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1 **Uranium retention in a Callovo-Oxfordian clay rock formation:**
2 **From laboratory-based models to *in natura* conditions**

3
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12
13 **Keywords:** U, COx, adsorption, lability, nuclear waste

14
15 **Abstract**

16 For the performance assessment of radioactive waste disposal, it is critical to predict the
17 mobility of radionuclides in the geological barrier that hosts it. A key challenge consists of
18 assessing the transferability of current knowledge on the retention properties deduced from
19 model systems to *in natura* situations. The case of the redox-sensitive element uranium in the
20 Callovo-Oxfordian clay formation (COx) is presented herein. Extensive experimental work
21 was carried out with respect to parameters affecting uranium speciation (pH, P_{CO2}, [Ca] and

22 redox potential) with illite, COx clay fraction and raw COx claystone. The "bottom-up"
23 approach implemented, with illite and montmorillonite as reactive phases, quantitatively
24 explains the adsorption results of U(VI) and U(IV) on COx. While retention is high for U(IV)
25 ($R_d \sim 10^4 \text{ L}\cdot\text{kg}^{-1}$), it remains very low for U(VI) ($R_d \sim 4 \text{ L}\cdot\text{kg}^{-1}$) due to the formation of soluble
26 ternary Ca(Mg)-U(VI)-carbonate complexes. The applicability of the sorption model was then
27 assessed by comparing predictive analyses with data characterizing the behavior of naturally-
28 occurring U ($< 3 \text{ mg}\cdot\text{kg}^{-1}$). The COx clay phase is the largest reservoir of naturally-occurring
29 U ($\sim 65\%$) but only a small fraction appears to be adsorbed ($\sim 1\%$). Under representative site
30 conditions (especially with respect to reducing conditions), we have concluded that ternary
31 U(VI) complexes control U speciation in solution while U(IV) surface species dominate U
32 adsorption, with R_d values $> 70 \text{ L}\cdot\text{kg}^{-1}$.

33

34 **1. INTRODUCTION**

35 The disposal of radioactive waste in deep geological formations has been a topic of debate for
36 many decades. The rationale lies in isolating the waste packages within specifically
37 engineered facilities located deep underground (often 400 to 600 m) in geological formations
38 in order to ensure that the time for fractions of the confined radionuclide inventories to
39 potentially migrate to the human accessible environment remains much longer than their half
40 life so as to protect future generations from radiological and chemotoxic risks. The main types
41 of host rock formations used are sedimentary (e.g. clay rocks), crystalline (e.g. granite) or salt
42 domes (Altmann, 2008).

43 Several countries are investigating claystone formations (e.g. Opalinus claystone (OPA) in
44 Switzerland, Boda claystone (BODA) in Hungary, Boom clay (BC) in Belgium and Callovo-
45 Oxfordian claystone (COx) in France). The COx formation is located in the eastern part of the

46 Paris Basin (depth: ~400-600 m, thickness: ~120-150 m) and is characterized by low
47 permeability, mineralogical homogeneity and a low variation in other physicochemical
48 properties over a large expanse (ANDRA, 2005).

49 Uranium and its decay series nuclides are a significant component of radioactive waste. Given
50 a reducing environment measured in COx in the natural state ($E_h \sim -190$ mV), geochemical
51 modeling exercises applied to radioactive waste disposal are able to predict U(IV) as the
52 principal redox state in the formation when using a thermodynamic database, such as the
53 Nuclear Energy Agency's Thermochemical Database (NEA-TDB) published in 2003
54 (Guillaumont et al., 2003). U(IV) species can be considered as immobile due to both their
55 very high reactivity with claystone (sorption on clay mineral surfaces) and poor solubility in
56 pore water. However, studies have identified soluble ternary complexes Ca-U(VI)-carbonate
57 with strong formation constants (Bernhard et al., 1996; Dong and Brooks, 2006), thus shifting
58 the U(VI)/(IV) stability field boundaries to more negative redox potentials; these complexes
59 are expected to stabilize the U(VI) form under reducing conditions like those in COx
60 claystone. Should the ternary aqueous complexes be stable and dominant, then uranium would
61 become more soluble and mobile, even if U(IV) remains dominant in the solid phase. This
62 outcome has recently been shown for OPA by means of a detailed modeling effort using one-
63 dimensional diffusion models (Hennig et al., 2020).

64 For this reason, special care must be taken when dealing with uranium, since it is considered
65 to be potentially the most mobile of the long-lived actinides in the performance assessment.

66 To date, just one work has been devoted to U(VI) retention on COx (Hartmann et al., 2008).
67 These authors showed the importance of ternary complexes in describing the data, but their
68 model was unable to satisfactorily reproduce the experimental results. Furthermore, all studies
69 on claystones (COx, BODA or OPA) focus on the retention of U(VI) (Amayri et al., 2016;

70 Bradbury and Baeyens, 2011; Hartmann et al., 2008; Joseph et al., 2011; Kautenburger et al.,
71 2019; Marques Fernandes et al., 2015), although to the best of our knowledge no experimental
72 data on U(IV) actually exist. Along the same lines, only the U(VI) retention model has been
73 taken into account to simulate the diffusion of U in the OPA formation (Hennig et al., 2020).
74 Given the highly reducing environment of deep formation, a major role played by the
75 tetravalent state of U on uranium retention cannot be excluded.

76 Given the limited number of published articles, lack of data (notably with respect to U(IV))
77 and importance of the topic, the objective of this work is to promote a better understanding of
78 uranium retention in the Callovo-Oxfordian formation. The term retention, as used herein,
79 means the adsorption of U on the clay fraction governed by kinetically fast and reversible
80 adsorption processes (ion-exchange, surface complexation) (OECD Nuclear Energy Agency,
81 2002). Throughout this paper, the term “labile fraction” will be used in order to characterize
82 the adsorbed U fraction in “equilibrium” with the water-soluble part over a short time scale,
83 whereby the formation can be described by local equilibrium approaches. The significance of
84 the clay fraction for metal adsorption by COx has already been shown for Cs (Chen et al.,
85 2014b), Ni (Chen et al., 2014a; Grangeon et al., 2015; Montavon et al., 2020) and Eu (Loni et
86 al., 2021). Such is also the case for Opalinus Clay, whose mineralogy is quite close (Bradbury
87 and Baeyens, 2011; Marques Fernandes et al., 2015).

88 In this exercise, extensive experimental work has been carried out under well-controlled
89 conditions with illite, the COx clay fraction and raw COx by studying the parameters
90 affecting uranium speciation (pH, P_{CO_2} , presence of divalent cations (Ca, Mg), redox
91 potential). The available 2-Site Protolysis, Non-Electrostatic Surface Complexation and
92 Cation Exchange model (2 SPNE SC/CE), proposed by Bradbury and Baeyens (1997) and
93 developed for OPA, has been used as a basis for describing the data in relying on the
94 simplifications proposed in the literature for COx (Chen et al., 2014a; Grangeon et al., 2015;

95 Tournassat et al., 2009). Special attention has been paid to the U(VI)/illite/carbonate system in
96 order to better assess the role of ternary surface complexes, $\equiv\text{S-U(VI)-carbonates}$, in light of
97 the absence of specific data for illite in the published models (Bradbury and Baeyens, 2017).
98 These data then make it possible to develop an operational model that allows predicting the
99 adsorption properties of uranium under *in natura* CO_x conditions by pursuing the "bottom-
100 up" approach with very good knowledge of the CO_x environment (Gaucher et al., 2009;
101 Lerouge et al., 2011).

102 Inspired by previous work focusing on uranium-contaminated areas (Curtis et al., 2004;
103 Kohler et al., 2004; Payne et al., 2001), these predictions were subsequently compared with
104 results describing the behavior of uranium naturally present in the formation at trace
105 concentrations. In addition to the analysis of U distribution in the various CO_x reservoirs,
106 emphasis has been placed on the labile part of this element, i.e. a fraction difficult to
107 characterize under conditions of trace concentrations, when spectroscopic tools are not
108 applicable (McKinley and Alexander, 1993). This fraction is needed in order to compare the
109 data with results obtained in the laboratory; it has been estimated by means of sequential
110 extraction (Claret et al., 2010), isotope exchange (Ahmed, 2014) and by following a
111 desorption method successfully tested on the Ni/CO_x system (Montavon et al., 2020).

112 In sum, the challenge of this work lies in linking the model system to the natural system and
113 drawing a conclusion on the degree to which uranium remains mobile in the CO_x formation;
114 the steps involved must identify which physicochemical parameters control uranium mobility
115 from a mechanistic point of view.

116 117 **2. EXPERIMENTAL DETAILS**

118 **2.1 Materials**

119 *2.1.1 Chemicals*

120 All solutions were prepared with ultra-pure deionized water (18.2 MΩ.cm) and commercially
121 available chemical products of analytical or ultra-trace metal grade. For HR-ICP-MS
122 measurements, nitric acid was purified using a sub-boiling distillation system (Savillex DST-
123 1000). U(VI) stock solution was prepared from ICP standard solution (1,000 µg/mL in 4%
124 HNO₃ or 10,000 µg/mL in 0.4% HNO₃) (SCP Science, France). The solution of ²³³U was
125 provided in 2N HNO₃ at an activity of 231 Bq (0.65 µg) of ²³³U per gram of solution
126 (Amersham). ²³²U (841 kBq.g⁻¹ in 2N HNO₃ - CERCA) was also used as a tracer for some of
127 the experiments performed under reducing conditions.

128 *2.1.2 CO_x samples*

129 All CO_x samples used in the present study are core samples from boreholes within the studied
130 area (Meuse/Haute-Marne, France). The claystone consists (in percentage by weight) of a
131 major clay fraction (40-60%), associated with both quartz (25-37%) and carbonate (12-32%)
132 fractions. The clay minerals are mainly illite (ILL) (14-29%) and ordered illite-smectite mixed
133 layers (IS) (18-33%), with minor kaolinite and chlorite contents (Lerouge et al., 2011).

