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1 **Seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios along continental margins: patterns and processes**
2 **in open and restricted shelf domains**

3

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18

19 **Highlights:**

- 20 • Homogeneous $^{87}\text{Sr}/^{86}\text{Sr}$ ratios similar to the oceanic value in open shelf contexts
- 21 • Noticeable heterogeneity of $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in lagoons and epeiric seas
- 22 • Importance of submarine groundwater discharges which may locally supply 10 to
23 60% of Sr in lagoons
- 24 • Oceanic Sr inputs and $^{87}\text{Sr}/^{86}\text{Sr}$ homogenization controlled by the coastal morphology

25 **Abstract**

26 To better constrain the Sr isotope budget in marginal domains without any fluvial inputs, we
27 analyzed the chemical composition and $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of waters and shells from four
28 locations: two coastal lagoons, one hemipelagic platform and one open marine shelf. Our
29 results highlight homogeneous $^{87}\text{Sr}/^{86}\text{Sr}$ ratios typical of oligotrophic oceanic waters (OOW)
30 (i.e., 0.709172 ± 0.000023) in the Pacific Tatakoto atoll and along a Mediterranean shore to
31 offshore transect (~25km off Banyuls-sur-Mer, BSM). This attests that oceanic inputs from
32 oligotrophic areas remain the main Sr source in open shelf areas compared with submarine
33 groundwater discharges (SGD) or particulate dissolution influences. In BSM, only foreshore
34 data are more radiogenic, possibly due to rainwater mixing, local groundwater springs or
35 more efficient particle dissolution in the intertidal zone. In restricted areas, we report variable
36 $^{87}\text{Sr}/^{86}\text{Sr}$ ratios between the Salses-Leucate (France) and Oualidia (Morocco) lagoons. The
37 first one has homogeneous $^{87}\text{Sr}/^{86}\text{Sr}$ ratio typical of OOW except close to groundwater
38 discharges. In Oualidia, $^{87}\text{Sr}/^{86}\text{Sr}$ ratios decrease by 1.2×10^{-3} from OOW values close to the
39 Atlantic inlet to progressively less radiogenic ones upstream within the interior of the lagoon.
40 These differences depend on several factors including the leaky, restricted or choked
41 morphology of lagoons modulating the oceanic Sr inputs, but also SGD fluxes whose $^{87}\text{Sr}/^{86}\text{Sr}$
42 ratios and Sr concentrations are highly variable according to the nature of rocks leached in
43 karstic aquifer. In Oualidia, the low $^{87}\text{Sr}/^{86}\text{Sr}$ ratios correspond to high Sr concentrations (up
44 to $150 \mu\text{mol}\cdot\text{l}^{-1}$) issued from the dissolution of Mesozoic evaporites, leading to SGD fluxes
45 accounting for 60% of the local Sr budget. Through data compilation, we show that similar
46 $^{87}\text{Sr}/^{86}\text{Sr}$ gradients and processes prevail at the whole Mediterranean scale. Finally, we
47 postulate that high coastal water retention times can also account for anomalous coastal
48 $^{87}\text{Sr}/^{86}\text{Sr}$ ratios and that the combination of water mass restriction, SGD, bioadsorption and
49 early diagenetic processes could decrease seawater Sr concentrations in some marginal areas.

50 **1. Introduction**

51 Although the strontium isotope compositions ($^{87}\text{Sr}/^{86}\text{Sr}$) of river waters and submarine
52 hydrothermal sources strongly vary across the globe (with respective averages of 0.7119 and
53 0.7037; Palmer and Edmond, 1989; 1992; Bach and Humphris, 1999; Davis et al., 2003;
54 Pearce et al., 2015; Chavagnac et al., 2018), the $^{87}\text{Sr}/^{86}\text{Sr}$ of oligotrophic oceanic waters
55 (OOW) remains homogeneous worldwide at 0.709172 ± 0.000023 (see El Meknassi et al.,
56 2018 for a recent synthesis). This paradigm is due to the long residence time of Sr in the
57 ocean (i.e., ~2.5 Myr; Hodell et al., 1990) compared to the global ocean mixing duration
58 ranging from 1000 to 2000 years (DeVries and Primeau, 2011). However, the OOW $^{87}\text{Sr}/^{86}\text{Sr}$
59 ratios fluctuated markedly through the Phanerozoic, as reported from Sr isotope data of fossil
60 archives (Peterman et al., 1970; Burke et al., 1982; Veizer, 1989; Zaky et al., 2018). These
61 secular variations are ascribed either to global geodynamic events modulating the respective
62 contribution of terrestrial (radiogenic) and hydrothermal (unradiogenic) Sr sources to oceans
63 or to paleogeographic and climatic changes modifying the continental weathering of rocks of
64 different age and lithology (Reeder et al., 1972; Brass, 1976; Goldstein and Jacobsen, 1987;
65 Bataille et al., 2017; Peucker-Ehrenbrink and Fiske, 2019). By assuming a homogeneous
66 OOW $^{87}\text{Sr}/^{86}\text{Sr}$ ratio at a global scale, these long-term $^{87}\text{Sr}/^{86}\text{Sr}$ variations are thus widely
67 used for dating sedimentary series through chemostratigraphic calibration of new fossil data
68 (McArthur, 1994; McArthur et al., 2012) or testing paleogeographic and climatic hypotheses
69 (Goddéris et al., 2017).

70 Despite the popularity of this paleoenvironmental proxy, the question of whether fossil
71 carbonate shells may be used to reconstruct past OOW $^{87}\text{Sr}/^{86}\text{Sr}$ fluctuations is still a matter of
72 debate as most of these organisms thrived in epeiric and shelf domains potentially
73 disconnected from the global oceanic Sr reservoir. This is partly supported by the apparent
74 heterogeneity of modern shell and (euhaline to brachyhaline) seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios from

75 worldwide marine shelves, with respective ranges of 0.707636–0.710483 and 0.704638–
76 710648 (precision on measured $^{87}\text{Sr}/^{86}\text{Sr}$ ratio is better than 10^{-5} ; El Meknassi et al., 2018).
77 Significant offsets between well-dated fossil (or bulk carbonate) $^{87}\text{Sr}/^{86}\text{Sr}$ ratios and the
78 Phanerozoic isotope curve are also regularly reported, even in sedimentary facies indicating
79 outer shelf contexts and/or normal marine environments (Cochran et al., 2003; Nieto et al.,
80 2008; Sessa et al., 2012; Wierzbowski et al., 2012; Eidvin et al., 2014; Schildgen et al., 2014;
81 Briard et al., 2020). While potential analytical, stratigraphical and diagenetic biases may be
82 discarded (Martin and Scher, 2004; Marcano et al., 2015; Bellefroid et al., 2018; Zaky et al.,
83 2018), these mismatches can indicate that, sporadically, seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of epeiric
84 and shelf domains are not necessarily representative of the global OOW value. In most case,
85 these variations result from dissolved Sr supplies delivered by river waters, with contrasted
86 influences on seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios according to their Sr concentrations and isotopic ratios
87 linked to the age and lithology of local bedrocks (Palmer and Edmond, 1989, 1992; Bryant et
88 al., 1995; Peucker-Ehrenbrink et al., 2010). This terrestrial Sr contribution is common in
89 restricted environments like epeiric seas, estuaries, fjords, lagoons, and bays where oceanic
90 influences are more restricted and where $^{87}\text{Sr}/^{86}\text{Sr}$ ratios covary landward with salinity
91 (Andersson et al., 1992; Ingram and Sloan, 1992; Israelson and Buchardt, 1999; Basu et al.,
92 2001; Major et al., 2006; Jones et al., 2014; Beck et al., 2013; Wang and You, 2013;
93 Chakrabarti et al., 2018; Shao et al., 2018). However, this river water influence can remain
94 limited in euhaline marginal contexts as, on average, measurable effects on the seawater
95 $^{87}\text{Sr}/^{86}\text{Sr}$ ratio are not expected to occur if freshwater inputs do not drop salinity below a
96 threshold value of 12 (Bryant et al., 1995). Only 5 % of world river systems have water
97 $^{87}\text{Sr}/^{86}\text{Sr}$ ratios sufficiently different from the OOW value to significantly modify the local
98 seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratio in euhaline marine environments.

99 Although poorly documented, seawater and shell $^{87}\text{Sr}/^{86}\text{Sr}$ ratios ranging from 0.70890
100 to 0.70921 have been yet reported in deep outer shelf domains and lagoons not directly
101 subject to major river supply (Müller et al., 1990a; 1990b; Peckmann et al. 2001; Major et al.
102 2006; Huang et al., 2011; El Meknassi et al., 2018). This suggests that submarine groundwater
103 discharges (SGD) or strontium released from particulate dissolution have a much stronger
104 impact on the coastal Sr budget than currently expected (Beck et al., 2013; Jones et al., 2014 ;
105 Trezzi et al., 2017). A direct consequence is that, depending on oceanic mixing and water
106 retention times on shelves (i.e., ranging from a couple of days to decades or centuries; Liu et
107 al., 2019), the Sr concentrations and $^{87}\text{Sr}/^{86}\text{Sr}$ of marginal areas, could not always be
108 representative of the global oceanic reservoir. In other words, seawater from open shelves and
109 epeiric domains could record Sr isotope anomalies just through cumulative effects of sporadic
110 terrestrial Sr inputs in coastal sub-reservoirs not sufficiently mixed with the global oceanic
111 reservoir.

112 Here, we test how submarine groundwater discharges (SGD) and water mass
113 restriction can influence the Sr isotope composition of shelf water devoid of river influences
114 by measuring and comparing the $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of waters and shells from four coastal
115 domains with different levels of connection to the open ocean. These include an open shelf
116 Mediterranean transect, two coastal lagoons influenced by groundwater discharges in the
117 Atlantic and Mediterranean contexts, and a Pacific hemipelagic platform. Through data
118 compilation, our aim is also to better document the worldwide variability of seawater $^{87}\text{Sr}/^{86}\text{Sr}$
119 ratios in the coastal ocean and to better constrain the environmental processes acting on
120 seawater $^{87}\text{Sr}/^{86}\text{Sr}$ patterns in these transition zones.

121

122

123 2. Geological Setting

124 The seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of three restricted lagoons exhibiting different
125 geomorphological, hydrogeological, hydrodynamical and climatic conditions is investigated
126 in this study (Fig. 1): 1) The Tatakoto atoll in the eastern Tuamotu Archipelago (French
127 Polynesia) in the southern part of the Pacific Ocean; 2) the Salses-Leucate lagoon (France) in
128 the northwestern Mediterranean domain; and 3) the Oualidia lagoon (Morocco) in the North
129 Atlantic domain. These three sites were chosen because they are under the influence of
130 different littoral or reefal barriers, their water resources are issued from various karstic
131 reservoirs, and they have different waves and tide forcing overprinted by different climatic
132 conditions. To get a comparison with an open shelf domain, sampling was also carried out in
133 the Mediterranean Sea along a 25 km-long transect from the rocky coastline of Banyuls-sur-
134 Mer to the submarine Lacaze-Duthiers Canyon (western part of Gulf of Lion).

