefficient (D)			Selectivity (S)	
		La^{3+}	Eu^{3+}	Fe^{3+}	$S_{\text{La/Fe}}$	$S_{\text{Eu/Fe}}$
[EOPip ⁺] [NTf ₂ ⁻]	0	<0.01	<0.01	<0.01	-	-
	0.05	0.03	0.07	0.04	0.7	1.6
	0.1	0.28	0.75	0.25	1.1	3.0
	0.2	2.7	6.5	0.43	6.4	15
	0.4	18	34	0.74	24	46
[EBPip ⁺] [NTf ₂ ⁻]	0	<0.01	<0.01	<0.01	-	-
	0.05	<0.01	0.04	<0.01	-	-
	0.1	0.32	1.69	0.01	62	327
	0.2	7.6	33	0.03	262	1138
	0.4	152	283	0.26	587	1092
Dodecane	0	<0.01	<0.01	<0.01	-	-
	0.05	<0.01	<0.01	<0.01	-	-
	0.1	0.11	0.07	0.04	3.1	1.9
	0.2	0.77	0.50	0.48	1.6	1.1
	0.4	3.17	2.01	1.84	1.7	1.1

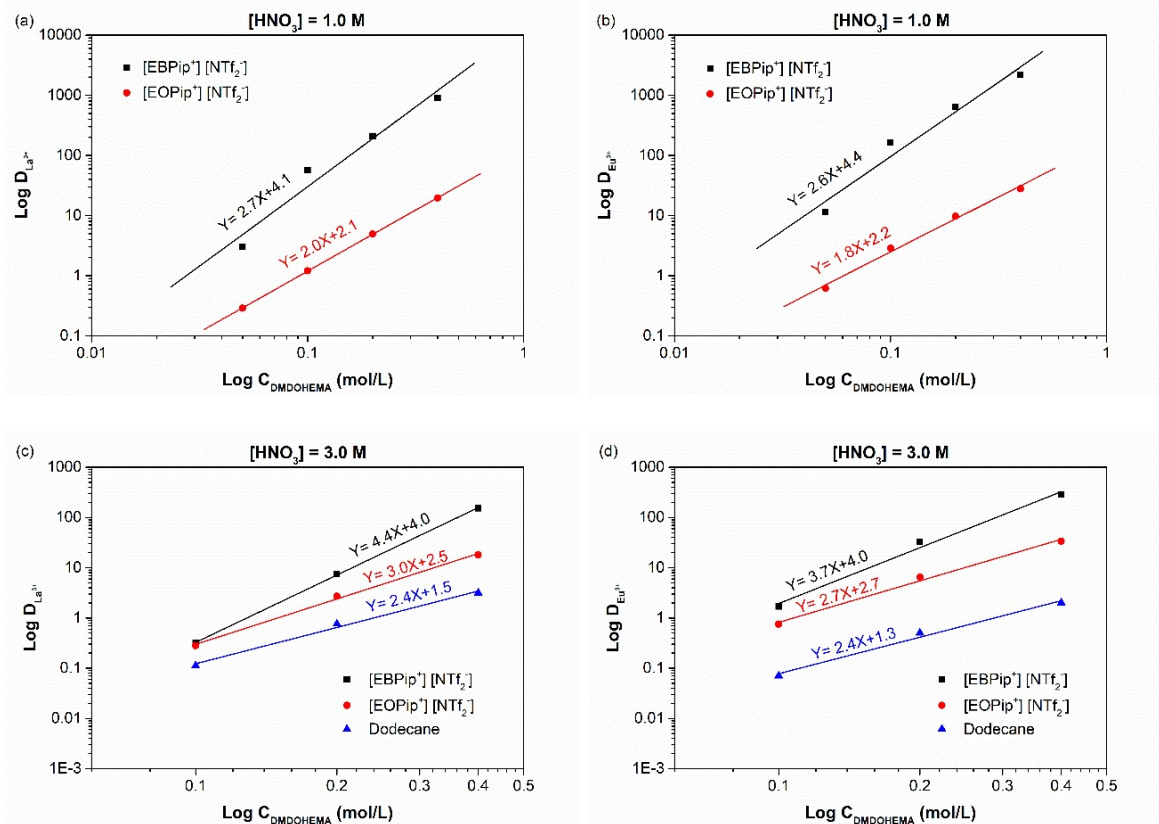


Fig. S1 Slope analysis in ILs and dodecane for: (a and c) La^{3+} and (b and d) Eu^{3+} . Nitric acid concentration 1 M (top) and 3 M (bottom).