134 This work has benefited from a collection of some 50 samples extracted from various units
135 and characterized in terms of composition and trace elements (Lerouge et al., 2006). Other
136 samples were selected for the experimental work with two objectives, namely: (i) evaluation
137 of the U distribution in the CO_x in order to identify the U-carrier phases by correlating the
138 measured concentrations with sample characteristics; and (ii) execution of the laboratory
139 experimental campaign. Detailed information on these samples is given in Table A-1. Except
140 for sample EST21400, which is rich in calcite, the samples were all extracted in a unit with
141 maximum clay content (40%-53%). After drilling, they were conditioned in airtight aluminum
142 foil filled with nitrogen, transported to the laboratory and then stored until their opening in a
143 N₂/CO₂ (1%) glove box.

144 The separation and purification of the COx clay fraction (COX-cf) (from EST51779) was
145 performed by elutriation and chemical purification, as provided by the BRGM Institute
146 (Claret et al., 2004). The sample labeled EST25687(ox) corresponds to a portion of the
147 previously preserved EST25687 sample analyzed in 2006 and not preserved since then.
148 Analyses conducted on both oxidized and non-oxidized samples allow assessing the potential
149 presence of reduced forms of uranium. Furthermore, sample EST26536 was selected in the
150 maximum clay zone for its phosphate nodules.

151 Water samples from three drillings, cored with inert gases (N₂ or Ar), were analyzed in order
152 to define the representative U concentration values in the pore water of the Callovo-Oxfordian
153 (PAC1002, POX1201, EPT1201) (Vinsot et al., 2013). Data were also collected from
154 GIS1002 (Grangeon et al., 2015); although initially drilled under unpreserved conditions, a
155 state of return to *in natura* equilibrium can now be observed. The effect of an oxidative
156 disturbance on the U pore water concentration has been evaluated from borehole KEY1001
157 (Grangeon et al., 2015).

158 159 2.1.3 Aqueous media

160 Synthetic pore water (SPW1) representative of the COx formation, and in equilibrium with
161 calcite, was prepared under controlled anoxic Ar/CO₂ conditions ($P_{\text{CO}_2} = 10^{-2}$ atm or 1%) for
162 the sorption experiments with illite, COx-cf and COx and for the purpose of evaluating the
163 labile part of naturally-occurring U (Table A-2).

164 These experiments were also carried out in weakly complexing media (NaNO₃ or NaCl) with
165 an ionic strength (0.1 M) close to that encountered *in natura*. Various atmospheres were
166 selected (Ar, air/CO₂ at atmospheric pressure ($10^{-3.5}$ atm), 1% CO₂ (10^{-2} atm) and 5% CO₂
167 ($10^{-1.3}$ atm)) in order to evaluate the impact of carbonates on the retention of uranium onto

168 illite. Equilibrium with P_{CO_2} was verified qualitatively, after at least one day of equilibration
169 while the pH was stable, and quantitatively via an alkalinity measurement.

170 The effect of redox potential was also evaluated under inert atmosphere (Ar) and in the
171 presence of carbonates (3,500 ppm or 9,500 ppm CO_2) in synthetic groundwater SPW2 (Table
172 A-2). In order to promote the conditions to have U(IV), a reducing agent (Na_2S) was added to
173 the system.

174

175 **2.2 Batch-type experiments**

176 A summary of the experimental campaign is given in Table 1 and detailed experimental
177 conditions are provided in the appendices (Tables A-3 and A-4).

178 All experiments were performed at room temperature ($T = 22^\circ \pm 3^\circ C$) in polypropylene
179 copolymer (PPCO) tubes either inside or outside the glove box, depending on the desired
180 atmosphere. Moreover, they were conducted under conditions whereby uranium did not
181 precipitate; these conditions were first deduced from simulations (see the "Modeling" section
182 below, 2.4) before being verified for an absence of precipitation by filtration using 0.45- μm
183 PolyTetraFluoroEthylene (PTFE) filters.

184 The data on adsorption experiments were acquired using the classic batch-type technique
185 (Montavon et al., 2006). The solid phase was equilibrated with the liquid phase before adding
186 U to the system. Note that for the experiments with Na_2S , the reducing agent was added both
187 to the U stock solution and to the suspension. In the end, the Eh value measured under
188 equilibrium conditions was taken into account for the data analysis. Preliminary sorption
189 experiments showed that a contact time of 3 days was sufficient to reach sorption equilibrium
190 conditions (data not shown). The solid and liquid phases were separated by means of filtration
191 using 0.2- μm PTFE or Surfactant-free Cellulose Acetate (SFCA) filters. Except for the

192 experiments carried out under inert conditions (Ar), no sorption at the surface of tubes or on
 193 filters could be detected (< 5%). When such was not the case (experiments under Ar
 194 atmosphere), a correction was introduced based on a "blank" analysis (i.e. under the same
 195 conditions but in the absence of the sorbent phase). The pH values of the solutions and
 196 alkalinity were measured at the end of the experiments. This step also applied to experiments
 197 carried out under Ar atmosphere, wherein carbonate traces can affect the speciation of U
 198 (Tournassat et al., 2018). The distribution coefficient between solid and liquid phases, R_d (in
 199 $L \cdot kg^{-1}$), was defined as follows:

$$200 \quad R_d = \frac{[U]_0 - [U]_{aq}}{[U]_{aq}} \frac{V}{m} \quad (1)$$

201 where $[U]_0$, $[U]_{aq}$, m and V denote the U concentration introduced in the system ($mol \cdot L^{-1}$), the
 202 U concentration measured in solution at equilibrium ($mol \cdot L^{-1}$), the solid mass (kg) and the
 203 solution volume (L), respectively.

204 This same batch-type method was followed for experiments carried out with naturally-
 205 occurring U in order to quantify the labile fraction, in considering an isotopic exchange
 206 approach (Ahmed, 2014) and a desorption study (Montavon et al., 2020). The equilibration
 207 criteria will be presented in the "Results" section. For the desorption experiments, if U in the
 208 aqueous phase is controlled by an adsorption equilibrium and given the assumption that the
 209 R_d model is applicable, the following relation is obtained for correlating the U concentration
 210 in the aqueous solution (in $\mu g \cdot L^{-1}$) with the solid-to-liquid ratio m/V (in $g \cdot L^{-1}$):

$$211 \quad [U]_{aq} = \frac{n \cdot \frac{m}{V} + [U]_i}{(1 + R_d \cdot \frac{m}{V})} \quad (2)$$

212 where $[U]_i$ represents the potential contamination of U initially present in SPW (in $\mu g \cdot L^{-1}$),
 213 and n the quantity of mobilizable element ($\mu g \cdot g^{-1}$), i.e. the labile quantity associated with COx

214 plus the quantity initially present in pore water and present on the raw solid (in $\mu\text{g}\cdot\text{g}^{-1}$). The
215 Rd unit of measurement used in this expression is $\text{L}\cdot\text{g}^{-1}$.

216

217 **2.3 Analytical procedures**

218 **2.3.1 Aqueous solutions**

219 Uranium was measured in aqueous solutions by ICP-MS (Thermo Xseries 2, Thermo
220 Scientific) when concentrations were above $2\ \mu\text{g}/\text{L}$ and HR-ICP-MS for U concentrations at
221 ultra-trace level (Element XR, Thermo Scientific). Quantitative analyses of uranium were
222 performed by external calibration with a set of gravimetric uranium standard solutions
223 prepared using an internal standard solution of thallium in 2% (v/v) HNO_3 for correction of
224 matrix effects and instrumental drift.

225 ^{232}U analyses were performed by liquid scintillation counting with a Packard TriCarb 3170
226 TR using an Ultima Gold LLT[®] (Perkin Elmer) scintillation cocktail. $^{232}\text{U} / ^{228}\text{Th}$ separation
227 was carried out prior to analysis using the anionic exchanger Dowex 50WX8-200 with 8M
228 HCl as elution medium for thorium while uranium remains on the resin. Then ^{232}U was
229 ultimately recovered with water.

230 Alkalinity was determined by colorimetric analysis using Bromocresol Green-Methyl Red
231 pH- indicator powder pillow and a Digital Titrator (Hach method 8203) with digital titrator
232 cartridges containing 0.16 N (or 0.02 N for a low concentration of carbonates in the system)
233 of H_2SO_4 .

234 pH values were measured by a pH-meter combined with a pH electrode and thermometer
235 detector (pHC2085, Hach Lange) of hydrogen ion activity. The pH electrode was composed
236 of glass combined with a red rod reference electrode of silver/silver chloride. The pH-meter
237 was automatically calibrated by PHM220 Lab pHmeter (Radiometer Analytical) at the

238 beginning of each experiment using two standard IUPAC buffer solutions of pH 4.005 and
239 7.000 (Radiometer Analytical).

240 The redox potential was measured by a pH-meter combined with a redox electrode (MC3051-
241 PT-9, Radiometer Analytical). This electrode had an Ag/AgCl reference system with a
242 refillable electrolyte (3 M KCl saturated with AgCl), porous pin junction and platinum ring
243 sensor. The electrode was calibrated with a redox standard solution; its values were given
244 with respect to the normal hydrogen electrode (NHE).

245

246 **2.3.2 CO_x phases**

247 The sequential extraction protocol adapted from Claret et al. (2010) was first used to assess
248 the U distribution in the CO_x clay. The extraction steps comprised the following:

249 Fraction 1 (F1 - water soluble fraction): The extraction was performed in the glove box (Ar
250 atmosphere), where 20 mL of deionized water were added to 4 g of solid sample and shaken
251 for 24 h. The resulting solution was then filtered using 0.22- μ m Surfactant-Free Cellulose
252 Acetate filters after centrifugation at 20,000 rpm for 10 minutes.

253 Fraction 2 (F2 – part of the adsorbed fraction): In this step, as well as for steps 3 through 5,
254 the extractions took place outside the glove box. 20 mL of hexamine trichloride (4.458 g·L⁻¹)
255 were mixed with the residue from the previous step and shaken for 30 minutes. After
256 centrifugation and filtration, the solid obtained was used for step 3.

257 Fraction 3 (F3 – part of the adsorbed fraction and U present in carbonate phases): 20 mL of an
258 acetate/acetic acid buffer at pH 5 were added to the residue from the previous step and then
259 shaken for 12 h. After centrifugation and filtration, the solid fraction was used for the next
260 step.

