

Supplementary Material

Rare earth elements as tracers of active colloidal organic matter composition

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Concentrations of REE as determined by ICP-MS

The ICP-MS used in the study was 7700x from Agilent. The operating conditions used during the measurements are presented in Table S1.

Table S1: Instrument operating conditions

Plasma conditions	
RF Power	1450-1550W
Carrier Gas	15 L/min
Auxiliary argon flow	1 L/min
Nebulizer micromist- argon flow	0.9- 1.0 L/min
CeO ⁺ /Ce ⁺	1%
Ce ²⁺ /Ce ⁺	<2 %

Calibrations were performed using synthetic multi-elemental solutions prepared in 2% HNO₃. Instrument drift was monitored and corrected by spiking each sample with an internal standard (Re, Rh) introduced on-line using a peristaltic pump. In order to suppress some spectroscopic interferences, mathematical corrections were required and calculated as a function of the oxide level. The choice of the measured isotopes and interferences applied is summarized in Table S2. The oxide production level was around 1 %.

Table S2. Preferred isotopes for REE analysis, isotopic abundance in %, oxides and hydroxides interference on analyte.

Element	Isotope Mass	Abundance (%)	(Hy)Oxide interferences	Interferences corrections applied
La	139	99.91		
Ce	140	88.48		
Pr	141	100		
Nd	146	17.19		
Sm	147-149	15	¹³⁰ BaOH	
Eu	151-153	47.8-52.2	¹³⁵ BaO, ¹³⁷ BaO, ¹³⁶ BaOH,	Eu (153) corrected from ¹³⁷ BaO Eu (151) corrected from ¹³⁵ BaO
Gd	157-158	15.65-24.84	¹⁴² CeO, ¹⁴² NdO, ¹⁴¹ PrO, ¹⁴¹ PrOH	Gd (157) corrected from ¹⁴⁰ CeO and ¹⁴¹ PrO
Tb	159	100	¹⁴³ NdO	Tb(159) corrected from ¹⁴³ NdO
Dy	163	24.9	¹⁴⁷ SmO, ¹⁴⁶ NdOH,	Dy(163) corrected from ¹⁴⁷ Sm
Ho	165	100	¹⁴⁹ SmO	Ho(165) corrected from ¹⁴⁹ SmO
Er	166	33.6	¹⁵¹ EuO	Er(166) corrected from ¹⁵⁰ SmO
Tm	169	100		
Yb	172-174	21.9-31.8	¹⁵⁷ GdOH	Yb(174) corrected from ¹⁵⁸ GdO
Lu	175	97.41	¹⁵⁹ TbO	Lu(175) corrected from ¹⁵⁹ TbO

The detection limits for all the REE are presented in Table S3.

Table S3. Detection limits obtained on HP 7700x ICP-MS, calculated from the AFNOR standards and based on the blank measurement

Element	Isotope	Detection limit (ppt)	Detection limit (mol L ⁻¹)
La	139	0.15	1.1 X 10 ⁻¹²
Ce	140	0.14	1.0 X 10 ⁻¹³
Pr	141	0.06	4.3 X 10 ⁻¹³
Nd	146	0.16	1.1 X 10 ⁻¹²
Sm	147	0.23	1.5 X 10 ⁻¹²
Eu	153	0.10	6.6 X 10 ⁻¹³
Gd	157	0.22	1.4 X 10 ⁻¹²
Gd	158	0.13	8.3 X 10 ⁻¹³
Tb	159	0.04	2.5 X 10 ⁻¹³
Dy	163	0.17	1.0 X 10 ⁻¹²
Ho	165	0.05	3.0 X 10 ⁻¹³
Er	166	0.15	9.0 X 10 ⁻¹³
Tm	169	0.04	2.4 X 10 ⁻¹³
Yb	174	0.17	9.8 X 10 ⁻¹³

ICP-MS analysis included three consecutive replicate measurements, producing a repeatability error less than 2%. To control the quality of the REE element measurements, the certificated reference material SLRS-6 standards (National Research Council –CNRC Canada) was used.

Concentrations of REE added and present naturally in the HS.

Table S4. Average concentrations of the REE added in the different experiments in mol L⁻¹.

	PLFA (mol L ⁻¹)	AHA (mol L ⁻¹)	LHA (mol L ⁻¹)
La	4.5E-09	4.4E-09	9.6E-10
Ce	4.4E-09	4.4E-09	9.5E-10
Pr	4.4E-09	4.3E-09	9.4E-10
Nd	4.3E-09	4.2E-09	9.2E-10
Sm	4.1E-09	4.1E-09	8.8E-10
Eu	4.1E-09	4.0E-09	8.8E-10
Gd	3.9E-09	3.9E-09	8.5E-10
Tb	3.9E-09	3.8E-09	8.4E-10
Dy	3.8E-09	3.8E-09	8.2E-10
Ho	3.8E-09	3.7E-09	8.1E-10
Er	3.7E-09	3.6E-09	8.0E-10
Tm	3.7E-09	3.6E-09	7.9E-10
Yb	3.6E-09	3.5E-09	7.7E-10
Lu	3.5E-09	3.5E-09	7.6E-10

Table S5. Average REE concentrations present naturally in PLFA, AHA and LHA (detection limit = DL).

	PLFA (mol L⁻¹)	AHA (μmol L⁻¹)	LHA (mol L⁻¹)
La	7.5E-10	2.0E-11	3.9E-10
Ce	2.7E-10	5.7E-11	7.4E-10
Pr	3.4E-11	4.8E-12	7.3E-11
Nd	1.3E-10	1.9E-11	3.1E-10
Sm	2.2E-11	3.6E-12	4.5E-11
Eu	8.4E-12	1.0E-12	1.1E-11
Gd	1.8E-11	3.4E-12	5.2E-11
Tb	4.8E-12	5.0E-13	7.6E-12
Dy	9.7E-12	2.7E-12	4.6E-11
Ho	3.4E-12	5.2E-13	3.0E-12
Er	7.8E-12	1.4E-12	2.5E-11
Tm	2.4E-12	2.0E-13	<DL
Yb	9.1E-12	1.1E-12	2.0E-11
Lu	2.4E-12	1.7E-13	<DL