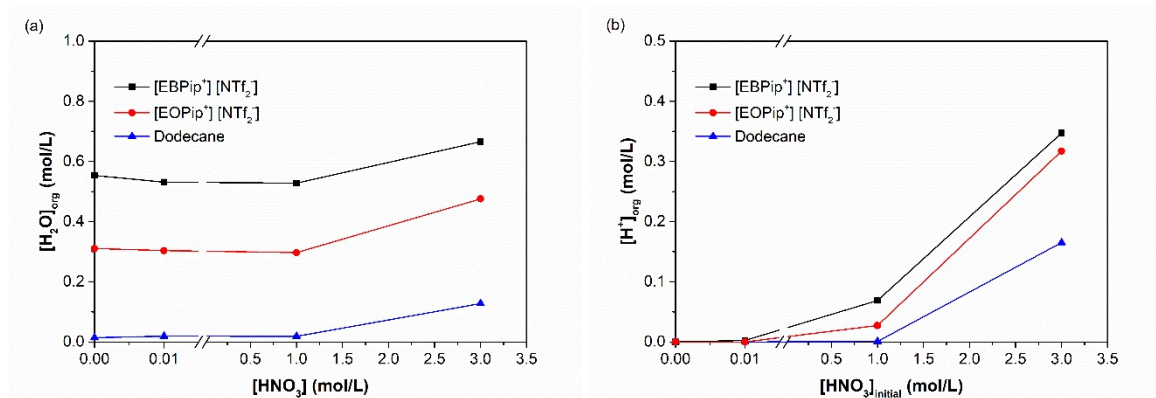


Fig. S2 (a) Water and (b) H^+ concentrations in the organic phase, after metal extraction, as a function of the nitric acid concentration for $[\text{DMDOHEMA}] = 0.2 \text{ M}$.

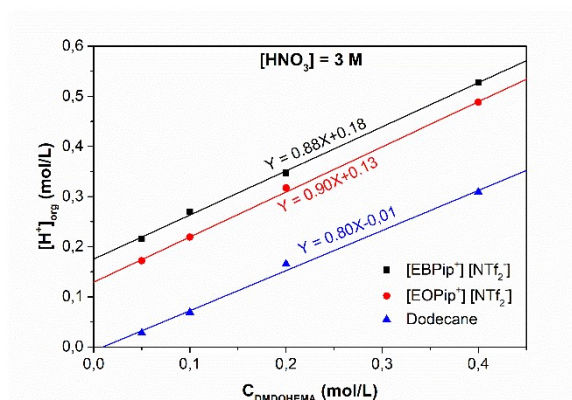


Fig. S3 H^+ concentration in the organic phase, after metal extraction, in function of the DMDOHEMA concentration for $[\text{HNO}_3] = 3 \text{ M}$.