135 2.1. Oualidia Lagoon (Morocco)

136 Formed around 8000 – 6500 BP (Ballouche and Carruesco, 1986), the Oualidia coastal
137 lagoon is located on the western coast of Morocco (Abda Doukkala region) (Fig. 1A) and
138 belongs to a coastal endorheic basin called the Coastal Sahel of Oualidia. The lagoon is 7 km
139 long, 400 m wide, and 4m water deep during the high tide season (Kaid Rassou et al., 2005).
140 It is organized around a main channel of 180 m wide parallel to the coastline and separated
141 from the Atlantic Ocean by a Plio-Quaternary shoal (Ballouche and Carruesco, 1986). The
142 lagoon is connected to the ocean by two inlets to the south (130 and 70 m wide) (El Khalidi et
143 al., 2011), but recent anthropic developments for salt exploitation and eutrophication
144 prevention along the northern part allowed punctual oceanic inflows (Hilmi et al., 2009). Its
145 hydrodynamic balance is influenced by: 1) Atlantic marine inflows linked to important semi-
146 diurnal tidal dynamics (i.e., average amplitude of 0.97 m and main currents $>1 \text{ m}\cdot\text{s}^{-1}$; Hilmi et
147 al., 2005; 2009), 2) submarine groundwater supplies estimated from 0.2 to $1.2 \text{ m}^3\cdot\text{s}^{-1}$ (Fakir et

148 al., 2019), and 3) evaporation rates of 1.3 to 1.5 m.yr⁻¹ characteristic of semi-arid conditions
149 in this region (with precipitation rates of 30 mm.yr⁻¹) (Hilmi et al., 2009). Along seasonal
150 climatic and hydrodynamic conditions, the recurrent groundwater discharge to the lagoon is
151 highlighted by a net salinity gradient ranging from 4 to 36 along a north to south transect
152 (Hilmi et al., 2017). These groundwater supplies occur all along the lagoon (Fakir et al., 2019)
153 and are predominantly linked to the extended karstic aquifer of the Coastal Sahel of Oualidia
154 (Fakir et al., 2002; Kaid Rassou et al., 2005; Bouchaou et al., 2017). Its upper part (i.e., the
155 most exploited for freshwater consumption) is composed of a 50 m thick Plio-Quaternary
156 limestone-sandstone overlying a less permeable level of upper Hauterivian red clays. Below,
157 the 30 m thick Dridrate limestone formation represents the lower aquifer which is less
158 exploited but still provide the most abundant freshwater resources (Fakir et al., 2002; Kaid
159 Rassou et al., 2005). Finally, the upper Valanginian marls constitute the substratum of the
160 Dridrate aquifer and covers the Berriasian limestones, the upper Jurassic limestones rich in
161 gypsum levels, and Triassic evaporites (Fakir et al., 2002; Fadili et al., 2015). According to
162 Fakir et al. (2002), the dissolution of evaporites would be responsible for high Sr
163 concentrations in local groundwater, reaching 2 to 2.5 times the OOW value.

164

165 *2.2. Salses-Leucate Lagoon (South of France)*

166 The Salses-Leucate lagoon formed during the Flandrian (i.e., 15 kyr BP; Arnaud and
167 Raimbault, 1969; Clanzig, 1987). It corresponds to a shallow coastal basin separated from the
168 Mediterranean Sea by a sandy barrier interrupted by three narrow and dispatched marine
169 inlets (two of them are artificial ones) (Ladagnous and Le Bec, 1997; Fig. 1B). The lagoon is
170 14 km long and 5 km wide, with an average water depth of 1.7 m reaching 3.7 m in its deepest
171 areas (Stieglitz et al., 2013). The region of Salses-Leucate is characterized by a dry season
172 typical of the Mediterranean climate, with annual rainfalls of 588 mm.yr⁻¹ and annual

173 evaporation rates of 1.5 m.yr^{-1} (Ladagnous and LeBec, 1997). Basically, wind-driven currents
174 control the hydrodynamic balance of the lagoon by provoking lagoon outflows and
175 Mediterranean seawater inflows, both of them overprinted by moderate tidal influences (i.e.,
176 tidal range of 0.40 m; Ladagnous and le Bec, 1997). The lagoon hydrology is affected by
177 freshwater inputs from two main karstic systems in its southwestern parts (i.e., the Font Dame
178 and Font Estramar discharges) that are responsible for 98% of freshwater inputs into the
179 lagoon, with fluxes of 8400 and 12500 $\text{m}^3.\text{h}^{-1}$ respectively (Ladagnous and Le Bec, 1997;
180 Stieglitz et al., 2013). Altogether, these parameters produce spatial and seasonal salinity
181 variations in the lagoon ranging from 20 to 30 (Bejannin et al., 2017). Finally, it is worth
182 noting that the Font Estramar and Font Dame karstic resurgences belong to the well-studied
183 Corbières karstic domain, which is composed of Jurassic and Cretaceous dolomites,
184 limestones and marls covering Triassic evaporites, and constitutes one of the biggest karstic
185 network from Europe (Ladouche and Dörfliger 2004; Aunay et al., 2003).

186 *2.3. Banyuls-sur-Mer transect (South of France)*

187 The Banyuls-sur-Mer (BSM) coast is located on the most eastern part of the Pyrenean
188 reliefs (i.e., Massif des Albères) where the substratum lithology is dominated by metamorphic
189 and crystalline continental rocks of Cambrian age (Fig. 1C; Got and Stanley, 1974).
190 Influenced by a Mediterranean climate (with annual rainfalls of 575 mm.yr^{-1}), this open rocky
191 coast displays no major freshwater input apart from the Baillaury river (mean flow rate of 0.2
192 $\text{m}^3.\text{s}^{-1}$) that can show very important freshwater discharges in rainy season (up to $165 \text{ m}^3.\text{s}^{-1}$
193 during floods). BSM is characterized by a low tidal range with an average of 0.40 m. From
194 the BSM beach, a 25 km-long W-E transect was carried out at sea up to the Lacaze-Duthiers
195 (LD) Canyon in order to collect surface and bottom seawater (i.e., up to 450 m water depth in
196 the canyon). The LD canyon is one of the numerous canyons of the Gulf of Lion, with a
197 strong bottom current and high-suspended sediment supplies from seasonal gravity flow

198 events (Heussner et al., 2006). The salinity and temperature of seawater is regularly measured
199 by the Oceanological Observatory of Banyuls-sur-Mer in the framework of the SOMLIT
200 program, both at SOLA (in the Banyuls Bay) and MOLA (offshore above the canyon) buoys.
201 The surface seawater salinity varies between 34 and 38 whereas bottom seawater varies
202 between 37 and 39. The salinity at 450 m depth fluctuates slightly around 38 (Durrieu de
203 Madron et al., 2013).

204 *2.4. The Tatakoto lagoon (French Polynesia, Pacific Ocean)*

205 The Tatakoto lagoon is a semi-enclosed hemipelagic carbonate platform located in the
206 eastern Tuamotu Archipelago in the Pacific Ocean (Fig. 1D). This archipelago spreads over
207 1600 km and is characterized by 77 atolls whose reef-barrier caps volcanic basement dated
208 between 44 and 37.5 Myrs (Rougerie, 1994; Pirazzoli, 1998). The climate is tropical and
209 dominated by a seasonal rainfall regime of 50 to 100 mm monthly from June to October and
210 100 to 200 mm monthly from November to May (Andréfouët et al., 2001). The Tatakoto atoll
211 is about 12 km long and 3 km large, and the lagoon covers an area of $\sim 12 \text{ km}^2$ (Gilbert et al.,
212 2006) with a maximal water depth measured at 18 m (C. Brahmi, personal communication).
213 The atoll rim is closed in the northern part and semi-opened in the southern part. Water
214 exchanges between the lagoon and the ocean occur through several shallow inlets located on
215 the southern side and seawater mixing depends on tide, wave and wind regimes (Andréfouët
216 et al., 2012; Van Wynsberge et al., 2017). The seawater exchanges between the atoll inner
217 parts and ocean are weak owing to the low tidal range (with a maximum amplitude of 0.2 to
218 0.35 m). In the Tuamotu islands, water percolation through the reef porosity (i.e., 30 to 60%)
219 is linked to erosion-dissolution and karstification processes (Rougerie, 1994). Note that
220 intermediate oceanic seawaters flow through the porosity network of the barrier by endo-
221 upwelling convection, leading to the formation of interstitial waters representing $\sim 50\%$ of the
222 barrier volume (Rougerie, 1994). Thus, the renewal of the atoll seawater volume requires

223 several weeks up to several months depending on the morphology, the porosity of the barrier
224 and the seasonal hydroclimatic conditions (Rougerie, 1994; Tartinville et al., 1997).
225 Combined with evaporation rates of 0.5 to 1 m.yr⁻¹, all these processes impose lagoonal
226 salinities fluctuating between 30 and 44 (Rougerie et al., 1984).

227

228 **3. Material and Methods**

229 *3.1. Water and mollusc sampling*

230 In order to check the spatial variability of seawater ⁸⁷Sr/⁸⁶Sr ratios in coastal domains,
231 38 water samples and 18 carbonate shells of living molluscs (i.e., bivalves and gastropods)
232 were collected at the four studied sites (Fig. 1; Tables 1 and 2). When and where possible, we
233 collected both the surface and bottom seawater together with molluscs for a given position.

234 In the field, the water samples were collected in 1L LDPE bottles previously cleaned
235 with ultrapure 1N HCl, filtered *in situ* using 60 ml sterile plastic syringes with 0.2 µm
236 Millipore filters (*in-situ* ultrafiltration was not possible in the field due to logistic constraints).
237 All water samples are void of any microparticles usually found as suspended particle matter in
238 the natural environment. *In-situ* filtration during water collection can significantly reduce
239 microparticle-water interaction and therefore improve the characterization of the dissolved
240 chemical pool (Cotte et al., 2015). They were stored at 4°C in a cold room prior to any sample
241 preparation in a clean laboratory for chemical analyses. In the Oualidia and Salses-Leucate
242 coastal lagoons, the water samples were collected (in April and June 2018, respectively) from
243 0 to 2 m water depth by diving from a chartered semi-rigid boat at various locations from
244 landward parts affected by groundwater discharges to marine inlets and beaches facing the
245 open marine realm (Fig. 1A,B). Bivalve shells from the Oualidia lagoon include *Crassostrea*
246 *gigas*, *Venerupis decussata*, and *Solen marginatus*, whereas those from Salses-Leucate
247 correspond to *Mytilus galloprovincialis*, *Crassostrea sp.*, and *Glycymeris glycymeris*. Note

248 that it was possible to collect directly the groundwaters from the Font-Dame and Font-
249 Estramar resurgences at Salses-Leucate lagoon. However, the groundwater discharges of the
250 Oualidia lagoon are mostly submerged, sparse and difficult to locate without *in-situ* chemical
251 and salinity monitoring (Fakir et al., 2019).

252 In the Tatakoto lagoon, two water samples and four *Tridacna maxima* shells have been
253 all collected at 2 m depth from two locations: one close to the southern oceanic inlets and the
254 other one in the northern quiescent part of the inner lagoon (Fig. 1D).

255 Finally, a proximal-distal transect between the BSM beach and LD Canyon was
256 carried out in May 2018 using the oceanographic research ship Néréis II (Station research
257 vessel of Banyuls-sur-Mer, TGIR French Oceanographic Fleet; Fig. 1C), with a regular
258 sampling of both surface and bottom seawater (down to 450 m below sea level; with *ex-situ*
259 water filtration) as well as shells (encompassing unidentified bivalves fragments, *Mytilus*
260 *galloprovincialis* and one gastropod *Turritella communis*) using a sediment grab.

261 3.2. Chemical composition of waters and shells

262 All the analytical steps have been performed at the Geosciences Environnement
263 Toulouse laboratory (France). First, the collected water samples were prepared in a clean
264 laboratory for chemical analyses including major (Mg, Ca, Na, K) and trace (Sr) element
265 concentrations as well as the anion (Cl, SO₄, Br) concentrations. Anions were determined by
266 anionic chromatography calibrated with an IAPSO seawater standard solution that was diluted
267 at different proportion with MQ-H₂O to cover the entire range of anion concentrations. The
268 IAPSO solution is a seawater standard solution provided by OSIL (UK) and certified for its
269 salinity. The major and trace elements (Ca, Mg, Na, K, Sr) concentrations were measured
270 using an Inductively Coupled Plasma optical emission spectrophotometer (ICP-OES), Horiba
271 Jobin Yvon Ultima 2. We used the IAPSO standard solution to calibrate the instruments
272 (Besson et al., 2014).