261 Fraction 4 (F4 - organic matter fraction): 20 mL of H₂O₂ at pH 2 (adjusted using HNO₃) were
262 added to the residue of the previous extraction and heated at 50°C for 3 h. After centrifugation
263 and filtration, the solid fraction was used for the final step.

264 Fraction 5 (F5 - residual fraction): In this last step, the sample was digested by a sequence of:
265 dissolution in a mixture of HNO₃ (2 mL), HF (10 mL) and HClO₄ (5 drops) at 120°C for one
266 week followed by the evaporation of HF, and then the addition of HCl 6N (10 mL) at 100°C
267 for one week.

268 The same digestion procedure was also carried out on CO_x samples of interest for the
269 determination of total concentration of U naturally present. It should be noted that a digestion
270 procedure was also conducted to dissolve the organic matter separated by handpicking from the
271 EST26536 sample under the binocular lens; to accomplish this, 20 mL of an H₂O₂ solution at
272 pH 2 (adjusted using HNO₃) were used at 50°C for 3 hours. Also note the absence of any
273 visible residue after extraction indicating the purity of the separated organic matter.

274 U solid analyses were performed by laser ablation HR-ICP-MS. Details of the instrumental
275 facility, optimization strategy of laser and HR-ICP-MS instrument, data acquisition and off-
276 line processing are reported elsewhere (Loni et al., 2021; Montavon et al., 2020). The certified
277 value of U concentration at 37.38 mg.kg⁻¹ of the NIST 612 standard reference material was
278 used for uranium quantification in CO_x mineral phases by external calibration and internal
279 standardization (Loni et al., 2021; Montavon et al., 2020). The total analysis time for laser
280 ablation sample and gas blank measurements is reported in Table S1. The sample had
281 previously been mounted on a glass blade with an epoxy-type glue, and its surface polished
282 by silicon carbide grinding paper (Struers) (grading size: from 80 to 4,000 Grit).

283
284 **2.4 Modeling considerations**

285 All calculations were carried out with the PhreeqC code (version 3.2.1) (Parkhurst and
286 Appelo, 2013). The Davies equation (Davies, 1962) was used for the ionic strength correction
287 of solutes with the corresponding THERMOCHEM v10a (2018) database (Grivé et al.,
288 2015), www.thermochimie.org . The stability constants of aqueous uranium species were
289 extracted from the NEA Thermodynamic Database (Grenthe et al., 2020), except for
290 $\text{CaUO}_2(\text{CO}_3)_3^{2-}$ and $\text{Ca}_2\text{UO}_2(\text{CO}_3)_3$ species, which were drawn from (Maia et al., 2021). . All
291 relevant data can be found in the supplementary data section of this article (Table S2).

292 For the adsorption model, it was not sought to build a new type of model but rather to use
293 existing models and test/adapt them to the CO_x conditions. We chose the 2SPNE SC/CE
294 model (Bradbury and Baeyens, 1997) since it is the most complete among those available in
295 the literature and had already been satisfactorily tested for both Ni/CO_x (Chen et al., 2014a;
296 Grangeon et al., 2015; Montavon et al., 2020) and Eu/CO_x (Loni et al., 2021) systems.
297 According to the 2SPNE SC/CE model, the surface complexation phenomenon is controlled
298 by both weak (i.e. high quantity, limited affinity) and strong (low quantity, high affinity) sites.
299 Previously published simplifications and assumptions (Chen et al., 2014a; Grangeon et al.,
300 2015; Montavon et al., 2020; Tournassat et al., 2009) are considered herein, namely: (i) 2:1
301 clay minerals control the retention, and the potential contributions of chlorite and kaolinite are
302 not taken into account in the overall sorption of uranium; (ii) the I/S interstratified layers are
303 assumed here as two phases, where each phase corresponds to 50% wt. of illite and smectite,
304 respectively; and (iii) montmorillonite is used as a proxy for smectite phases since it is an
305 example of a well-studied smectite.

306 For the quantitative description of these adsorption experiments, the mineralogical
307 compositions listed in Table A1 have been taken into account.

308 The adsorption parameters for both illite and montmorillonite are summarized in Table A5.
309 Values for the U(VI) surface complexation reactions were taken from (Baeyens and Bradbury,
310 2017) for montmorillonite and from (Bradbury and Baeyens, 2017) for illite. Values for the
311 U(IV) surface complexation reactions were calculated from the Linear Free Energy
312 Relationship equations (Baeyens and Bradbury, 2017; Bradbury and Baeyens, 2017), as
313 established from sorption data with Th(IV).

314 For the various modeling predictions, the average clay concentrations encountered in the
315 formation (see previous section) have been used. As regards model sensitivity, unless
316 otherwise stated, calculations were performed with the lower and upper clay content limits.

317

318 **3. RESULTS**

319 Based on literature data (Chen et al., 2014a; Grangeon et al., 2015; Montavon et al., 2020;
320 Tournassat et al., 2009), it has been assumed that the COx clay fraction controls uranium
321 adsorption. The adsorption mechanism, including both surface complexation and cation
322 exchange, leads to the formation of so-called "labile" chemical forms, i.e. in rapid equilibrium
323 with the fraction in solution. Other retention mechanisms such as incorporation or co-
324 precipitation, which generally lead to so-called "inert" forms, were therefore neglected in our
325 approach. The experimental and simulation approaches were oriented along these lines. The
326 relevance of these assumptions, in particular when linking the model system to the natural
327 system, will be discussed in Section 4.

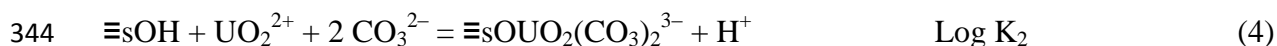
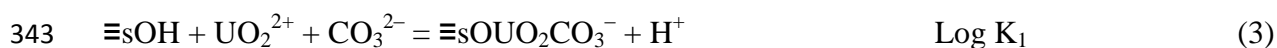
328 **3.1. Which model for describing U retention on COx?**

329 *3.1.1. Conditions where U(VI) species dominate*

330 The experimental data obtained for illite under various experimental conditions are given in
331 Figure 1. Maximum sorption was reached at a pH range between about 6.5 and 7.5. Under

332 conditions with a low carbonate level and simple background electrolyte, a retention value of
 333 $(4 \pm 0.3) 10^5 \text{ L}\cdot\text{kg}^{-1}$ was calculated. With an increasing carbonate level (i.e. atmospheric CO_2
 334 and 1% CO_2), a decrease in the R_d value was observed ($R_d \sim (3 \pm 0.5) 10^4 \text{ L}\cdot\text{kg}^{-1}$ and
 335 $\sim (2 \pm 0.4) 10^3 \text{ L}\cdot\text{kg}^{-1}$, respectively). In SPW containing divalent cations and a 1% CO_2
 336 atmosphere, this decrease was even greater, with R_d values of approx. $45 \pm 9 \text{ L}\cdot\text{kg}^{-1}$.

337 This trend has previously been observed for the retention of U(VI) on smectite in the presence
 338 of CO_2 (Marques Fernandes et al., 2012). If the formation of carbonate complexes (and in
 339 particular ternary complexes with divalent cations) lies at the origin of the retention decrease,
 340 then the data cannot be modeled when ignoring the sorption of uranyl-carbonate complexes.
 341 Similarly to the study by (Marques Fernandes et al., 2012) the formation of two surface
 342 complexes for illite are considered (Table A5):



345 These constants have been (manually) adjusted to obtain a satisfactory level of agreement
 346 between the model and the experiment.

347 The model was then applied predictively to the COx clay fraction, under various experimental
 348 conditions, as well as to the COx clay rock in the presence of synthetic water (Fig. 2). For the
 349 clay fraction, in a simple background electrolyte, R_d values of $(8.0 \pm 0.8) 10^3 \text{ L}\cdot\text{kg}^{-1}$ and
 350 $(2.0 \pm 0.2) 10^3 \text{ L}\cdot\text{kg}^{-1}$ were obtained in close to atmospheric CO_2 pressure and 1% CO_2 ,
 351 respectively. In the presence of SPW, R_d decreased to about $50 \pm 25 \text{ L}\cdot\text{kg}^{-1}$. For the COx clay
 352 rock experiment, the retention coefficient decreased to low values, i.e. approx.
 353 $3.6 \pm 1.5 \text{ L}\cdot\text{kg}^{-1}$.

354 In sum, the retention coefficient varies in the following order: illite ~ CO_x-cf > CO_x, which
355 qualitatively shows that the clay fraction of CO_x is, in terms of reactivity, close to illite. These
356 data are on the whole well reproduced by the model, including the two new surface
357 complexation reactions (lines in Fig. 2).

358 The model was ultimately tested with data published for U/clayey rock systems (Fig. A1,
359 Table S3); the output provides a satisfactory explanation of the experimental data, within the
360 uncertainty, for BODA and OPA, as do the modeling approaches proposed in the referenced
361 papers. Despite producing better results, the experiment / calculation agreement was still not
362 perfect for the data held by Hartmann et al. (2008) (Fig. A1c). Note that in their work, the
363 sorption of uranyl on CO_x claystone was studied at 0.1 M NaClO₄ in atmospheric CO₂
364 pressure as a function of pH and with an assumption of calcite equilibrium.

365 *3.1.2 Impact of reducing conditions*

366 Given the highly reducing environment for the deep formation of CO_x, a significant role
367 played by the tetravalent state of U on uranium retention cannot be excluded. This hypothesis
368 has therefore been tested from batch-type experiments with the EST05640 sample. The
369 corresponding experimental conditions are shown in the appendices (Table A4).

370 The experimental data are given in Figure 3, along with the model predictions. As expected,
371 retention was strong with Rd values on the order of 10⁴ L·kg⁻¹. Although the experimental
372 data were rather scattered, the agreement between experiment and calculation appears to be
373 relatively good, with no parameter adjustment being required. Modeling results indicate that
374 U(IV) dominates the surface speciation for all studied conditions (> 99.6%). It is also the
375 major redox state in solution for the conditions presented in Figures 3a and 3b (99% and 90%,
376 respectively) but not in the presence of 9,500 ppm CO₂, where it corresponds to only 10%
377 (Fig. 3c).