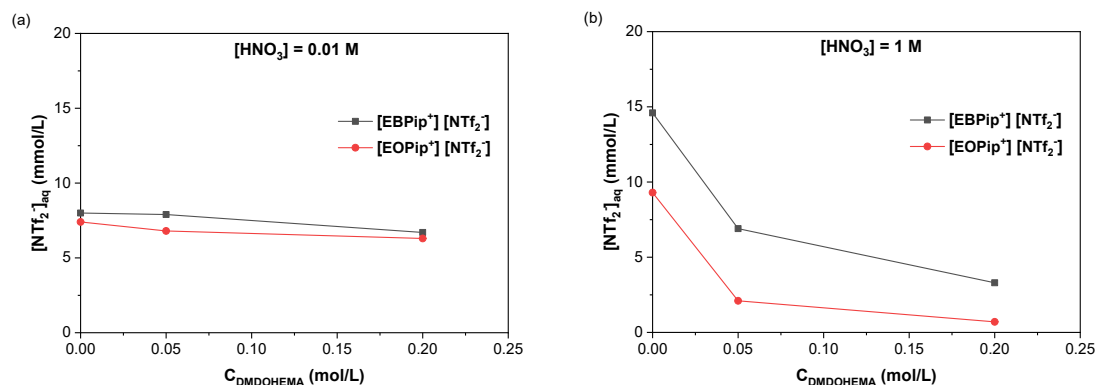


Fig. S4 Bistriflimide anion concentration in the aqueous phase as a function of the $[\text{DMDOHEMA}]$ when $[\text{EBPip}^+][\text{NTf}_2^-]$ or $[\text{EOPip}^+][\text{NTf}_2^-]$ are used as organic solvent, for an initial nitric acid concentration of: (a) 0.01 M and (b) 1.0 M.

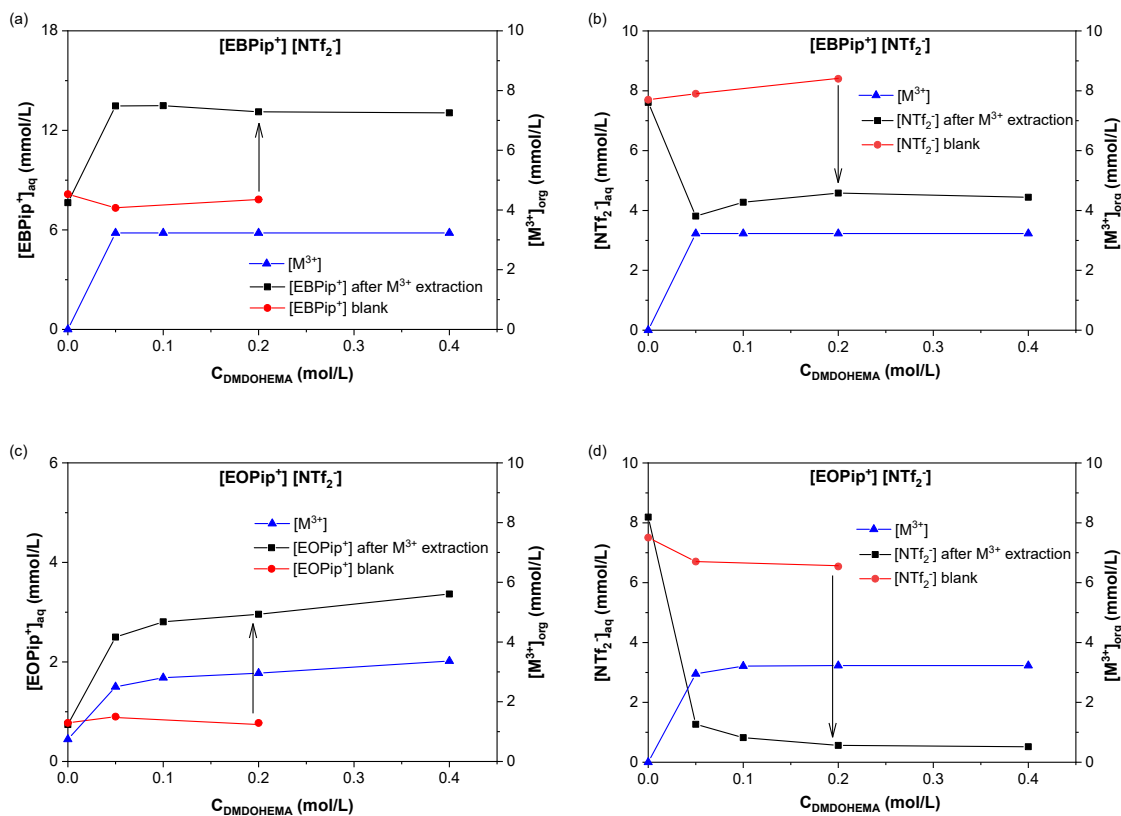


Fig. S5 a) Pip⁺ and b) NTf₂⁻ concentrations in the aqueous phase for [EBPip⁺] [NTf₂⁻]; c) Pip⁺ and d) NTf₂⁻ concentrations in the aqueous phase for [EOPip⁺] [NTf₂⁻], as a function of the DMDOHEMA concentration and for an initial nitric acid concentration of 0 M in the aqueous phase.