273 The shell samples were abraded, cleaned with Suprapur® acetic acid, and rinsed in
274 Milli-Q water to remove external impurities and periostracum. Smaller bivalves and
275 gastropods samples were entirely crushed with an agate mortar to get a powder, whereas only
276 the inner parts of the umbo (recording the entire life span of the organism) were microdrilled
277 for larger bivalve shells. For major element concentrations of molluscs, we dissolved 10 mg
278 of sample using double-distilled 1 M HNO₃ on a hot plate at 70°C. After evaporation of the
279 solution to dryness, it was dissolved in 2N HNO₃ and diluted 100 times with MQ-H₂O. The
280 solution was then analyzed for Mg, Ca, and Sr concentrations using an ICP-OES. The
281 dissolved elements are 1000 times more concentrated than their limit of detection (>10ppb).
282 The ICP-OES instrument was calibrated using a matrix-matched synthetic standard. One of
283 the calibration solutions was measured every 5 samples to check the instrumental drift during
284 the course of the analyses. The measurements achieved a precision of 2.3% for Ca, 2.8% for
285 Mg and 4.3% for Sr, in line with previous analytical precision on Ca-rich sample (Chavagnac
286 et al., 2005).

287 *3.3. Sr isotopic compositions*

288 To determine the ⁸⁷Sr/⁸⁶Sr ratios, 1 ml of filtered water was transferred to a Savillex
289 beaker prior to evaporation on a hot plate at 70 °C and the residue was dissolved in 0.5 ml of
290 2N HNO₃. For the shells, ~10 mg of powder was dissolved in a Savillex beaker with 2N
291 HNO₃ on a hot plate at 70°C and centrifuged to exclude any residual fraction. Strontium was
292 isolated from resulting solutions using Eichrom Sr-Spec column, a conventional elution
293 protocol (Pin et al., 2014). The isolated Sr fraction was dissolved with 2µl of 0.5N H₃PO₄
294 solution of which 1µl was loaded on a tungsten filament together with 1µl of Tantalum
295 activator. The Sr isotopic composition was measured using a Thermo Triton Plus thermal
296 ionization mass spectrometer. The ⁸⁷Sr/⁸⁶Sr ratios were defined as the average of 150
297 measurements of ion intensities following the static multi-collection mode normalized to

298 $^{86}\text{Sr}/^{88}\text{Sr} = 0.1194$. During the measurement period, the standard NBS 987 gave $^{87}\text{Sr}/^{86}\text{Sr}$
299 ratios of 0.710280 ± 0.000003 (2SD, standard deviations, $n = 5$). Over the 2018 year, the
300 measured NBS 987 isotopic compositions provided an external reproducibility of $0.710280 \pm$
301 0.000012 (2SD; $n = 26$). Regarding international seawater standards (IAPSO and NASS-6),
302 they were processed in the clean room laboratory in the same manner as for our samples for
303 Sr chemical isolation. The external $^{87}\text{Sr}/^{86}\text{Sr}$ reproducibilities are 0.709179 ± 0.000007 (2SD;
304 $n = 7$) for IAPSO and 0.709180 ± 0.000014 (2SD; $n = 5$) for NASS-6, which agree very well
305 with their certified values of 0.709179 ± 0.000005 and 0.709173 ± 0.000018 , respectively
306 (Neymark et al., 2014).

307 For the bivalve shells, we analyzed at repeated times the Sr isotopic composition of
308 the JcT-1 international standard (a giant clam *Tridacna gigas*) treated in the clean room as for
309 our shell samples. The measurements provide an external reproducibility of $0.709176 \pm$
310 0.000015 (2SD; $n = 11$) for a certified value of 0.709150 ± 0.000050 (Ohno and Hirata, 2007).
311 For comparison purposes between the $^{87}\text{Sr}/^{86}\text{Sr}$ ratios reported in this study and the literature
312 data (tables 1 and 2), the $^{87}\text{Sr}/^{86}\text{Sr}$ ratios were corrected by adjusting the measured NBS 987
313 standard reference value to a certified value of 0.710250.

314 3.4. Geochemical and Sr isotopic database

315 To better depict potential influences of groundwater in the Oualidia and Salses-
316 Leucate lagoons, we compiled a dataset of published major and trace elements concentrations
317 (Ca, Mg, Na, K, SO_4 , Cl, Br and Sr) and $^{87}\text{Sr}/^{86}\text{Sr}$ ratios corresponding to groundwaters
318 representative of Moroccan aquifers ($n = 112$) and the Corbières aquifer in France ($n = 141$)
319 (see supplementary data). In order to analyze the geochemical patterns at a worldwide scale,
320 we also compiled a dataset gathering 522 $^{87}\text{Sr}/^{86}\text{Sr}$ ratios and 571 Sr concentrations
321 representative of OOW and shelf waters (SW) (see supplementary data). As for our samples,

322 all $^{87}\text{Sr}/^{86}\text{Sr}$ ratios reported in the literature were corrected by adjusting their respective NBS
323 987 standard reference value to a certified value of 0.710250.

324

325 **4. Results**

326 *4.1. $^{87}\text{Sr}/^{86}\text{Sr}$ of waters and shells*

327 The $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of lagoon waters and shells are reported in Tables 1 and 2. First, at
328 the Tatakoto lagoon, waters and shells exhibit $^{87}\text{Sr}/^{86}\text{Sr}$ ratios at 0.709165 - 0.709171 and
329 0.709164 – 0.709182, respectively, which are both indistinguishable from the average OOW
330 value of 0.709172 (Figs. 1D and 2A). Similarly, seawater collected at different water depths
331 along the BSM transect presents a slight dispersion ranging from 0.709166 to 0.709176, but
332 still around the OOW value (Figs. 1C). The only sample deviating from this world value is
333 BA9 collected in the Banyuls-sur-Mer beach: it presents a low salinity of 20 and a $^{87}\text{Sr}/^{86}\text{Sr}$
334 ratio of 0.709206 slightly more radiogenic than the OOW range (Fig. 2A). Molluscs from the
335 BSM transect present Sr isotope compositions between 0.709164 and 0.709175, all identical
336 to those of surrounding water (Fig. 2A).

337 In the Salses-Leucate lagoon, the seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios ranging from 0.709155 and
338 0.709170 are typical of the OOW variability range. Contrastingly, three water samples (LEU
339 6, 7, 8) from the western part of the lagoon display values ranging from 0.708903 to 0.709132,
340 that is below the OOW variability range. However, these values are trending towards the
341 signatures of the Font Dame and Font Estramar discharges (LEU 10, 11) at 0.708618 and
342 0.708401, respectively (Fig. 1B). Again, these relatively low isotopic ratios occur along a
343 salinity drop from 37.4 to 11.8 (Fig. 2A). In this context, the least radiogenic Sr isotope
344 composition is measured for the bottom water sample (LEU 8) compared to the surface one
345 (LEU 7). Mollusc shells sampled in this lagoon present $^{87}\text{Sr}/^{86}\text{Sr}$ ratios varying between
346 0.709125 and 0.709177, being either similar to or less radiogenic than the OOW value (Table

347 2; Fig. 2A). Finally, it is worth mentioning that our isotopic results are within the variability
348 of those obtained for local seawater and groundwater sampled in 1998, 2002 and 2012 by
349 Ladouche et al. (2000), Ladouche and Dörfliger (2004) and Petelet-Giraud et al. (2016).

350 At the Oualidia lagoon, both seawater and mollusc shells display highly variable
351 $^{87}\text{Sr}/^{86}\text{Sr}$ ratios ranging from 0.708389 to 0.709168 and 0.708939 to 0.709179, respectively
352 (Tables 1 and 2; Figs. 1A and 2A). Downstream, close to Atlantic seawater incursions by the
353 inlets, the $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of waters stay within the OOW variability range, while they are less
354 and less radiogenic upstream between the OU12 and OU8 locations (Table 1, Figs. 1 and 2).
355 As in Salses-Leucate and Banyuls bay, this isotopic shift is accompanied by a salinity
356 decrease from 37.2 to 9.9 (Fig. 2A). However, the water sample OU6 deviates from this
357 general tendency: although close to the northern dam, it records a $^{87}\text{Sr}/^{86}\text{Sr}$ ratio typical of
358 OOW values (Fig. 1A). Finally, we note that most bivalve shells record $^{87}\text{Sr}/^{86}\text{Sr}$ ratios typical
359 of OOW values but different of surrounding water values (Fig. 2A, Table 2). Only the O4
360 sample (i.e., an endobenthic *Solen marginatus*) fits the general $^{87}\text{Sr}/^{86}\text{Sr}$ decrease of lagoonal
361 waters.

362 4.2. Geochemical composition of waters

363 All concentrations of anions and major and trace elements measured on the collected
364 waters are reported in Table 1. In order to determine the origin of the waters influencing the
365 $^{87}\text{Sr}/^{86}\text{Sr}$ of local seawater, only the most discriminating elements such as Sr, SO_4 , and Ca are
366 used here, plotted as function of salinity, and compared to published values of regional
367 groundwater aquifers (Fig. 2).

368 In the Pacific Tatakoto atoll, seawater has salinity and $^{87}\text{Sr}/^{86}\text{Sr}$ ratios typical of OOW
369 while they display lower Sr contents ($\sim 73.5 \mu\text{mol}\cdot\text{l}^{-1}$ against $88 \mu\text{mol}\cdot\text{l}^{-1}$ for OOW; Millero et
370 al. 2008). Slight enrichments in SO_4 ($30.8 \text{ mmol}\cdot\text{l}^{-1}$) and Ca ($10.3 \text{ mmol}\cdot\text{l}^{-1}$) are recorded but
371 still within the OOW variability range. These geochemical characteristics also concern most

372 water samples from the Mediterranean BSM transect, whatever the water depth, with
373 respective Sr, SO₄ and Ca concentrations of 72.2 – 74.5 μmol·l⁻¹, 30.9 – 32 mmol·l⁻¹, and 11.2
374 – 12.2 mmol·l⁻¹. The only exception is the beach sample BA9 which low salinity value of 26.4
375 is mirrored by low element concentrations compared to those of OOW (Table 1 and Fig. 2).

376 Euhaline waters from the inner parts of the Oualidia and Salses-Leucate lagoons are
377 also characterized by similar depletions in Sr and slight enrichments in Ca and SO₄ when
378 compared to the areas described above. However, the element concentrations evolve
379 differently in Salses Leucate and Oualidia while the salinity and ⁸⁷Sr/⁸⁶Sr are decreasing
380 landward. In Salses-Leucate, the Sr, SO₄ and Ca contents are dropping down to 17.2 μmol·l⁻¹,
381 3.8 mmol·l⁻¹ and 3.5 mmol·l⁻¹, respectively, close to the Font Dame and Font Estramar
382 discharges. In agreement with previous analyses of Ladouche et al. (2000) and Ladouche and
383 Dörfliger (2004), these low concentrations are similar to those prevailing at the scale of the
384 regional aquifer and the covariations between salinity, ⁸⁷Sr/⁸⁶Sr ratio and these elemental
385 concentrations are consistent with freshwater-seawater binary mixing. In Oualidia, the salinity
386 and ⁸⁷Sr/⁸⁶Sr decrease while Sr and Ca concentrations remain relatively high and stable (i.e.,
387 ~73 μmol·l⁻¹ and ~11.5 mmol·l⁻¹, respectively), except in the salt marshes where values are
388 much higher (i.e., 117 μmol·l⁻¹ and 20.5 mmol·l⁻¹, respectively). Contrastingly the SO₄
389 contents decrease with salinity but toward a brackish groundwater end-member with
390 anomalously high SO₄ concentrations (~15 mmol·l⁻¹). Oualidia geochemical patterns are
391 compatible with the high Sr, SO₄, and Ca contents recorded in some Moroccan groundwater
392 aquifers, and especially those related to the dissolution of Mesozoic carbonate and gypsum
393 levels (see Supplementary data).