378
379

3.2. Distribution of naturally-occurring U in COx

380 In the COx formation, uranium contents are low, i.e. between 0.5 and 2.4 mg·kg⁻¹ (Fig. 4a).
381 Among the various samples whose U amount has been analyzed, a distinct correlation exists
382 with the organic matter: the higher the organic matter (OM) concentration, the greater the
383 amount of U (Fig. 4a). For the samples used in laboratory studies, the upper concentration
384 limit is reached, i.e. between 2 and 3 mg·kg⁻¹.

385 Among the various phases, uranium is expected to be incorporated in minor phases, e.g.
386 phosphates (Regenspurg et al., 2010). Calcite might be another possible reservoir due to its
387 high reactivity and ability to trap metals (Zhao and Zheng, 2014). These minerals were
388 therefore singled out in the present study, which also considered organic matter (in relation to
389 the correlation presented above) as well as the clay fraction, given its predominant role in the
390 metal adsorption process. Analysis details can be found in Table S1.

391 The analyses of individual mineral phases carried out on sample EST51769 yielded U
392 concentration values of 2.7 ± 0.8 , 1.0 ± 0.3 and 0.013 ± 0.004 mg·kg⁻¹ for clay, calcite and
393 pyrite fractions, respectively. Similar results were observed for biocalcite and diagenetic
394 calcite on the EST05738 and EST26536 samples; the values were 0.4 ± 0.2 and 0.5 ± 0.3 mg
395 kg⁻¹, respectively. Four calcium phosphate analyses (EST26536) were performed in line mode
396 and the average uranium content equaled 32 ± 4 mg·kg⁻¹. Lastly, the uranium concentration
397 measured in organic matter led to a U content of 69.2 µg·kg⁻¹.

398 Considering the average composition of COx in the clay-rich zone and the U concentration
399 measured for EST51769 (whose main values were obtained), the distribution diagram shown
400 in Figure 4c is derived (due to the low levels measured, U concentrations in pyrite and organic
401 matter are not reported).

402 The U concentration of the bulk rock (called total (ind.) in Fig. 4c), as recalculated from the U
403 concentrations measured in the various phases and their proportion in the CO_x, corresponds
404 (within the range of uncertainties) to the U concentration measured directly in the bulk rock
405 (called total in Fig. 4c). This finding indicates that the major carrier phases have been
406 identified. While the highest U concentrations were found in phosphate minerals, as expected,
407 the clay fraction actually corresponds to the greatest reservoir with ~65% of the uranium
408 present. On the other hand, Figure 4a serves as an "optical illusion" since the organic matter,
409 due to its low quantity in the CO_x and low U concentration measured in this phase, is clearly
410 not a U-bearing phase. The origin of this visual correlation remains unclear and has not been
411 further investigated herein, given the objectives of this work.

412 The uranium distribution was then indirectly evaluated by means of sequential extraction;
413 results are shown in Figure 4b. Distributions are similar for all clay-rich samples (~3 mg·kg⁻¹
414 in U). The U in fractions F1 (water soluble fraction), F2 (part of the adsorbed fraction) and F4
415 (organic matter fraction) accounts for less than 5% of total U. This result is consistent with
416 previous data; U in the organic fraction (F4) can be neglected. Therefore, U is mainly
417 distributed between fractions F3 (part of the adsorbed fraction and U present in carbonate
418 phases) and F5 (residual fraction), with a relatively constant distribution of ~12%:86% F3:F5.
419 Moreover, no significant effect was observed relative to the state of conservation of the CO_x.

420 At this stage, it is interesting to compare these sequential extraction results with analyses
421 conducted directly on solids. The uranium fractions present in carbonate phases and on clays
422 (assuming that U is adsorbed) should mostly be found in F3 (the adsorbed part of U found in
423 F2 being considered negligible). If we consider that the uranium in phosphates is trapped in
424 the matrix, a proportion between fractions F3 and F5 on the order of 90%:10% is thus
425 obtained from the direct solid analyses presented in Fig. 4c. While this analysis is consistent
426 with sequential extraction results from a qualitative point of view, such is not the case from a

427 quantitative point of view; in fact, the inverse trend is observed, i.e. ~12%:86% F3:F5. Since
428 the carbonate phases are completely dissolved after step 3, a sizable fraction of the U directly
429 detected in the clay phase is present in a “non-labile” form, i.e. not desorbed with the pH
430 decrease and thus recovered in F5. The U content measured in the F3 fraction therefore stems
431 mainly from the dissolution of carbonate phases.

432 This result is consistent with the fact that the F3 fraction increases with the amount of
433 carbonate phases, that is ~35% and ~12 % for EST21400 (~81% of carbonate phases) and
434 EST26536, EST25687 and EST26480 (~25% of carbonate phases), respectively. From a
435 quantitative point of view, assuming that the uranium in the clay fraction is completely found
436 in F5, in knowing the carbonates phases content of the CO_x samples (Table 1), a U content of
437 ~1.3 mg·kg⁻¹ and ~0.9 mg·kg⁻¹ in the carbonate phases can be recalculated from the
438 sequential extraction results (F3) for samples rich in clay and carbonate phases, respectively.
439 The order of magnitude is consistent with that found from direct analysis of calcite (the
440 majority phase of carbonate phases), i.e. ~0.4-1 mg·kg⁻¹.

441 In conclusion, if uranium is mainly present in the CO_x clay fraction, it is not retained (for a
442 large part) in a form expected by our model, namely retained by a surface complexation
443 mechanism.

444
445 **3.3. Retention coefficient and labile fraction of naturally-occurring U under both**
446 **oxidizing and reducing conditions**

447 Experiments were conducted with CO_x powder in the presence of SPW1 under Ar/CO₂
448 atmosphere.

449 Results of the isotope exchange method are given in Figure 5a. A fraction of the naturally-
450 occurring U was rapidly resupplied from the clay rock samples in solution, with steady state

451 achieved after less than 1 day of contact. Regarding the samples doped with ^{233}U , about 3
452 days were required to reach steady state. We consider that this contact time is insufficient to
453 allow for the exchange of U-233 with U-238 incorporated into the mineral matrix; since this
454 process is controlled by solid diffusion, the kinetics are expected to be very slow. An analysis
455 of the data yields an R_d value of $7.6 \pm 2.3 \text{ L}\cdot\text{kg}^{-1}$ and an amount of labile U present in the
456 COx of $26 \pm 7 \mu\text{g}\cdot\text{kg}^{-1}$.

457 For the desorption method, like for the isotope exchange method, contact time is a critical
458 parameter to be assigned. While the time to reach steady state is quick at $m/V=50 \text{ g}\cdot\text{L}^{-1}$, it still
459 might vary with respect to the solid-to-liquid ratio. Two other kinetic experiments were
460 therefore performed for lower solid-to-liquid ratios (i.e. 0.5 and $1 \text{ g}\cdot\text{L}^{-1}$). The equilibration
461 time was longer (i.e. 4 days instead of 1 day) (Fig. 6b), and a contact time of 7 days was thus
462 set for all experiments.

463 The desorption results obtained as a function of the solid-to-liquid ratio are provided in Figure
464 5c. Like with the sequential extraction results, no difference has been found between the
465 preserved (black symbols) and unpreserved (white symbols) COx samples. The experimental
466 data can be well explained in considering our simple model (Eq. 2) with R_d and n values at
467 $8.0 \pm 4.6 \text{ L}\cdot\text{kg}^{-1}$ and $30 \pm 12 \mu\text{g}\cdot\text{kg}^{-1}$, respectively (solid lines in Fig. 5c). Note that the
468 uranium content present in SPW1 as well as the amount of U in COx originating from pore
469 water are both negligible quantities. The good agreement of results between the two
470 equilibrium-based methods have given us confidence in the results. It should also be noted
471 that although the isotope exchange method does make it possible to return to a value for the
472 distribution of uranium (VI) between solid and liquid phases (as translated by an R_d value),
473 the desorption method, owing to the curve shape, exposes this as an adsorption phenomenon.
474 If in fact it was controlled by a solubility mechanism, then the U concentration would not
475 evolve with the mass-to-volume ratio (under saturated conditions).

476 A final experiment was carried out in the laboratory to assess the relevance of redox potential.
477 A suspension at $50 \text{ g}\cdot\text{L}^{-1}$ of COx (leading to a measured U concentration of $0.99 \text{ }\mu\text{g}\cdot\text{L}^{-1}$ for
478 $E_h \sim 0 \text{ mV} / \text{NHE}$) remained in the glove box (Ar/CO_2) for nearly 3 years. The measured
479 potential dropped back to the characteristic value of the site (i.e. $E_h \sim -200 \text{ mV}$) and the U
480 concentration decreased by a factor of nearly 10 (gray symbols in Fig. 5c). By considering the
481 labile fraction of U as a fixed parameter, this implies that U speciation at the surface has
482 changed, which results in an increase in the Rd value from 8.0 ± 4.6 to $500 \pm 200 \text{ L}\cdot\text{kg}^{-1}$.

483

484 **4. DISCUSSION**

485 An operational model, based on both a bottom-up approach and the 2SPNE SC/CE model, has
486 made it possible to quantitatively describe uranium sorption onto illite, the clay fraction of
487 COx and COx clay rock for a wide variety of conditions sensitive to U specification (pH, E_h ,
488 pCO_2 , concentration of divalent cations (Ca^{2+} , Mg^{2+})). Compared to the available adsorption
489 database, two parameters had to be fitted, namely the constants describing formation of the
490 ternary surface complexes with illite and U(VI) (Eqs. (3) and (4)). Note that the existence of
491 such carbonato surface complexes has recently been questioned in describing the U(VI) /
492 montmorillonite system (Tournassat et al., 2018). The formation of secondary sites
493 $\equiv\text{sOH}_2\text{CO}_3^-$ and $\equiv\text{sOH}_2\text{HCO}_3$ able to assist modifying metal retention (Rieder, 2019) can also
494 be envisaged. Given the objective of the present work, this aspect has not been further
495 explored.