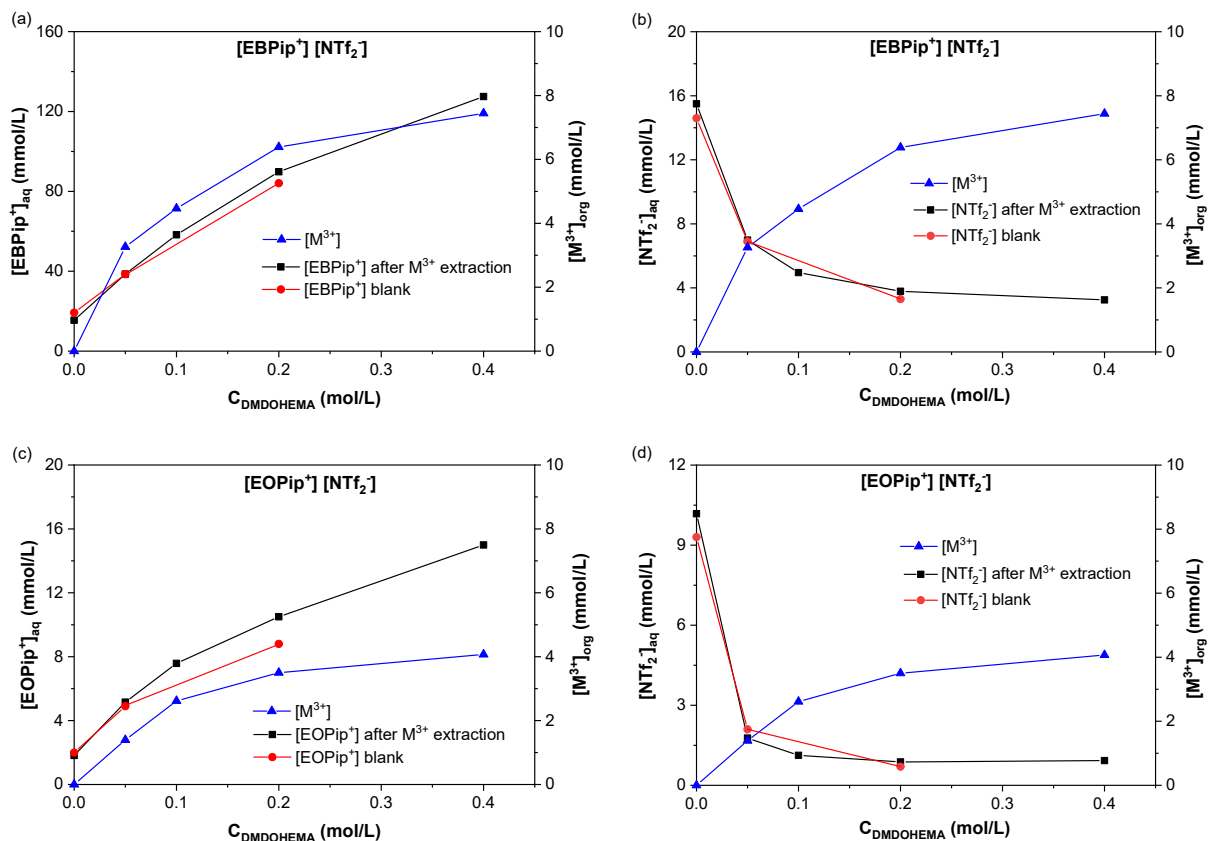


Fig. S6 a) Pip⁺ and b) NTf₂⁻ concentrations in the aqueous phase for [EBPip⁺] [NTf₂⁻]; c) Pip⁺ and d) NTf₂⁻ concentrations in the aqueous phase for [EOPip⁺] [NTf₂⁻], as a function of the DMDOHEMA concentration and for an initial nitric acid concentration of 1 M in the aqueous phase.