394

395 **5. Discussion**

396 Studies with high precision $^{87}\text{Sr}/^{86}\text{Sr}$ ratio have confirmed that modern oligotrophic
397 oceanic waters (OOW) without terrestrial inputs have a quite homogeneous $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of
398 0.709172 with a variability range of only ± 0.000023 at the global scale (Andersson et al.,
399 1992; Winter et al., 1997; Huang and You, 2007; Huang et al., 2011; Mokadem et al., 2015;
400 Pearce et al., 2015; Trezzi et al., 2017; El Meknassi et al., 2018). This homogeneity is yet
401 regularly challenged for shelf domains as more radiogenic or unradiogenic terrestrial Sr
402 supplies from rivers or SGD can introduce a dispersion to the measured $^{87}\text{Sr}/^{86}\text{Sr}$ of 6×10^{-3} or
403 more (El Meknassi et al., 2018) (Fig. 3A). Nevertheless, this estimate mainly concerns
404 estuaries, fjords, bays or epeiric seas supplied by major river systems for which data are
405 relatively abundant (e.g., Ingram and Sloan, 1992; Jørgensen and Banoeng-Yakubo, 2001;
406 Négrel et al., 2005; Jørgensen et al., 2008; Wang and You, 2013; Eissa et al., 2016; Casse et
407 al., 2019). In this context, new Sr isotope data from open shelves and coastal lagoons become
408 crucial to get a better view of processes at play in neritic domains, especially those locations
409 deprived of river and delta influences. These areas represent 40 – 50% of the world coasts
410 (Dürr et al., 2011).

411

412 *5.1. Seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in oceanic and open shelf domains*

413 With an average Sr concentration of $87.4 \mu\text{mol}\cdot\text{l}^{-1}$ (de Villiers 1999), oceanic inflows
414 are the main dissolved Sr source to shelf domains and hemipelagic carbonate platforms. If
415 these oceanic inflows remain constant through time and sufficiently mixed with coastal waters,
416 locally homogeneous seawater and shell $^{87}\text{Sr}/^{86}\text{Sr}$ ratios close to OOW value of 0.709172
417 would be expected, even in semi-enclosed areas. Far from any terrestrial influence, this is
418 clearly illustrated in the isolated Tatakoto atoll where Pacific seawater is the main dissolved

419 Sr source, conveyed to the shelf by endo-upwelling circulation through the reef porosity
420 followed by surface currents (Rougerie, 1994). Rainwater infiltrations responsible for fossil
421 reef karstification and possibly volcanic basement weathering are important in the Tuamotu
422 archipelago (Guy et al., 1992; Waljeski, 2003). However, unradiogenic Sr supplied from local
423 SGD appears too limited to modify the OOW $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of studied waters and giant clams
424 in this Pacific atoll (Table 1 and 2).

425 In the nearshore domain exemplified by the 25 km-long Mediterranean BSM transect,
426 seawater and shells from surface down to ~450 mbsl display a homogeneous $^{87}\text{Sr}/^{86}\text{Sr}$ ratio
427 with an average value at 0.709171 ± 0.000008 (2SD; n=12) indistinguishable from the OOW
428 value (El Meknassi et al., 2018). In the absence of noticeable dissolved Sr inputs from rivers
429 or SGD, this implies that the Sr inputs resulting from boundary-exchange processes acting at
430 the sediment-water interface are not sufficient to modify the seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratio in these
431 well-mixed open shelf conditions. This contrasts with estuaries or deltas where dissolved Sr
432 released from suspended particulate dissolution may alter the $^{87}\text{Sr}/^{86}\text{Sr}$ of seawater in a few
433 months (Jones et al., 2014; Jeandel and Oelkers, 2015). Moreover, the shell data from the
434 BSM transect indicate that the Sr isotope composition of local seawater did not significantly
435 change over life spans of 2 to 3 years. However, these results do not preclude caution in
436 paleoenvironmental interpretation of shell $^{87}\text{Sr}/^{86}\text{Sr}$ ratios from the BSM coast and more
437 generally nearshore contexts. This is because SGD influence may strongly vary in time and
438 space along continental margins (Burnett et al., 2006), and especially in the Gulf of Lion
439 where SGD are very important (Rodellas et al., 2015).

440 In the foreshore zone, additional factors can impact the seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios. This
441 is illustrated by the BA9 water sample collected on the BSM sandy beach, characterized by a
442 more radiogenic value and a much lower salinity (i.e., 26.4) than the other samples. We
443 propose that this more radiogenic Sr value can be related to three factors typical of intertidal

444 zones: 1) mixing with rainwater whose regional $^{87}\text{Sr}/^{86}\text{Sr}$ ratios range from 0.708993 to
445 0.709225 (Khaska et al., 2013), 2) sparse groundwater supplies whose Sr concentrations and
446 radiogenic $^{87}\text{Sr}/^{86}\text{Sr}$ ratios mirror those from Quaternary, Pliocene, and Paleozoic aquifers (i.e.,
447 0.708142 to 0.713782; Petelet et al., 1998; Petelet-Giraud et al., 2016), and 3) important
448 dissolved Sr amounts released from suspended sandy particles continuously shaken in the surf
449 zone (Kalnejais et al., 2010; Jeandel, 2016; Fabre et al., 2019) with particles $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of
450 0.72232 – 0.72901 reflecting their metamorphic substratum (Brems et al., 2013). At broader
451 scale, similar local processes can explain part of isotopic deviations reported in the rare water
452 and shell data from open shelves (e.g., North Sea, Gulf of Guinea, South China Sea;
453 Jørgensen and Banoeng-Yakubo, 2001; Jørgensen et al., 2008; Huang et al., 2011; El
454 Meknassi et al., 2018) (Fig. 3). Moreover, marginal areas are transition zones where the
455 oceanic mixing is not instantaneous and may take several years according to littoral currents,
456 coastal physiography, and shelf width (Liu et al., 2019) (Fig. 3B). Terrestrial dissolved Sr
457 sources, even remote, could slightly impact local seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios through cumulative
458 effects. Thus, we advocate to reject any water and/or shell sample from beach or intertidal
459 zones to infer modern or past OOW $^{87}\text{Sr}/^{86}\text{Sr}$ ratios.

460

461 *5.2. $^{87}\text{Sr}/^{86}\text{Sr}$ patterns in restricted coastal domains*

462 Supplementing sparse literature data (e.g., Müller et al. 1990b; Beck et al. 2013; Shao
463 et al. 2018), the seawater and shell results from the Oualidia and Salses-Leucate lagoons show
464 clear evidence that restricted coastal domains, even without river influence, display very
465 distinct $^{87}\text{Sr}/^{86}\text{Sr}$ ratios depending on the respective contribution of oceanic and SGD influxes
466 (Fig. 1): 1) quite homogeneous OOW $^{87}\text{Sr}/^{86}\text{Sr}$ ratios except in close vicinity of SGD in
467 Salses-Leucate; or 2) variable $^{87}\text{Sr}/^{86}\text{Sr}$ ratios ranging from OOW values at oceanic inlets to
468 unradiogenic signals in the upstream parts of the Oualidia lagoon.

469 At broader scale, such Sr isotopic variabilities are also noticeable in epeiric seas. By
470 compiling literature data and excluding the anomalous $^{87}\text{Sr}/^{86}\text{Sr}$ ratios (i.e., samples from
471 beach or close to SGD) from Salses-Leucate, Banyuls-sur-Mer and Venice (Fig.4A), we
472 observe that seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios are typical of OOW values in western Mediterranean
473 Sea (i.e., Spain, France, Italy), whereas those from eastern domains like the Bannock Basin,
474 Aegean Sea, Marmara Sea and Black Sea decrease down to 0.70912 (this study; Müller et al.,
475 1990a; Clauer et al., 2000; Peckmann et al., 2001; Major et al., 2006; Beck et al., 2013;
476 Petelet-Giraud et al., 2016; Trezzi et al., 2017; Teichert et al., 2018). Highlighted here for the
477 first time, these regional $^{87}\text{Sr}/^{86}\text{Sr}$ variations are consistent with the isotopic variability of
478 Mediterranean water masses illustrated by the Nd isotopes (Tachikawa et al. 2004). Whether
479 local or supra-regional, spatial Sr isotopic variability is insightful to decipher the respective
480 influence of SGD and oceanic inputs on the Sr isotope budget of semi-enclosed areas.

481

482 *5.2.1. SGD influences*

483 With relatively high Sr concentrations (i.e., $2.9 \mu\text{mol}\cdot\text{l}^{-1}$) compared to river waters (i.e.,
484 0.5 to $1.2 \mu\text{mol}\cdot\text{l}^{-1}$), SGDs represent an unradiogenic Sr source that can account for 13 - 30%
485 of the global Sr ocean budget (Basu et al., 2001; Krabbenhöft et al., 2010; Beck et al., 2013).
486 Depending on their geochemical and hydrodynamical characteristics related to the age and
487 nature of weathered rocks but also oceanic intrusions, meteoric recharges, and water residence
488 time in the aquifers (e.g., Négrel et al., 2003; Fadili et al., 2016, 2015; Santoni et al., 2016),
489 their impact on the elemental and isotopic Sr budgets are, however, highly variable in coastal
490 domains (e.g., Beck et al., 2013; Trezzi et al., 2017; Shao et al., 2018). This is illustrated by
491 comparing the Salses-Leucate and Oualidia lagoons.

492 In the Salses-Leucate lagoon, the low salinities and the low Sr, SO_4 and Ca
493 concentrations of Font Estramar and Font Dame discharges show that these groundwaters are

494 typical of meteoric freshwaters percolating through the local karstic aquifer (Fig. 2B,C,D). To
 495 better constrain the origin of the collected seawaters in the lagoon, we applied the isotopic
 496 mixing equation (Albarède, 1996) to geochemical data of regional end members presented in
 497 Table 3:

$$498 \quad {}^{87}\text{Sr}/{}^{86}\text{Sr}_{(\text{mix})} = \left(\frac{{}^{87}\text{Sr}/{}^{86}\text{Sr}_{(\text{SW})} \times [\text{Sr}]_{(\text{SW})} \times \%q_{(\text{SW})}}{[\text{Sr}]_{(\text{mix})}} \right) + \left(\frac{{}^{87}\text{Sr}/{}^{86}\text{Sr}_{(\text{GW})} \times [\text{Sr}]_{(\text{GW})} \times \%q_{(\text{GW})}}{[\text{Sr}]_{(\text{mix})}} \right)$$

499 ,where the mixed water ${}^{87}\text{Sr}/{}^{86}\text{Sr}_{(\text{mix})}$ ratio varies between the local seawater (SW) and
 500 groundwater (GW) end members of given Sr concentrations and ${}^{87}\text{Sr}/{}^{86}\text{Sr}$ ratios, using mixed
 501 water Sr concentrations $[\text{Sr}]_{(\text{mix})}$ calculated from cumulative 10% increments of GW to SW.
 502 The low ${}^{87}\text{Sr}/{}^{86}\text{Sr}$ ratios of Font Estramar and Font Dame discharges correspond to the lowest
 503 isotopic values of regional groundwaters and indicate a preferential dissolution of Jurassic and
 504 Cretaceous carbonates characterized by low ${}^{87}\text{Sr}/{}^{86}\text{Sr}$ ratios compared to the Pliocene and
 505 Quaternary silicates ones (Fig. 4A; Petelet et al., 1998; Ladouche et al. 2000; Aquilina et al.,
 506 2002; Ladouche and Dörfliger 2004; Khaska et al., 2013; Petelet-Giraud et al., 2016). Given
 507 the low Sr concentrations of Jurassic and Cretaceous GW endmembers (Table 3; Fig. 2B), our
 508 results suggest that the SGD contribution to the lagoon Sr budget is broadly less than 15%
 509 despite local SGD fluxes exceeding $5.55 \text{ m}^3 \cdot \text{s}^{-1}$ (Fig. 4B; Ladagnous and Le Bec, 1997;
 510 Stieglitz et al., 2013).