496 Thanks to the operational model, it was possible to analyze the results of equilibrium-based
497 methodologies describing the behavior of the naturally-occurring U labile fraction for a
498 COx/SPW1 system representative of the natural environment. In terms of “oxidizing”

499 conditions, this labile fraction is characterized by an R_d value of $\sim 8 \text{ L.kg}^{-1}$. According to the
500 model, this finding reflects the presence of U(VI) adsorbed onto the clay surface. Such a
501 result validates the hypothesis of the clay fraction controlling the U adsorption process. The
502 labile fraction however remains low compared to that measured directly by LA-HR-ICP-MS
503 in the clay phases ($\sim 1\%$ of total U content). This result confirms our previous hypothesis that
504 a significant fraction of uranium found in the COx clay fraction is in fact incorporated into the
505 matrix in an “inert” form, as found in the residual fraction of the sequential extraction
506 experiments.

507 In conclusion, we can consider a labile fraction of naturally-occurring U in the formation on
508 the order of $30 \pm 12 \mu\text{g.kg}^{-1}$, to be associated with the clay fraction in an adsorbed form.

509 At this stage, the purpose of this work is to correlate the results obtained with the information
510 available at the geological formation level, i.e. the concentrations of U in the pore water.

511 Since no uranium phase has been observed and the labile fraction of naturally-occurring U
512 was found to be adsorbed onto the clay fraction (in the form of U(VI)), we can forward the
513 assumption that *in natura* pore water concentration is also controlled by an adsorption
514 phenomenon. Based on this assumption, the range of *in natura* concentrations in pore water
515 can be predicted from Eq. (2), in knowing: the labile fraction of U, the R_d value (as
516 characterized by the presence of U(VI)), and the high solid-to-liquid ratio characterizing the
517 site ($\sim 8\%$ wt. of water, i.e. $\sim 11,500 \text{ g.L}^{-1}$). A range of aqueous U concentrations between 1.4
518 and $12 \mu\text{g.L}^{-1}$ can thus be calculated (solid lines, Fig. 5c).

519 It is now important to compare this concentration range, deduced from the laboratory
520 experiments, with that measured directly in the pore water collected at the various boreholes
521 (see Section 2.1.2). A value of $0.13 \mu\text{g.L}^{-1}$ has been selected by ANDRA; this value lies close
522 to that measured in this study for the GIS1002 borehole, i.e. $0.25 \pm 0.02 \mu\text{g.L}^{-1}$. It also lies

523 close to the value measured under the undisturbed conditions of the DIR experiment (0.24
524 $\mu\text{g.L}^{-1}$) (Fralova, 2020). The results are therefore very consistent, and a concentration range of
525 0.13-0.25 $\mu\text{g.L}^{-1}$ can be given as representative values of the COx formation.

526 This range appears to be more than 10 times lower than the values deduced from Eq. (2); this
527 information immediately suggests that laboratory experiments are not representative of the
528 natural system.

529 The uncontrolled parameters in the laboratory are the COx state and redox potential. Three
530 articles focus on the effect of compaction for metal retention parameters on COx clay rocks
531 (Chen et al., 2014a; Loni et al., 2021; Montavon et al., 2020); overall, no significant effect
532 was observed. Redox potential is certainly the key parameter; it proves to be even more
533 consistent since uranium is sensitive to redox and the adsorption properties of U(IV) and
534 U(VI) are quite distinct. The potential measured in the solutions prepared in the glove box
535 (and in contact with the clay phases) are oxidizing while that expected under *in-natura*
536 conditions is of the order of -190 mV. This result also aligns with the fact that *in natura*
537 aqueous U concentrations measured in KEY1001 under disturbed oxidizing conditions are
538 higher, i.e. around 2 $\mu\text{g.L}^{-1}$, than those representative of the “natural” conditions (0.13-0.25
539 $\mu\text{g L}^{-1}$). These concentrations are more closely resembling those predicted by our simple
540 model based on laboratory results (1.4 - 12 $\mu\text{g.L}^{-1}$, Fig. 6c).

541 The following scenario can now be proposed, namely that a more retained form of uranium is
542 present under *in natura* conditions, i.e. the U(IV) form. When the sample is removed from its
543 environment, the U(IV) adsorbed on the surface rapidly oxidizes to U(VI), despite the
544 controlled conditions (with no differences in results between preserved and unpreserved COx
545 samples). In the laboratory, this situation will have the effect of increasing the concentration
546 of aqueous U in suspension since U(VI) is less retained than U(IV). It also explains the

547 decrease in U concentration in equilibrium with the CO_x as the solution potential decreases
548 over time; this is the opposite case where the equilibria shift towards the sorption of U(IV)
549 retained to a greater extent by the CO_x clay fraction than U(VI), even in the presence of the
550 soluble ternary complexes of U(VI) (grey symbols in Fig. 6b).

551 According to this reasoning, it becomes possible at this stage to return to Rd values more
552 representative of *in natura*/reducing conditions. Using the labile fraction determined from
553 laboratory experiments and the concentration of U found *in natura* (0.25 - 0.13 μg.L⁻¹), a
554 range of *in natura* Rd between 72 and 323 L·kg⁻¹ can be calculated. This Rd value range is in
555 agreement with the value deduced from the desorption experiments performed at -200
556 mV/NHE and Eq. (2), i.e. 500 ± 200 L kg⁻¹.

557 This reasoning must now be compared with the predictive calculations derived from the
558 operational model under *in natura* conditions. Results are presented in Figure 6 as a function
559 of P_{CO2} and redox potential, the two key parameters controlling U speciation in the formation.
560 In these calculations, pH is determined such that calcite is considered in equilibrium in a
561 SPW1 type solution. Under conditions expected in the formation (hatched area in the figure),
562 Rd values vary greatly, from 640 to 7,400 L.kg⁻¹. From a more mechanistic point of view, the
563 speciation in solution is mainly controlled by U(VI) complexed species, whereas at the
564 surface U(IV) surface complexes govern U speciation (Supplementary data, Fig. S1).

565 On the one hand, the predicted retention range overestimates the one previously discussed
566 based on experimental data (~70-700 L kg⁻¹). Yet on the other, the same conclusion could still
567 be drawn: ternary U(VI) complexes control U speciation in solution while U(IV) surface
568 species dominate U adsorption. It is also important to recall that predictive approaches give
569 trends, since in addition to working with a complex and heterogeneous natural system (other
570 parameters like clay content affect retention values), uncertainties in the model parameters are

571 not taken into account. Let's also recall that the parameters for the U(IV)/CO_x system are
572 extrapolated using the LFER. Therefore, the overlap area between the predicted Rd domain
573 and the Rd domain deduced from experiments allows us to build confidence in our reasoning.

574 **5. CONCLUSION**

575 The aim of this work has been to describe and understand the behavior of U under CO_x
576 conditions. An operational model based on both a bottom-up approach and the 2SPNE SC/CE
577 model has made it possible to describe the reactivity of U(VI) and U(IV) with respect to CO_x
578 under laboratory conditions from batch-type experiments. Compared to the accessible
579 databases, it was however necessary to adjust the constant describing the formation $\equiv\text{S-UO}_2\text{-}$
580 CO₃ ternary surface complexes for illite. Ultimately, it has been well confirmed that the
581 adsorption of uranium onto CO_x claystone is controlled by the clay phase (and more
582 specifically, illite). Moreover, the CO_x clay phase is the largest reservoir of naturally-
583 occurring U. In contrast, only a small fraction appears to have been adsorbed (~1 %, namely
584 ~30 $\mu\text{g}\cdot\text{kg}^{-1}$), i.e. the fraction in rapid exchange equilibrium with the fraction in solution.
585 Depending on the experimental conditions imposed in the laboratory, particularly on the
586 redox potential of the solution, Rd values between ~3 and 700 $\text{L}\cdot\text{kg}^{-1}$ were obtained in order
587 to characterize the behavior of this labile naturally-occurring U fraction. This variability can
588 mainly be explained by a change in U speciation at the surface, from U(VI) for the lowest Rd
589 values to U(IV) for the highest. Under *in natura* conditions, the results of this work indicate a
590 Rd value $> 70 \text{ L}\cdot\text{kg}^{-1}$, thus reflecting a control of U(IV) at the clay CO_x surface. This result
591 now needs to be validated by diffusion experiments on intact CO_x samples.

592

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 599 information and CO_x water samples for analysis, as is S. Betelu from the BRGM Institute for
 600 measuring the redox potential of the long-term experiment.

601

602 **Appendices**

603 **Table A-1:** Characteristics of the CO_x samples; further details on the characterization method
 604 can be found elsewhere (Lerouge et al., 2011)

| Sample | Borehole | Depth (m) | Clay fraction (%) | Clay fraction (wt.%) | | | | Carbonate phases | Quartz + Feldspar | Minor phases | |
|---------------------|----------|-----------|-------------------|----------------------|----------------------|-----------|----------|------------------|-------------------|--------------|------|
| | | | | Illite | I/S _(R=0) | kaolinite | chlorite | | | Pyrite | OM |
| EST51779 / EST51769 | OHZ6126 | 469.72 | 50 | 17 | 28 | 2 | 3 | 25 | 25 | 0.8 | n.m. |
| EST05738 | EST205 | 501.50 | 48.5 | 20.5 | 20.5 | 5 | 2.5 | 25 | 23 | 0.6 | 0.8 |
| EST25687 | PAC1002 | 475.92 | 39.9 | 3.7 | 33 | 1.9 | 1.3 | 28 | 29 | 1.6 | 3.9 |
| EST26536 | TSF1001 | 494.32 | 50 | 18 | 31 | 4.5 | 3 | 22 | 19 | 0.8 | n.m. |
| EST44346 | OHZ1607 | 489.76 | 49.5 | 31.2 | 16.2 | < 1 | 2.1 | 24.4 | 25.5 | 0.2 | 1 |
| EST21400 | PAC2002 | 430.10 | 9.2 | 2 | 7.2 | < 1 | < 1 | 81 | 9.1 | 0.5 | 0.07 |
| EST26480 | FOR1118 | 490.20 | 53 | 21.5 | 25.5 | 3 | 3 | 24 | 22 | 0.8 | n.m. |
| EST05640 | EST205 | 476.57 | 45 | 20 | 23 | <1 | 2 | 25 | 27 | 1.7 | n.m. |

605 OM = organic matter; n.m.: not measured

606

607

608 **Table A-2:** Compositions of the Synthetic Pore Water; concentrations are given in mmol L⁻¹

| | pH | Na | K | Mg | Ca | Sr | Cl | SO ₄ | S(-2) | C(4) | Si | Al | Fe |
|------|-----|------|-----|-----|-----|-----|------|-----------------|-----------------------|----------|-------|-----------------------|--------|
| SPW1 | 7.2 | 40.9 | 1.1 | 5.3 | 8.4 | 0.2 | 40.2 | 13.6 | — | 2.6 | n.m. | n.m. | n.m. |
| SPW2 | 7 | 41.7 | 5.4 | 9.7 | 7.7 | | 73.3 | 4.3 | 3.8 10 ⁻¹⁰ | variable | 0.094 | 9.26 10 ⁻³ | 0.0644 |

609 n.m.: not measured

610

611 **Table A-3:** Detailed experimental conditions in the U(VI) batch experiments

| Sample | Atm-osphere | Solution | m/V (g L ⁻¹) | Type | Alkalinity (meq/L) | FIG |
|---------------|--------------------|------------------------|--------------------------|---------------------------------------------|--------------------|-----|
| Illite | Ar(g) | NaNO ₃ 0.1M | 2 | Edge; [U] ₀ = 10 ⁻⁷ M | 0.04 | 1A |
| Illite | Air | NaCl 0.1M | 0.5 – 1 | Edge; [U] ₀ = 10 ⁻⁷ M | 0.17 | 1A |
| Illite | Ar(g) | NaNO ₃ 0.1M | 2 | Isotherm; pH 6.5±0.2 | 0.07 | 1B |
| Illite | Air | NaNO ₃ 0.1M | 2 | Isotherm; pH 7.3±0.2 | 0.23 | 1B |
| Illite | 1% CO ₂ | NaNO ₃ 0.1M | 2 | Isotherm; pH 7.3±0.1 | 2.4 | 1B |
| Illite | 1% CO ₂ | SPW1 | 10 | Isotherm; pH 7.1±0.2 | 2.4 | 1B |
| Clay fraction | Air | NaNO ₃ 0.1M | 2 | Isotherm; pH 7.1 | 0.22 | 2A |
| Clay fraction | 1% CO ₂ | NaNO ₃ 0.1M | 2 | Isotherm; pH 7.2 | 2.4 | 2A |
| Clay fraction | 1% CO ₂ | NaCl 0.1M | 2 | Isotherm; pH 7.3 | 1.6 | 2A |
| Clay fraction | 1% CO ₂ | SPW1 | 2 | Isotherm; pH 7.1 | 2.5 | 2A |
| EST51779 | 1% CO ₂ | SPW1 | 40 | Isotherm; pH 7.4 | 2.5 | 2B |

612

613 **Table A-4:** Detailed experimental conditions in the U(IV) experiments

| Sample | Atmosphere | Solution | m/V (g L ⁻¹) | pH | Eh | [U] ₀ | FIG |
|----------|---------------------------|----------|--------------------------|-----|---------|--------------------------------------------|-----|
| EST05640 | N ₂ (g) | SPW2 | 0.098 | 8.3 | -270 mV | 3 · 10 ⁻¹² - 10 ⁻⁸ M | 3A |
| EST05640 | 3,500 ppm CO ₂ | SPW2 | 0.098 | 7.6 | -210 mV | 5 · 10 ⁻¹² - 10 ⁻⁹ M | 3B |
| EST05640 | 9,500 ppm CO ₂ | SPW2 | 0.098 | 7.1 | -203 mV | 4 · 10 ⁻¹² - 10 ⁻⁹ M | 3C |

614

615

616

617 **Table A-5:** Surface complexation and cation exchange reactions for the uranium used in
 618 modeling. Unless otherwise specified, the constants have been extracted from the Bradbury
 619 and Baeyens model (Baeyens and Bradbury, 2017; Bradbury and Baeyens, 2017).

| Reactions | Illite (Log K) | Montmorillonite (Log K) |
|-------------------------------------------------------------|----------------|-------------------------|
| Hexavalent reactions | | |
| $sOH + UO_2^{2+} = sOUO_2^+ + H^+$ | 2 | 3.1 |
| $sOH + UO_2^{2+} + H_2O = sOUO_2OH + 2 H^+$ | -3.9 | -4.6 |
| $sOH + UO_2^{2+} + 2 H_2O = sOUO_2(OH)_2^- + 3H^+$ | -10.8 | -12.6 |
| $sOH + UO_2^{2+} + 3 H_2O = sOUO_2(OH)_3^{2-} + 4H^+$ | -18.7 | -20.9 |
| $w_1OH + UO_2^{2+} = w_1OUO_2^+ + H^+$ | 0 | 0.5 |
| $w_1OH + UO_2^{2+} + H_2O = w_1OUO_2OH + 2 H^+$ | -5.8 | -5.7 |
| $w_1OH + UO_2^{2+} + 2 H_2O = w_1OUO_2(OH)_2^- + 3 H^+$ | (***) | |
| $sOH + UO_2^{2+} + CO_3^{2-} = sOUO_2CO_3^- + H^+$ | 11.5 (*) | 9.8 |
| $sOH + UO_2^{2+} + 2 CO_3^{2-} = sOUO_2(CO_3)_2^{3-} + H^+$ | 17.3 (*) | 15.5 |
| $w_1OH + UO_2^{2+} + CO_3^{2-} = w_1OUO_2CO_3^- + H^+$ | | 9.3 |
| $2 NaX + UO_2^{2+} = UO_2X_2 + 2 Na^+$ | 0.65 | 0.146 |
| Tetravalent reactions | | |
| | (**) | (**) |
| $sOH + U^{4+} = sOU^{3+} + H^+$ | 7.62 | 8.24 |
| $sOH + U^{4+} + H_2O = sOUOH^{2+} + 2 H^+$ | 7.1 | 7.7 |
| $sOH + U^{4+} + 2 H_2O = sOU(OH)_2^+ + 3H^+$ | 3.6 | 4.0 |
| $sOH + U^{4+} + 3 H_2O = sOU(OH)_3 + 4H^+$ | -1.6 | -1.4 |
| Amphoteric reactions | | |
| $s/w_1OH + H^+ = s/w_1OH_2^+$ | 4.0 | 4.5 |
| $s/w_1OH = s/w_1O^- + H^+$ | -6.2 | -7.9 |
| $w_2OH + H^+ = w_2OH_2^+$ | 8.5 | 6.0 |
| $w_2OH = w_2O^- + H^+$ | -10.5 | -10.5 |

620 $sOH =$ strong site (density = 2 $\mu\text{mol/g}$); $w_1OH =$ weak site (density = 40 $\mu\text{mol/g}$); $w_2OH =$ weak site (density =
 621 40 $\mu\text{mol/g}$)

622 $NaX =$ ion exchange site (CEC illite = $2.25 \cdot 10^{-4}$ eq/g, montmorillonite = $8.7 \cdot 10^{-4}$ eq/g)

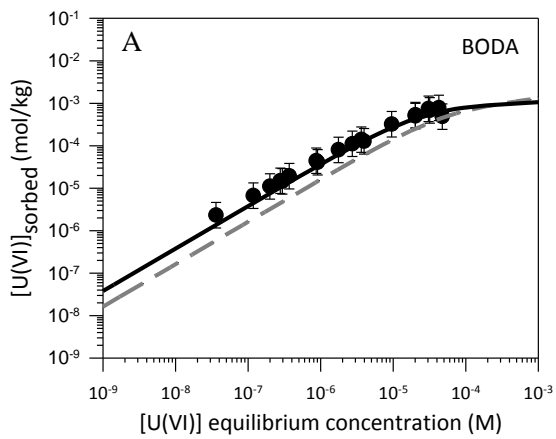
623 (*) this work

624 (***) LFER associated with Th^{4+} data: (Montmorillonite: $\text{Log K}(S_{x-1}) = 1.02 \cdot \text{Log K}(OH_x) + 8.79$; illite: $\text{Log K}(S_{x-1}) = 0.97 \cdot \text{Log K}(OH_x) + 8.14$)

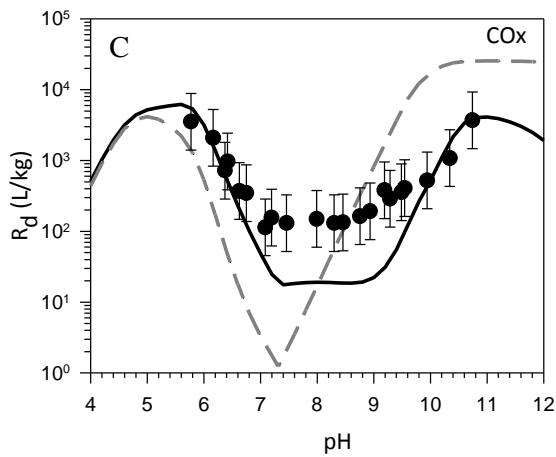
625 (***) Note that the equilibrium leading to the surface species $w_1UO_2(OH)_2$ was not considered
 626 because its contribution in our experimental conditions could be neglected