511 In Oualidia, the dissolved Sr supply of SGD with much lower estimated fluxes of $0.2 -$
 512 $1.6 \text{ m}^3 \cdot \text{s}^{-1}$ (Hilmi et al., 2005; Fakir et al. 2019) markedly modifies the seawater ${}^{87}\text{Sr}/{}^{86}\text{Sr}$
 513 ratios within the upstream parts of the lagoon (Figs. 1A). These induced modifications can be
 514 related to considerable unradiogenic Sr inputs while maintaining constant dissolved Sr
 515 concentrations (Fig. 2A,B). Indeed, groundwaters from the Coastal Sahel of Oualidia exhibit
 516 variable but generally high Sr and Ca concentrations reaching 2 to 2.5 times the OOW value
 517 as well as high SO_4 amounts (Fakir et al., 2002; Hilmi et al., 2005; Fig. 2B,C,D). Even if
 518 oceanic water intrusions are not excluded in the coastal aquifer, specific Ca and SO_4 excesses

519 indicate that the higher Sr contents would result from a preferential dissolution of Sr rich
520 gypsum and celestite (SrSO_4) present in the Jurassic (and Cretaceous) limestones (Fakir et al.,
521 2002; Hilmi et al., 2005; Kaid Rassou et al., 2005; Bouchaou et al., 2017). This hypothesis is
522 in agreement with the unradiogenic $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of lagoon waters (i.e., down to 0.708389)
523 whose values are close to Jurassic aquifers of Morocco (Bouchaou et al., 2017; Vinson et al.,
524 2013) (Fig. 4A). We estimate that groundwater fluxes dissolving Mesozoic sulphates spatially
525 contribute from ~10 to 60 % to the lagoon Sr budget, the higher Sr concentrations being
526 related to high evaporation rates in the salt marshes (Fig. 4A). This Sr supply markedly varied
527 in a few years, as shown by the isotopic scatter between local seawater and shell data at a
528 same location or shells sampled in 2006 (El Meknassi et al., 2018) and 2018 (this study) (Fig.
529 2A). In contrast, the low $^{87}\text{Sr}/^{86}\text{Sr}$ ratio recorded by one endobenthic *Solen marginatus*
530 suggests that groundwater influence remained more constant in the interstitial waters at 80 cm
531 sediment depth (i.e., the maximal burial depth of this species) than in the lagoon waters.

532 With Sr concentrations up to $150 \mu\text{mol}\cdot\text{l}^{-1}$ leading to local contributions of 60% to the
533 Sr budget of lagoons, our results call to reconsider the importance of SGD in link to the
534 dissolution of evaporitic deposits. In association with geothermal brines, their impact on the
535 global ocean Sr budget still remains difficult to appraise but could be regionally important
536 given the proximity of Messinian or Triassic evaporites such as those of Mediterranean
537 margins (e.g., Stein et al., 2000). Considering their elevated Sr concentrations, it is also
538 important to note that, contrarily to river-dominated contexts requiring considerable
539 freshwater inputs to alter the coastal seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratio, SGD may quickly change this
540 isotopic ratio in euhaline conditions. Indeed, most major world river systems produce
541 measurable effects on seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios when salinity drops below an average
542 threshold of 12 (Bryant et al., 1995). Only ~15, 10 and 5 % of the rivers have significant
543 effects at salinities above 20, 25 and 30, respectively. In contrast, significant $^{87}\text{Sr}/^{86}\text{Sr}$ shifts

544 occur at salinities close to 32–35 in the studied lagoons (Fig. 2A). From a paleoenvironmental
545 point of view, these results call to be very careful in any paleosalinity deduction from coastal
546 fossil $^{87}\text{Sr}/^{86}\text{Sr}$ data, especially since saline SGD are difficult to detect from biosedimentary
547 facies or seismic data compared with freshwater SGD (Lecher and Mackey, 2018; Goff, 2019).

548

549 *5.3. Influence of water mass restriction*

550 From small lagoons to larger epeiric areas like Mediterranean and Baltic seas, oceanic
551 mixing is essential to buffer regional SGD and river effects and maintain homogeneous
552 seawater $^{87}\text{Sr}/^{86}\text{Sr}$ patterns typical of OOW. However, water mass restriction imposed by
553 geographic barriers and water column stratification may be highly variable from one domain
554 to another one. In lagoons, the local OOW renewal is more or less perennial and important
555 depending on their leaky, choked or restricted geomorphology (Kjerfve and Magill, 1989;
556 Umgiesser et al., 2014). Basically, restricted and leaky lagoons are parallel to the coast and
557 present, respectively, few to numerous inlets that allow an efficient oceanic water turnover
558 (e.g. every one to a few days) driven by tidal dynamics and/or wind forcing (Kjerfve and
559 Magill, 1989; Umgiesser et al., 2014). These characteristics contribute therefore to
560 homogenize seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratio close to the OOW one like in the Salses-Leucate or
561 Venice lagoons (Beck et al. 2013) (Fig. 1). To the extreme, choked lagoons are characterized
562 by only one or two oceanic inlets with a dominant wind forcing implying several days to
563 several months for a total water renewal in the lagoon (Kjerfve and Magill, 1989; Umgiesser
564 et al., 2014). This situation applies to the Oualidia lagoon, with the exception that tides are an
565 important factor in the local hydrodynamics (Hilmi et al. 2017). Based on numerical
566 modelling of tidal currents at Oualidia, the time necessary for full lagoon water renewal was
567 estimated to be one or a few days in close vicinity of oceanic inlet increasing up to 15 to 30
568 days in the upstream channel and salt marshes, respectively (Hilmi et al., 2005, 2009, 2017).

569 As a result, the duality and rhythmicity between SGD discharges and seawater tidal influxes
570 explain the $^{87}\text{Sr}/^{86}\text{Sr}$ gradient observed in the Oualidia lagoon from the oceanic inlet towards
571 the interior of the lagoon.

572 In epeiric seas, similar water mass restriction processes appear relevant to explain the
573 variability of seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios. The Mediterranean basin is very interesting in this
574 respect as SGD with elevated Sr concentrations ($5 - 12 \mu\text{mol}\cdot\text{l}^{-1}$) and low $^{87}\text{Sr}/^{86}\text{Sr}$ ratios (i.e.,
575 $0.7078 - 0.7080$) prevail all along the northern Mediterranean margin (Rodellas et al., 2015;
576 Trezzi et al. 2017). However, only the eastern Mediterranean waters are significantly affected
577 by these unradiogenic terrestrial Sr inputs (Fig. 3A). We propose that this spatial
578 heterogeneity can be due to an efficient buffering of SGD in the western Mediterranean areas
579 due to Atlantic OOW inflows from the strait of Gibraltar. Indeed, the renewal time of
580 intermediate and deep waters is estimated to be 2.5 to 5 times faster in the western
581 Mediterranean basin than it is in the eastern basin owing to the Sicily Strait (Tanhua et al.,
582 2013).

583 By summarizing, we identify significant deviations from the OOW $^{87}\text{Sr}/^{86}\text{Sr}$ ratio in
584 various marine contexts where the interplay between oceanic and terrestrial Sr inputs is a key
585 parameter. This relative variability attests that, depending on their isolation, shelf domains
586 may be regarded as dynamic and transitory Sr sub-reservoirs not always representative of the
587 global ocean reservoir for which the modern Sr residence time is estimated around 2.5 Myr
588 (Hodell et al., 1990). This spatial decoupling in the Sr budget was suggested by de Villiers
589 (1999) for the upper ocean by estimating partial Sr residence times of only 5800–700,000
590 years in the upper 400 m of the water column due to a group of celestite (SrSO_4)-secreting
591 radiolarians (i.e., Acantharia). In semi-enclosed coastal domains with transient dynamics like
592 lagoons or large epeiric seas, we expect that the partial residence time (*sensu* Lin and Liu,
593 2019) could be in the same order of magnitude as water renewal time in marginal basins (i.e.,

594 one day to half a month for studied lagoons up to 30–130 years, 2–40 years, and 5–625 years
595 in the Mediterranean, Baltic and Black Sea basins, respectively) (Lee et al., 2002; Stieglitz et
596 al., 2017; Tanhua et al., 2013; Omstedt et al., 2014; Hilmi et al., 2017).

597 Finally, it worth noting that the Sr concentrations measured in the studied marine areas
598 are relatively low (i.e., 60 – 70 $\mu\text{mol}\cdot\text{l}^{-1}$) compared with those of OOW (i.e., 80 – 90 $\mu\text{mol}\cdot\text{l}^{-1}$)
599 (Fig. 2B). However, these results are in agreement with the large variability of coastal Sr
600 concentrations (i.e., $\pm 20 \mu\text{mol}\cdot\text{l}^{-1}$) reported in worldwide coastal contexts (Fig. 5), sometimes
601 with important annual fluctuations and local heterogeneities in euhaline waters (Ladouche et
602 al., 2000; Brunskill et al., 2003; Elsdon and Gillanders, 2006). This suggests that the slightly
603 non-conservative behaviour of Sr in seawater could be locally enhanced in some coastal and
604 hemipelagic domains. Whether perennial or temporary, the cause of Sr depletions in euhaline
605 waters – already reported the Salses-Leucate lagoon (Ladouche et al. 2000) but also
606 sporadically observed along the Panama coast, in the Venice lagoon (Italy) or in the Gulf of
607 Papua (Papua New Guinea) (Brunskill et al. 2003; Beck et al. 2013) - remains obscure. We
608 hypothesize that, in some water mass restriction contexts, three factors could be involved and
609 combined: 1) binding of Sr at the surface of sedimentary particles depending on grain size,
610 iron and manganese oxides, and organic matter (Takada et al. 2014); 2) fast biological
611 adsorption rates by specific organisms (i.e., see Bowen 1956, de Villiers 1999); or 3)
612 influences of saline SGD (i.e., similar to superficial porewaters) slightly depleted in Sr and
613 with isotopic ratios close to the one of seawater (e.g., Kastner et al. 1990).