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630 **Figure A-1:** Adsorption of uranium on clay minerals in equilibrium atmospheric CO₂.
 631 (A) BODA Claystone (Marques Fernandes et al., 2015); (B) OPA Claystone (Marques
 632 Fernandes et al., 2015) (MF15) and (Joseph et al., 2011) (JOS11); (C) COx claystone in 0.1 M
 633 NaClO₄ (Hartmann et al., 2008). Symbols = experimental data. The grey dashed lines
 634 represent the published model. The black lines correspond to our adjusted model (Table A5).
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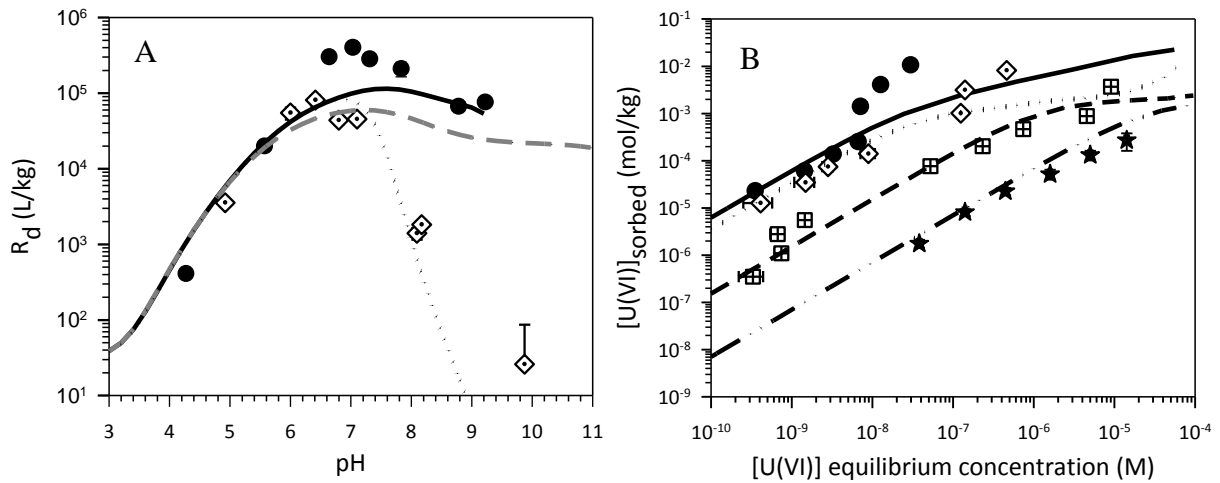
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Table 1: Summary of the experiments conducted in this study

| Objective | Samples | | Tasks | Conditions | Corresponding Figure(s) and/or Table |
|------------------------------------|-------------------|----------------------------------------------------------|----------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------|
| Model building | Illite | Purified | Evaluation of the carbonate effect on U adsorption onto illite | SPW1 / 0.1M NaCl / 0.1M NaNO ₃ Ar / P _{CO2} = 10 ^{-3.5} / 10 ⁻² / 10 ^{-1.3} pH: 4 - 9.8 m/V: 2 g/L [U] ₀ = 10 ⁻⁹ - 2·10 ⁻⁵ M | Figure 1 Table A |
| | COx-clay fraction | Purified from EST51779 | Model testing on the COx clay fraction | 0.1M NaNO ₃ / SPW1 P _{CO2} = 10 ^{-3.5} / 10 ⁻² pH: 6.7 - 7.1 m/V: 2 g/L [U] ₀ = 5·10 ⁻⁸ - 2·10 ⁻⁵ M | Figure 2A Table A |
| | COx claystone | EST51779 | Model testing on the COx under oxidizing conditions | SPW1 P _{CO2} = 10 ⁻² pH: 7.4 m/V: 40 g/L [U] ₀ = 5·10 ⁻⁸ - 2·10 ⁻⁵ M | Figure 2B Table A |
| | COx claystone | EST05640 | Model testing on the COx under reducing conditions | SPW2 N ₂ / P _{CO2} = 10 ^{-3.5} / 10 ⁻² Eh -185 mV pH: 7.1 - 8.3 m/V: 0.1 g/L [U] ₀ = 3·10 ⁻¹² - 10 ⁻⁸ M | Figure 3 Table B |
| Naturally-occurring Uranium | COx claystone | EST51769 / EST05738 / EST26536 | Uranium distribution and solid analyses | — | Table S1 Figure 4C |
| | | EST25687 / EST25687(ox) / EST21400 / EST26536 / EST26480 | Sequential extraction | See Section 2.2 | Figure 4 |
| | | EST44346 | Determination of the labile fraction | SPW1 [²³³ U] = 3.10 ⁻¹⁰ M Isotope exchange method | Figure 5 |
| | | EST25687(ox) / EST51779 | | SPW1 Desorption method | |

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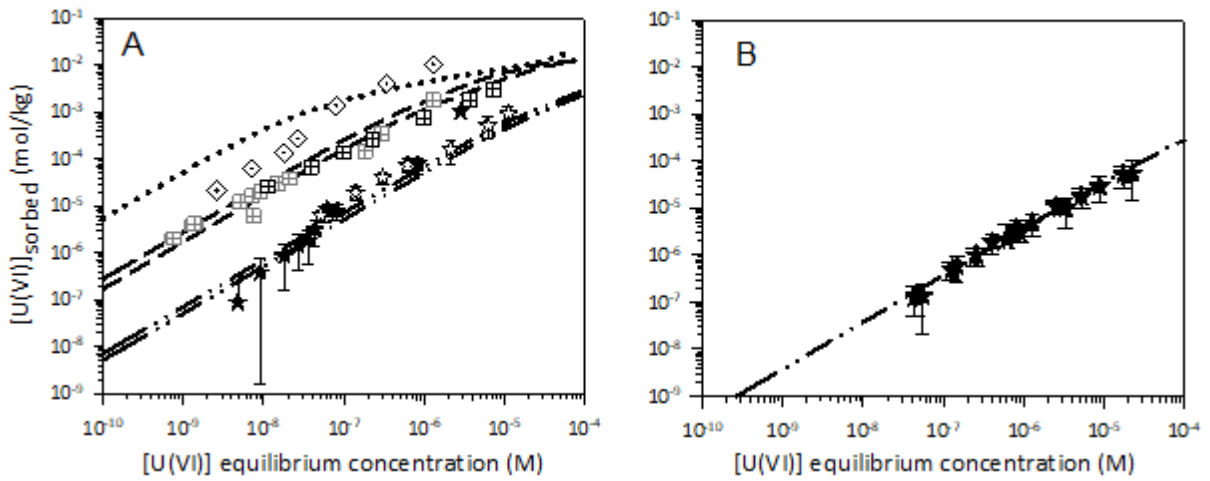
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772 Figure 1: Adsorption of U(VI) onto illite under various experimental conditions (see Tables 1
773 and A3). (A) variation vs. pH, (B) concentration isotherms. Symbols = experimental data
774 (circle: 0.1 M NaNO₃ in ~Ar atmosphere, diamond: 0.1 M NaNO₃ in ~atmospheric CO₂;
775 square: 0.1 M NaNO₃ or NaCl in ~1% CO₂, star = SPW1 in ~1% CO₂). The grey dashed line
776 depicts the model for 0.1 M NaClO₄ and CO₂ free atmosphere without considering the surface
777 complexes with carbonates. Black lines correspond to our adjusted model (Table A5).

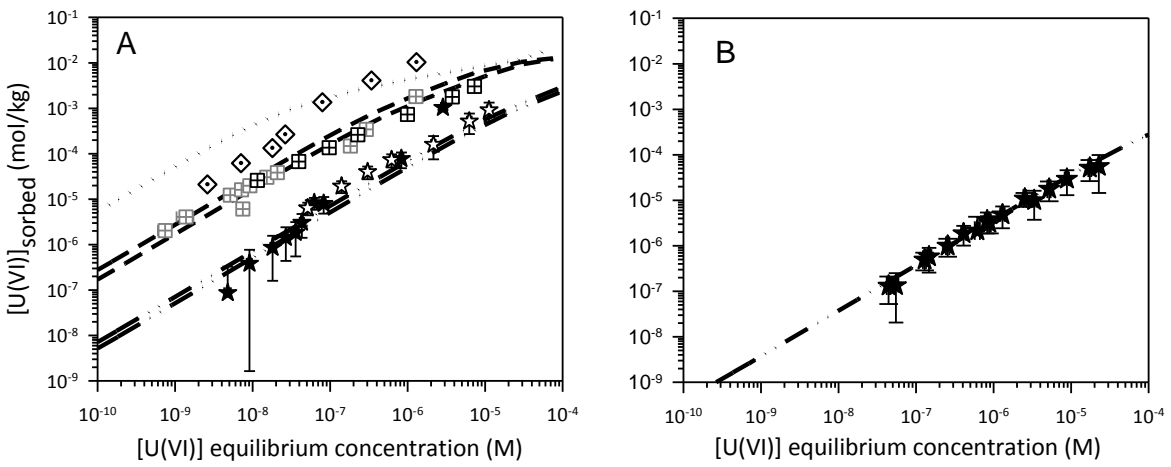
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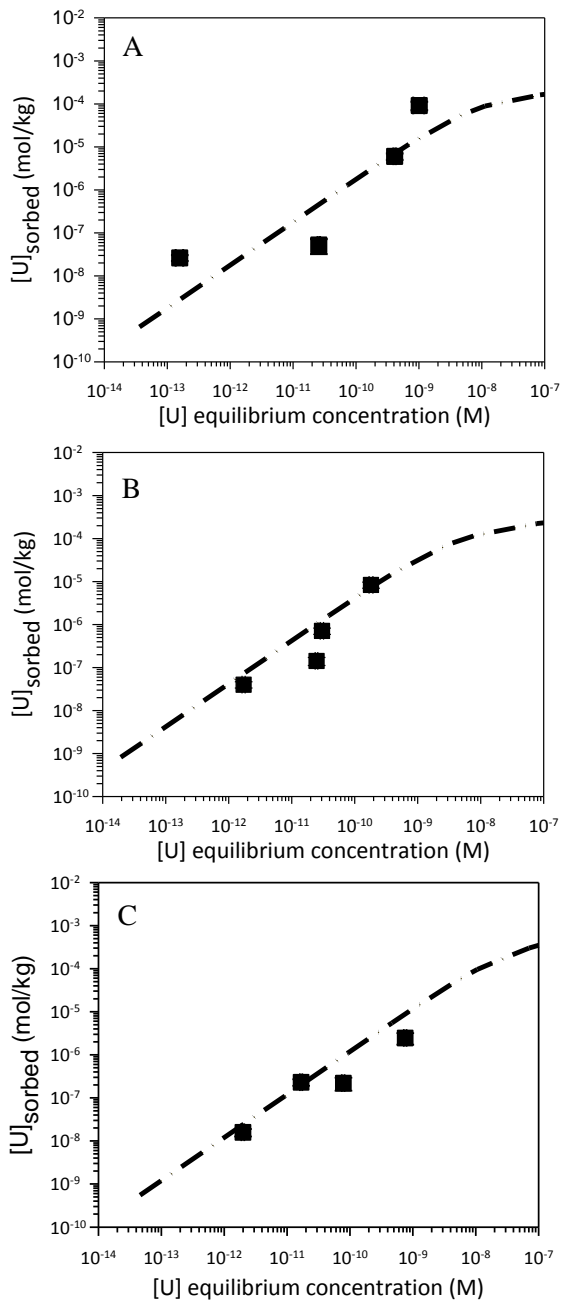
783 Figure 2: Adsorption of uranium(VI) on: (A) COx clay fraction, and (B) COx rock (see
784 Tables 1 and A3). Symbols = experimental data (diamond: ~atmospheric CO₂ pressure,
785 square: 0.1 M NaNO₃ or NaCl in ~1% CO₂, star = SPW1 composition in 1% CO₂). The lines
786 correspond to our adjusted model (Table A5).

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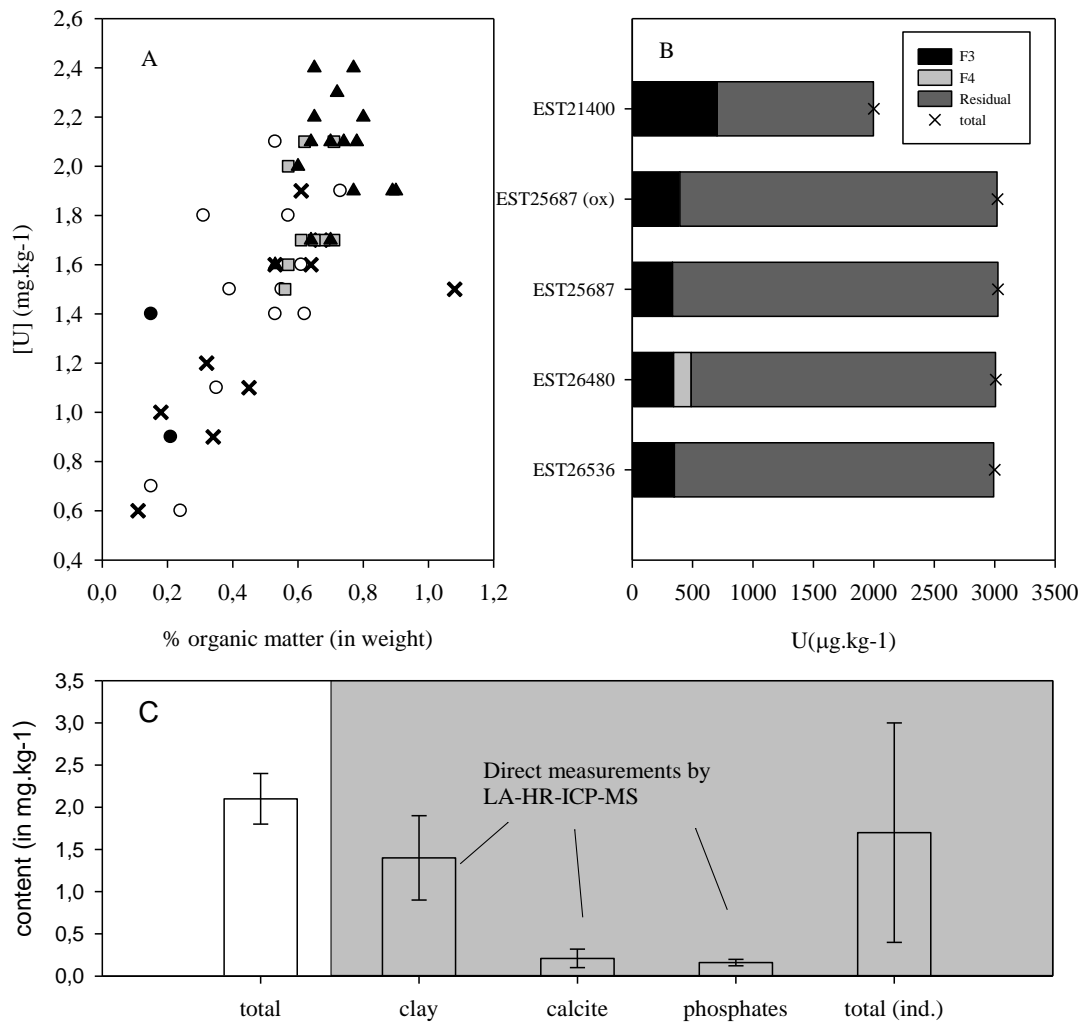
791 Figure 3: Adsorption of uranium on CO_x under reducing conditions for various atmospheres:
792 (A) Ar (pH = 8.3, Eh = -265 mV), (B) air (pH = 7.5, Eh = -215 mV), and (C) 1% CO₂ (pH =
793 7.1, Eh = -201 mV) (see Table B). The lines depict the model results (modeling parameters
794 are given in Table A5).

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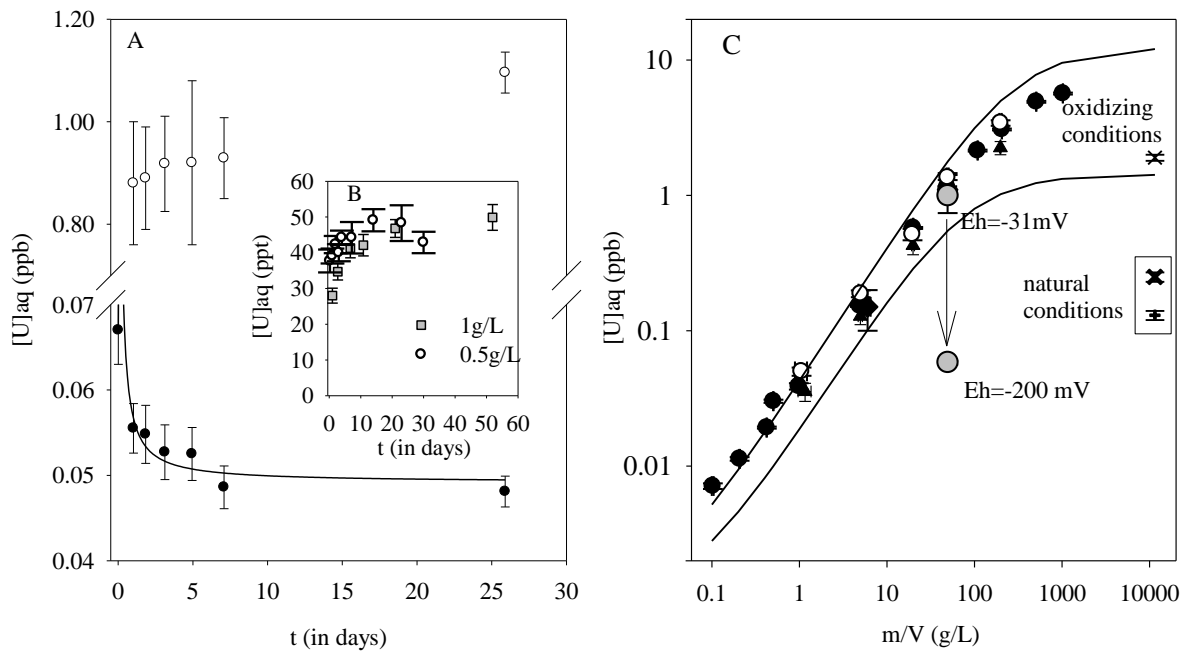


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800 **Figure 4:** Naturally-occurring U in the COx: (A) U content vs. organic matter content. (B)
801 Sequential extraction results; the U concentrations in Fractions 1 and 2 are not shown here
802 due to their weak contribution. (C) Distribution of U between the various selected phases; all
803 values are given in Table S3.

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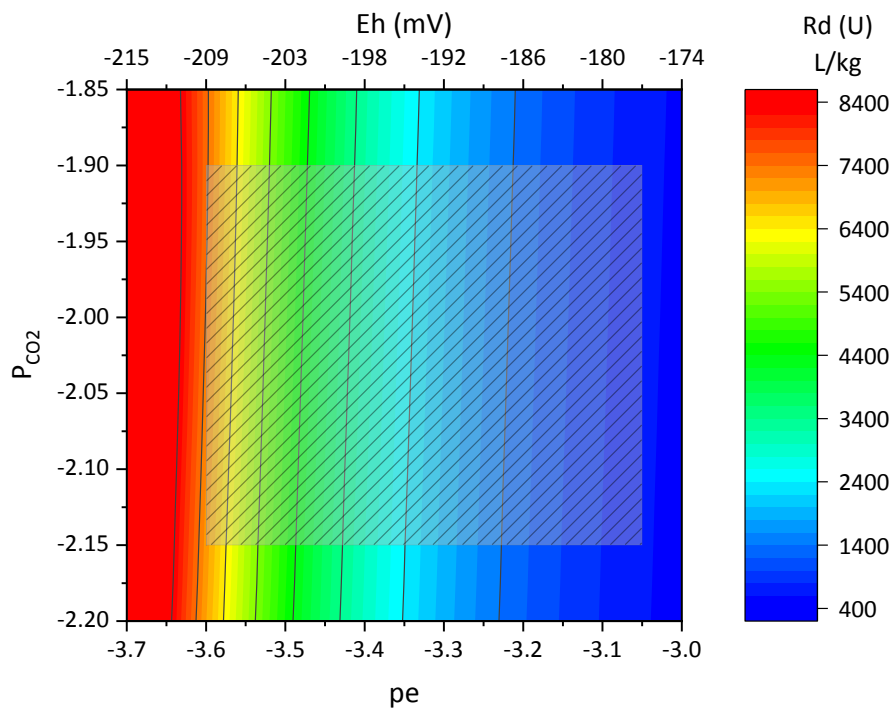
809 **Figure 5:** Results of equilibrium-based methods. (A) Isotopic exchange method with $S/L =$
 810 49.5 g/L ; U-238 (O) and U-233 (●) concentrations measured over time. The line shows a
 811 trend curve. (B) U-238 concentration measured over time. (C) Desorption method; the dark
 812 symbols correspond to three data series (EST51779, etc.); the white symbol represents the
 813 series with EST25687(ox); and the gray symbol denotes the experiment performed over a
 814 long contact time. The lines depict the fitting/calculation results (Eq. 2) (see text). The
 815 symbols on the right reflect concentrations measured directly in the pore water under
 816 preserved and unpreserved conditions.

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822 **Figure 6:** Variation in the distribution coefficient under *in natura* conditions vs. pe and P_{CO_2}
823 (pH determined for calcite equilibrium, with $[Ca]=8.4$ mmol/L). The hatched area
824 corresponds to expected *in natura* conditions.

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