614

615 **Conclusion**

616 In this study, we analyzed and compared the $^{87}\text{Sr}/^{86}\text{Sr}$ ratios and the major and trace
617 element concentrations of seawaters and shells from marginal and hemipelagic contexts with
618 different degrees of water mass restriction. Homogeneous $^{87}\text{Sr}/^{86}\text{Sr}$ ratios typical of OOW (i.e.,

619 0.709172 ± 0.000023) are recorded in the Tatakoto atoll and along the BSM transect. This
620 suggests that, in open shelf context without any river inputs, alternative Sr source like SGD or
621 particulate dissolution are too limited and likely buffered by oceanic Sr inputs. This comforts
622 the use of carbonate fossils from these domains to infer past OOW compositions. However,
623 slight isotopic shifts may be observed in the foreshore area (e.g., BSM beach) that we
624 interpret as resulting from rainwater mixing, local groundwater discharges or particle
625 dissolution in the surf zone. Elevated coastal water retention times could also account for
626 sporadic anomalous $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in open shelves. In semi-enclosed domains, we report
627 variable Sr isotope patterns between the studied karstic lagoons, with homogeneous seawater
628 $^{87}\text{Sr}/^{86}\text{Sr}$ ratios typical of OOW (i.e., 0.709155 to 0.709170) in the Salses-Leucate lagoon
629 (except close to groundwater discharges where the values drop to 0.708903) and an important
630 $^{87}\text{Sr}/^{86}\text{Sr}$ gradient from OOW values close to the Atlantic inlet to less radiogenic values of
631 0.707957 in the upstream parts of the Oualidia lagoon. We relate these differences to two
632 main factors. First, we note that, despite lower fluxes, the Oualidia lagoon is supplied by SGD
633 with unradiogenic $^{87}\text{Sr}/^{86}\text{Sr}$ ratios and very high Sr concentrations (up to 150 $\mu\text{mol}\cdot\text{l}^{-1}$) linked
634 to the dissolution of Mesozoic evaporites. These inputs lead to maximal contributions of 60%
635 to the local Sr budget. The second parameter concerns the leaky, restricted or choked
636 morphology of lagoons which controls the oceanic Sr inputs and thus buffers more or less any
637 terrestrial Sr inputs through water mass homogenization. These two parameters appear also
638 relevant to explain the $^{87}\text{Sr}/^{86}\text{Sr}$ gradient prevailing from west to east in the Mediterranean
639 basin.

640

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646 reviewers for their valuable remarks which considerably improved the manuscript.

647

648 **Figure Captions**

649

650 Fig. 1: Location and $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of seawater and groundwater samples collected in the four
651 studied sites: Oualidia lagoon (A), Salses-Leucate lagoon (B), Banyuls-sur-Mer (BSM)
652 transect (C), Tatakoto atoll (D). Dashed lines indicate groundwater discharges.

653

654 Fig. 2: $^{87}\text{Sr}/^{86}\text{Sr}$, Sr, Ca, and SO_4 concentrations of waters as a function of salinity. (A) For a
655 same sampling position with a given salinity, the $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of living mollusc shells are
656 compared to $^{87}\text{Sr}/^{86}\text{Sr}$ ratios of ambient seawater as well as oysters sampled in 2006 by El
657 Meknassi et al. (2018) (see Table 2 for samples correspondence). Additional literature data
658 from Salses-Leucate (i.e., water sampled in 1998, 2002 and 2012) are represented by squares
659 with white contours (Ladouche et al. 2000; Ladouche and Dörfliger 2004; Petelet-Giraud et al.
660 2016). The isotopic range and the average elemental concentrations of OOW are from El
661 Meknassi et al. (2018) and Millero et al., (2008), respectively. The salinity threshold defining
662 significant influence of river water (RW) inputs on seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratios is from Bryant et
663 al. (1995). Seawater and SGD values from the Oualidia and Salses-Leucates lagoons are
664 compared to groundwater data available at the scale of Moroccan aquifers (Fakir et al., 2002;
665 Kaid Rassou et al., 2005; Vinson et al. 2013; Fadili et al., 2015; Bouchaou et al., 2017) and
666 the Corbières aquifer (southern France) (Petelet et al., 1998; Aquilina et al., 2002; Khaska et
667 al., 2013; Petelet-Giraud et al., 2016). The analytical uncertainty bars are smaller than the
668 width of symbols.

669

670 Fig. 3: (A) Worldwide distribution of published and new $^{87}\text{Sr}/^{86}\text{Sr}$ ratios for euhaline to
671 brackish shelf waters and oligotrophic oceanic waters (OOW). If locally variable, the
672 maximal and minimal $^{87}\text{Sr}/^{86}\text{Sr}$ ratios are represented by two colours. Box-and-whisker plot
673 corresponds to the OOW $^{87}\text{Sr}/^{86}\text{Sr}$ variability range published by El Meknassi et al. (2018).
674 Literature data from Müller et al. (1990a, 1990b), Andersson et al. (1992), Ingram and Sloan
675 (1992), Winter et al. (1997), Barker et al. (1998), Israelson and Buchardt (1999), Clauer et al.
676 (2000), Ladouche et al. (2000), Jørgensen and Banoeng-Yakubo (2001), Peckmann et al.
677 (2001), Wang et al. (2001), Négrel et al. (2005), Major et al. (2006), Huang and You (2007),
678 Sharma et al. (2017), Xu and Marcantonio (2017), Jørgensen et al. (2008), Martin and moore
679 (2008), Huang et al. (2011), Patra et al. (2012), Rahaman and Singh (2012), Beck et al. (2013),
680 Uddin et al. (2013), Wang and You (2013), Jones et al. (2014), Mokadem et al. (2015), Pearce
681 et al. (2015), Eissa et al. (2016), Petelet-Giraud et al. (2016), Trezzi et al. (2017), Shao et al.
682 (2018), Teichert et al. (2018), Casse et al. (2019), Danish et al. (2020).

683

684 Fig. 4: $^{87}\text{Sr}/^{86}\text{Sr}$ vs. $1/\text{Sr}$ diagram for the Salses-Leucate (A) and Oualidia (B) lagoon waters.
685 The data are compared to mixing trends between local seawater and different groundwaters
686 (GW) from Moroccan (A) and Corbières aquifers (B). End members data are presented in
687 Table 3. The mixing curves show the $^{87}\text{Sr}/^{86}\text{Sr}$ ratios for each 10 % increment. The analytical
688 uncertainty bars are smaller than the width of symbols.

689

690 Fig. 5: Sr concentration of shelf (SW) and oligotrophic oceanic (OOW) waters as a function
691 of salinity. The OOW data are from de Villiers (1999) and the SW data are compiled from
692 various sources (see supplementary data). The red and grey lines represent the linear
693 regression and the 95% confidence interval of Sr concentration values, respectively.

694

695 Table 1: $^{87}\text{Sr}/^{86}\text{Sr}$ and elemental concentrations of surface and bottom seawater from the
696 studied sites.

697

698 Table 2: $^{87}\text{Sr}/^{86}\text{Sr}$ and elemental concentrations of studied shells.

699

700 Table 3: Average, minimal and maximal $^{87}\text{Sr}/^{86}\text{Sr}$ and Sr concentrations of seawater (SW) and
701 groundwater (GW) end members used for mixing model of Fig. 4. GW data refer to waters
702 percolating through different aquifers of the Corbières region (South France) and Morocco.
703 As GW $^{87}\text{Sr}/^{86}\text{Sr}$ and Sr concentration data are missing or sparse, respectively, for the Coastal
704 Sahel of Oualidia, we used GW data from Moroccan aquifers with host rocks of the same age
705 and lithology (see supplementary data).

706

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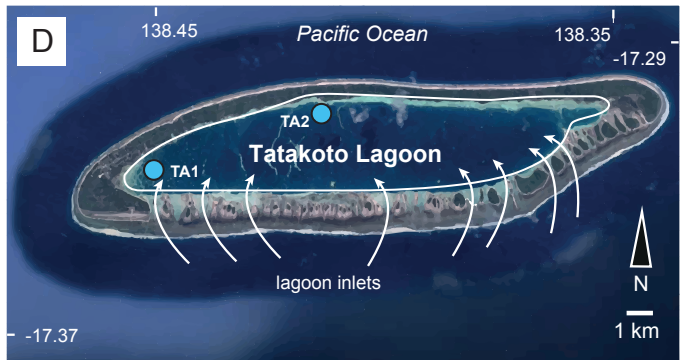
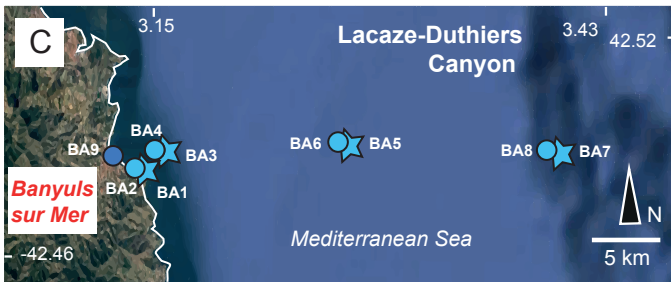
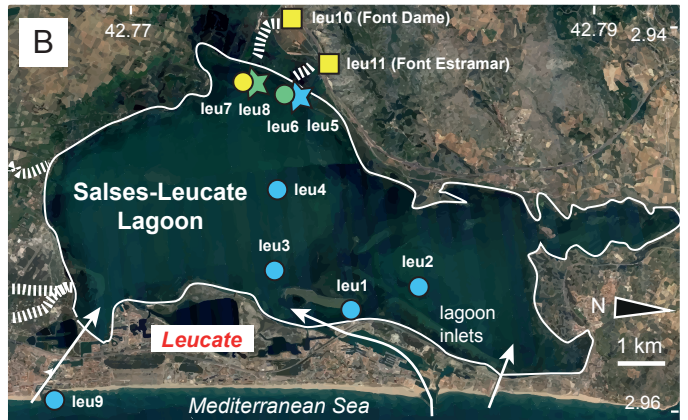
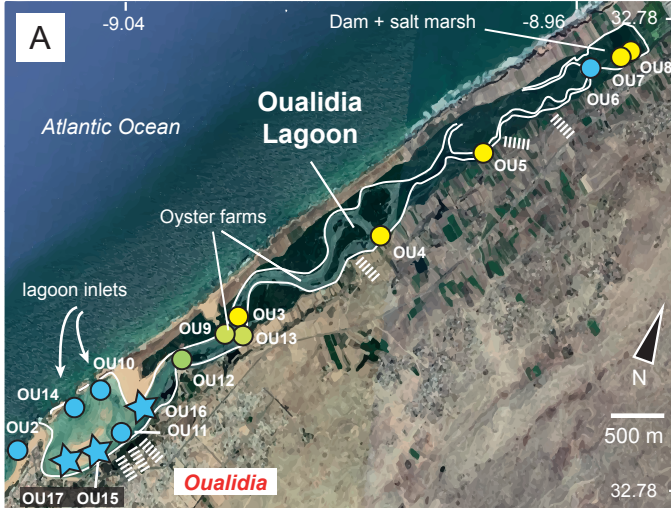
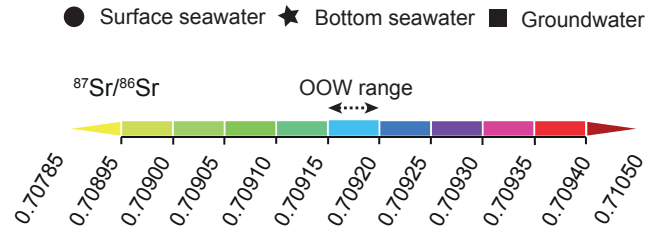
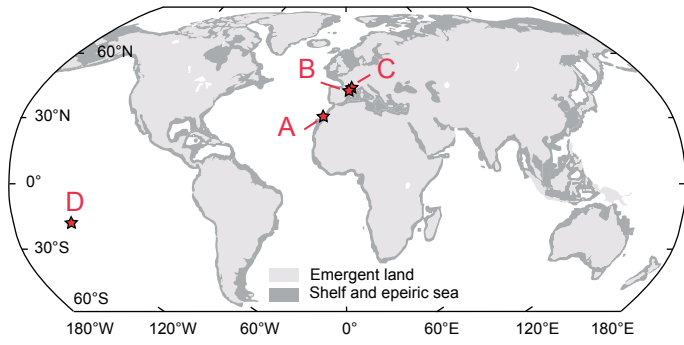
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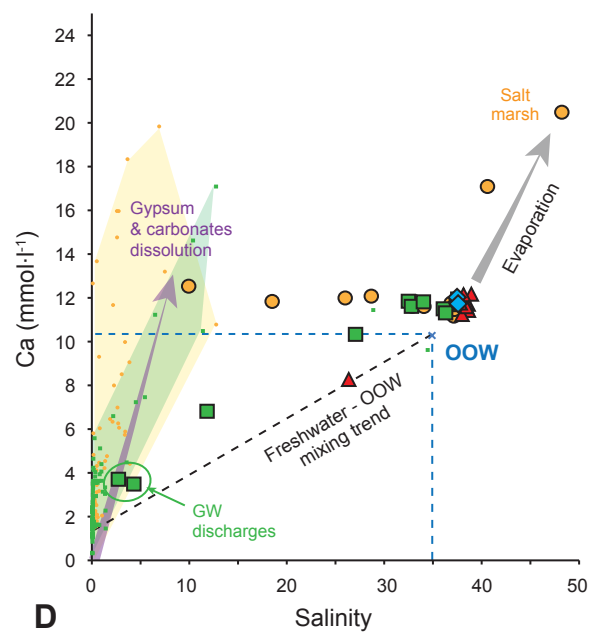
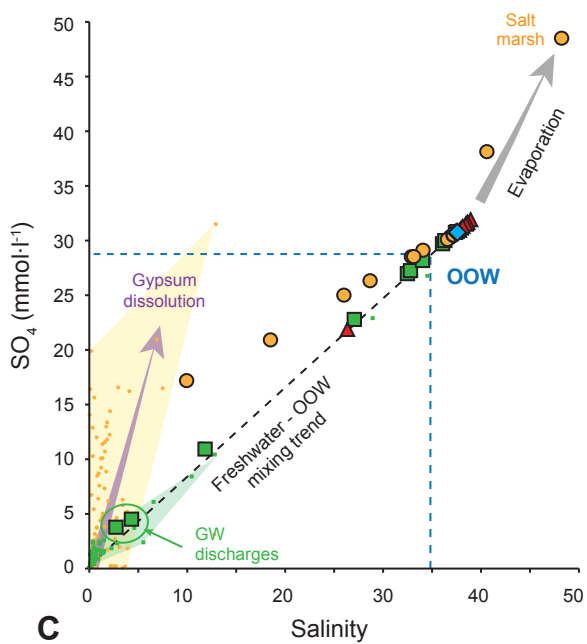
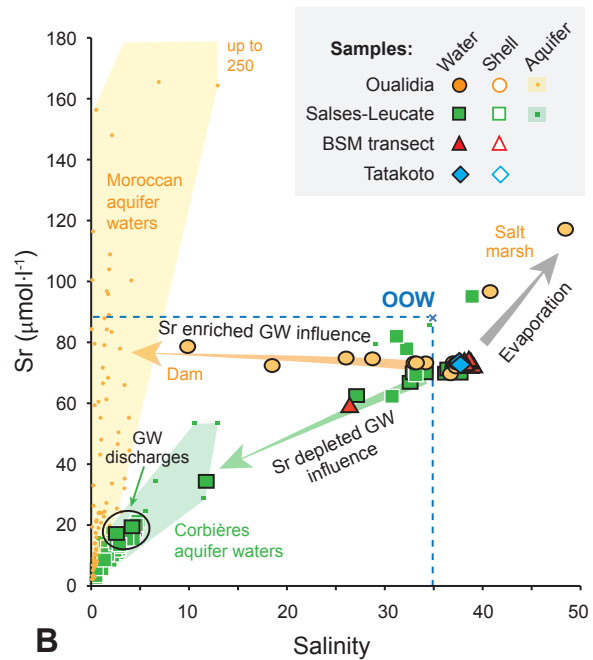
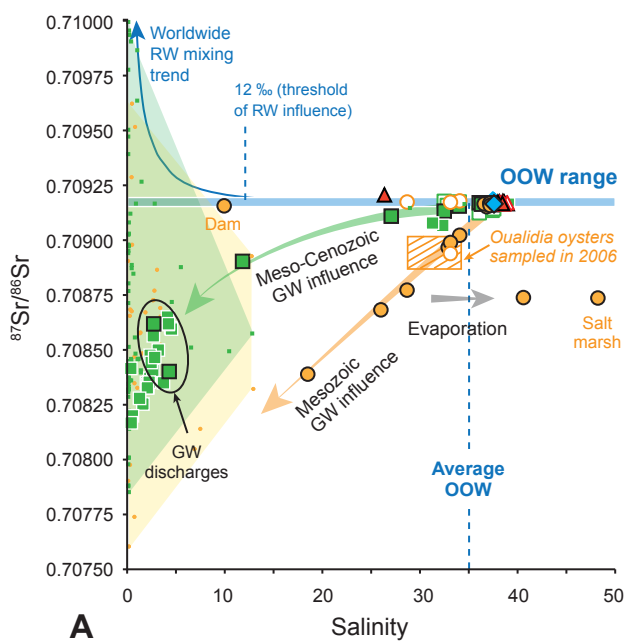
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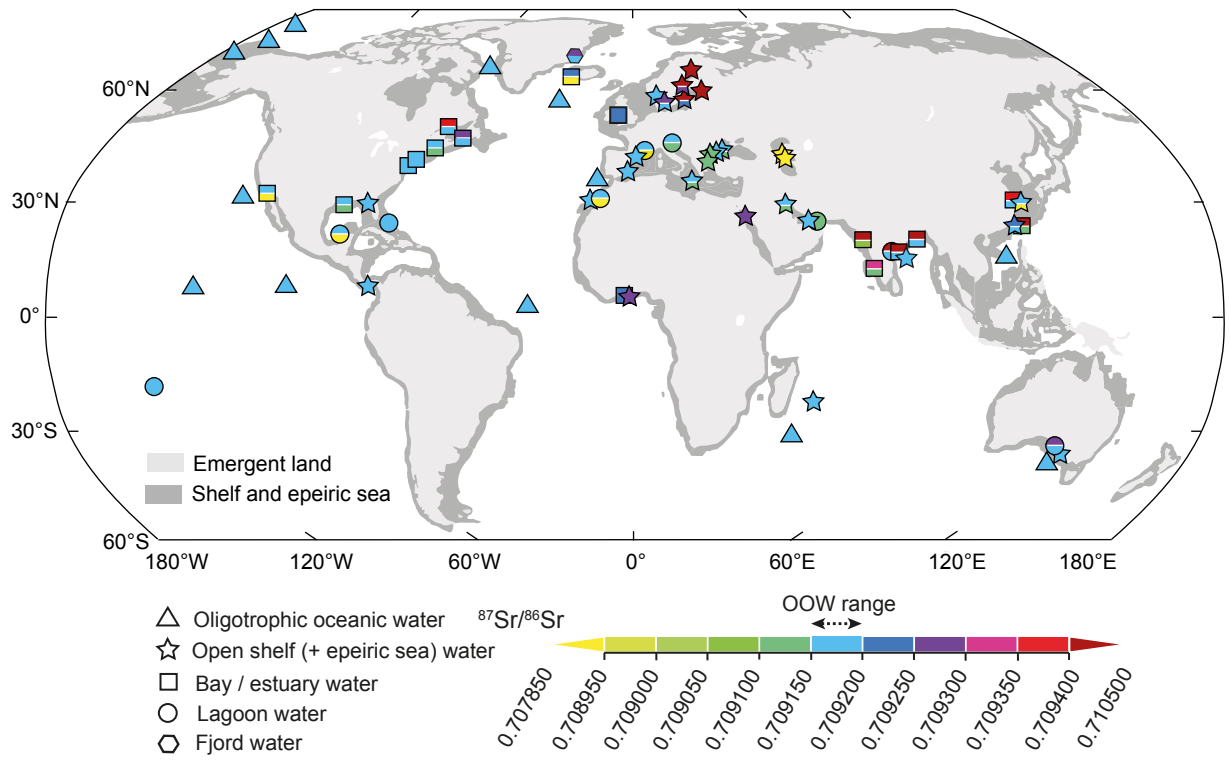
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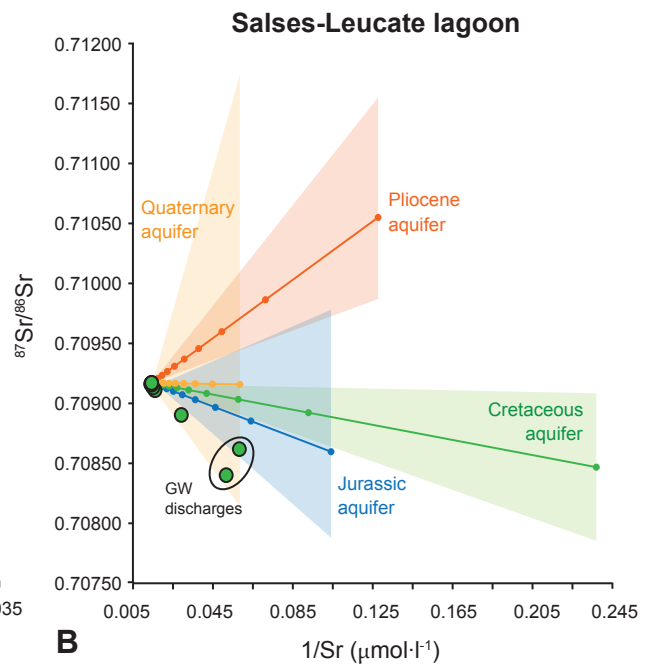
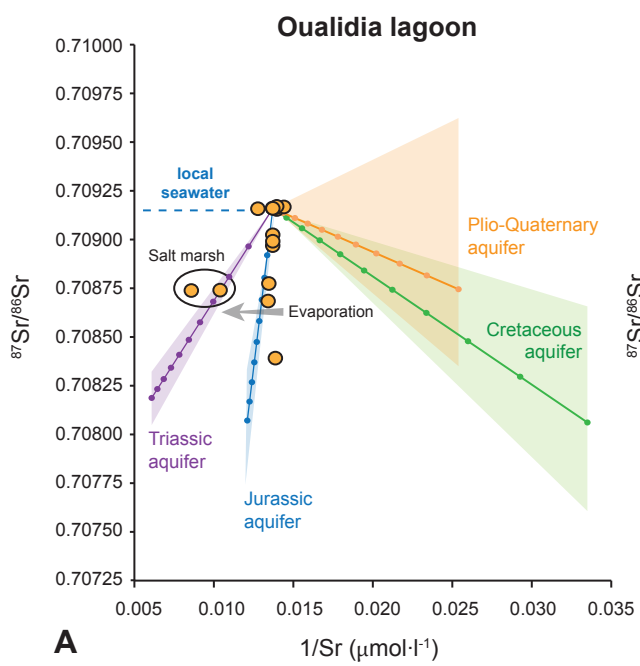
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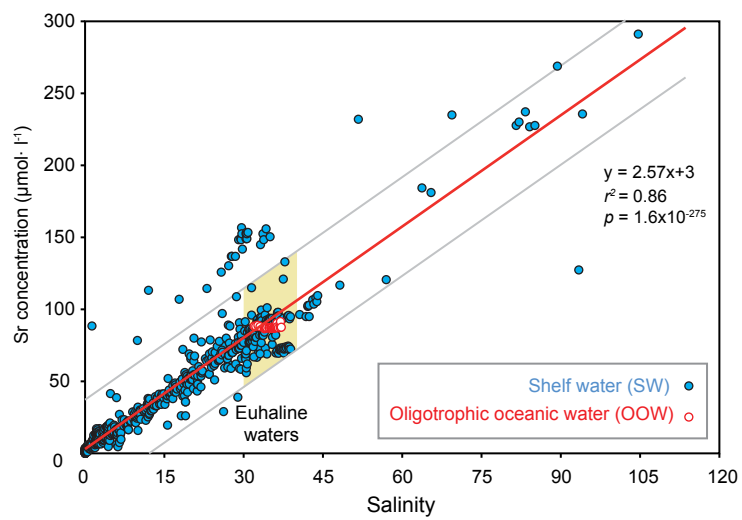
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Studied site	Sample	Water type (environment)	Water depth (mbsl)	Salinity	Ca mmol·l ⁻¹	Mg mmol·l ⁻¹	Na mmol·l ⁻¹	K mmol·l ⁻¹	SO ₄ mmol·l ⁻¹	Cl mmol·l ⁻¹	Li μmol·l ⁻¹	Br μmol·l ⁻¹	Sr μmol·l ⁻¹	⁸⁷ Sr/ ⁸⁶ Sr	2 std dev
Oualidia lagoon, Morocco	OU2	Atlantic seawater (beach)	0	37.2	11.47	59.23	567	9.97	30.51	581	48.72	865	73.0	0.709158	0.000004
	OU3	Lagoon water	0	28.7	12.08	47.85	453	7.83	26.33	448	36.26	667	74.4	0.708772	0.000004
	OU4	Lagoon water	0	26.0	11.99	42.47	399	6.96	25.01	406	31.53	597	74.6	0.708682	0.000004
	OU5	Lagoon water	0	18.5	11.84	30.88	276	4.85	20.92	289	23.17	420	72.2	0.708389	0.000004
	OU6	Lagoon water (close to the dam)	0	9.9	12.53	18.48	138	2.60	17.19	155	15.85	223	78.4	0.709156	0.000004
	OU7	Lagoon water (salt marsh)	0	40.6	17.09	71.32	653	11.55	38.13	634	54.53	1012	96.4	0.708738	0.000004
	OU8	Lagoon water (salt marsh)	0	48.2	20.49	92.47	753	15.36	48.51	753	80.57	1302	116.8	0.708736	0.000004
	OU9	Lagoon water	0	33.1	11.83	53.14	535	9.37	28.53	517	42.62	768	73.0	0.708989	0.000004
	OU10	Lagoon water	0	37.1	11.16	57.34	586	10.08	30.49	580	47.44	862	71.7	0.709168	0.000004
	OU11	Lagoon water	0	36.6	11.38	58.32	580	10.09	30.10	571	45.66	862	69.5	0.709165	0.000004
	OU12	Lagoon water	0	34.1	11.60	53.28	542	9.34	29.12	532	43.26	786	73.0	0.709022	0.000004
	OU13	Lagoon water	0	32.9	11.61	52.32	524	9.04	28.53	514	41.77	762	73.0	0.708967	0.000004
	OU14	Lagoon water	0	37.0	11.54	58.44	591	10.41	30.38	578	48.10	856	72.1	0.709162	0.000004
	OU15	Lagoon water	3	37.1	11.34	57.23	598	10.20	30.47	579	49.57	855	71.8	0.709164	0.000004
	OU16	Lagoon water	3	36.8	11.51	57.00	601	10.22	30.26	575	46.99	846	71.7	0.709152	0.000004
	OU17	Lagoon water	3	36.9	11.76	58.65	600	10.26	30.35	576	47.06	833	73.1	0.709159	0.000004
	Salses-Leucate lagoon, France	LEU9	Mediterranean seawater (beach)	0	37.6	11.95	61.04	598	10.71	30.82	586	46.71	871	69.7	0.709170
LEU1		Lagoon water (mussel farm)	0.5	36.3	11.31	57.40	577	10.24	30.00	567	45.72	848	71.0	0.709164	0.000004
LEU2		Lagoon water	0	36.1	11.50	57.25	579	10.39	29.71	563	46.54	849	69.7	0.709170	0.000004
LEU3		Lagoon water	0	37.4	11.69	59.52	593	10.61	30.75	584	47.81	880	71.9	0.709161	0.000004
LEU4		Lagoon water	0	34.0	11.82	56.83	576	9.95	28.17	531	42.88	786	69.8	0.709155	0.000004
LEU5		Lagoon water	1.5	32.8	11.61	54.76	553	9.65	27.25	512	39.48	762	69.8	0.709169	0.000004
LEU6		Lagoon water	0	32.5	11.85	54.75	556	9.67	26.97	507	41.36	746	66.7	0.709132	0.000004
LEU7		Lagoon water	0	11.8	6.82	20.23	186	3.25	10.93	185	11.42	235	34.2	0.708903	0.000004
LEU8		Lagoon water	2.5	27.1	10.33	44.94	454	7.73	22.79	422	30.55	618	62.3	0.709109	0.000004
LEU11		Groundwater (Font Estramar discharge)	0	4.3	3.49	6.83	39	1.07	4.52	67	4.78	86	19.3	0.708401	0.000004
LEU10		Groundwater (Font Dame discharge)	0	2.7	3.71	4.37	9	0.62	3.78	42	2.77	51	17.2	0.708618	0.000004
Banyuls-sur-Mer transect, France	BA1	Mediterranean seawater (open shelf)	22	38.1	12.15	60.73	622	10.57	31.29	595	45.09	894	73.1	0.709174	0.000004
	BA2	Mediterranean seawater (open shelf)	0	37.8	11.76	58.50	602	10.25	30.94	590	48.45	884	72.7	0.709167	0.000004
	BA3	Mediterranean seawater (open shelf)	40	38.4	11.60	59.69	588	10.46	31.49	600	47.76	913	74.5	0.709167	0.000005
	BA4	Mediterranean seawater (open shelf)	0	38.0	11.24	59.91	586	10.33	31.14	593	45.55	893	74.0	0.709173	0.000004
	BA5	Mediterranean seawater (open shelf)	140	38.6	11.43	59.52	579	10.40	31.66	603	50.66	900	73.9	0.709170	0.000004
	BA6	Mediterranean seawater (open shelf)	0	38.5	11.75	60.35	594	10.60	31.58	601	49.11	914	73.6	0.709176	0.000004
	BA7	Mediterranean seawater (open shelf)	510	38.9	12.17	63.26	615	11.19	31.91	608	49.62	904	72.4	0.709166	0.000004
	BA8	Mediterranean seawater (open shelf)	0	38.6	11.71	60.22	612	10.75	31.64	603	47.38	911	72.2	0.709174	0.000004
	BA9	Mediterranean seawater (beach)	0	26.4	8.27	42.21	432	7.40	21.87	412	30.34	605	59.3	0.709206	0.000004
Tatakoto lagoon, French Polynesia	T1	Lagoon water	2	37.6	11.73	61.81	599	10.84	30.75	587	50.84	889	72.4	0.709165	0.000004
	T2	Lagoon water	2	37.5	12.02	62.77	619	11.16	30.75	585	50.03	887	73.5	0.709171	0.000004
Global ocean#	-	Oligotrophic Oceanic water (OOW)	-	34.9	10.28	54.10	468	10.21	28.30	545	24.6	873	88.0	0.709172	0.000023

Studied site	Shell sample	Corresponding water sample	Species	Ca (mg·g ⁻¹)	Mg (µg·g ⁻¹)	Sr (µg·g ⁻¹)	⁸⁷ Sr/ ⁸⁶ Sr	2 std dev
Oualidia lagoon, Morocco	O 1	OU 3	<i>Crassostrea gigas</i>	370	3888	1091	0.709174	0.000004
	O 2	OU 9	<i>Crassostrea gigas</i>	371	1361	575	0.709173	0.000004
	O 3	OU 12	<i>Venerupis decussata</i>	334	118	1689	0.709179	0.000004
	O 4	OU 9	<i>Solen marginatus</i>	361	268	2797	0.708939	0.000004
Salses-Leucate lagoon, France	S1	LEU 9	<i>Mytilus galloprovincialis</i>	354	483	746	0.709143	0.000004
	S2	LEU 2	<i>Glycymeris glycymeris</i>	na	na	na	0.709125	0.000004
	S3	LEU 3	<i>Crassostrea sp.</i>	na	na	na	0.709136	0.000004
	S4	LEU 4	<i>Mytilus galloprovincialis</i>	364	945	969	0.709174	0.000004
	S5	LEU 5	<i>Mytilus galloprovincialis</i>	378	1291	932	0.709158	0.000004
	S6	LEU 6	<i>Mytilus galloprovincialis</i>	349	135	1511	0.709177	0.000004
Banyuls-sur-Mer, France	Bs 1	B 1	Bivalve fragments	360	792	1513	0.709174	0.000004
	Bs 2	B 3	<i>Turritella communis</i>	362	413	1645	0.709173	0.000004
	Bs 3	B 5	Bivalve fragments	na	na	na	0.709165	0.000004
	Bs 4	B 7	Bivalve fragments	381	11554	2214	0.709171	0.000004
Tatakoto lagoon, French Polynesia	Ts 1	T1	<i>Tridacna maxima</i>	na	na	na	0.709178	0.000004
	Ts 2	T1	<i>Tridacna maxima</i>	365	810	1587	0.709164	0.000004
	Ts 3	T2	<i>Tridacna maxima</i>	na	na	na	0.709182	0.000004
	Ts 4	T2	<i>Tridacna maxima</i>	357	205	1644	0.709172	0.000004

Studied sites	End-members	Water type	Average Sr ($\mu\text{mol}\cdot\text{l}^{-1}$)	Minimal Sr ($\mu\text{mol}\cdot\text{l}^{-1}$)	Maximal Sr ($\mu\text{mol}\cdot\text{l}^{-1}$)	Average $^{87}\text{Sr}/^{86}\text{Sr}$	Minimal $^{87}\text{Sr}/^{86}\text{Sr}$	Maximal $^{87}\text{Sr}/^{86}\text{Sr}$	References
Morocco	Plio-Quaternary aquifer	GW	39.4	1.4	148.0	0.70874	0.70835	0.70962	Fakir et al., 2002; Kaid Rassou et al., 2005; Vinson et al., 2005
	Cretaceous aquifer	GW	29.8	3.0	116.4	0.70760	0.70760	0.70866	Fakir et al., 2002; Fadili et al., 2015; Bouchaou et al., 2017
	Jurassic aquifer	GW	82.7	36.3	249.9	0.70807	0.70774	0.70833	Kaid Rassou et al., 2005; Bouchaou et al., 2017
	Triassic aquifer	GW	164.9	164.4	165.5	0.70819	0.70805	0.70832	Bouchaou et al., 2017
	Atlantic water at Oualidia	SW	73.0	-	-	0.70916	-	-	<i>this study</i>
Corbières region, France	Quaternary aquifer	GW	17.1	1.8	85.5	0.70916	0.70814	0.71174	Petelet et al., 1998; Petelet-Giraud et al., 2016
	Pliocene aquifer	GW	7.8	4.7	15.2	0.71055	0.70987	0.71155	Petelet-Giraud et al., 2016
	Cretaceous aquifer	GW	4.2	1.2	9.0	0.70847	0.70753	0.70908	Khaska et al., 2013
	Jurassic aquifer	GW	9.6	0.3	53.0	0.70860	0.70788	0.70978	Aquilina et al., 2003; Petelet-Giraud et al., 2016
	Mediterranean water at Salses-Leucate	SW	69.7	-	-	0.70917	-	-	<i>this study</i>

