- 1 Constraints of applying strontium isotope stratigraphy in coastal and shallow marine
- 2 carbonates: insights from Lower Cretaceous carbonates deposited in an active tectonic
- 3 setting (N Iberian Basin, Spain).
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13 Abstract

14 The Lower Cretaceous Leza Fm is an essentially carbonate unit deposited at the 15 northernmost active margin of the Cameros Basin (N Spain) under an extensional tectonic 16 regime. This unit is composed of freshwater, marine-influenced, marginal-marine and 17 hypersaline marine carbonate facies, interbedded with variable amounts of alluvial deposits, mainly derived from the erosion of the Jurassic substrate. ⁸⁷Sr/⁸⁶Sr, δ^{18} O and δ^{13} C analyses 18 19 were obtained from carbonate facies of the Eastern and Western sectors of the basin. $\delta^{18}O$ 20 values follow the expected trend in both sectors: they are more negative (down to -7.9‰) in 21 freshwater carbonates and more positive (up to +2.8%) in marginal-marine to hypersaline 22 facies. However, independently of the seawater or freshwater influence, in the Western Sector 23 the ⁸⁷Sr/⁸⁶Sr values (0.707373-0.707801) are significantly lower and closer to the published Lower Cretaceous seawater ⁸⁷Sr/⁸⁶Sr ratios, than those of the Eastern Sector (0.707988-24 25 0.709033), where the overall marine influence was relatively high and the alluvial input low. 26 These data strongly suggest that ⁸⁷Sr/⁸⁶Sr ratios were mainly controlled by those of the riverine 27 freshwater arriving to the coastal and marine areas after the weathering and erosion of the 28 Jurassic carbonates or siliciclastic rocks, in the Western and Eastern sectors, respectively. 29 Thus, data indicate that, in coastal and shallow marine carbonates the influence of the riverine 30 water on the ⁸⁷Sr/⁸⁶Sr ratios should be systematically evaluated. This is particularly necessary in 31 active tectonic settings, where the uplifted areas are significantly prone to weathering and 32 erosion and where alluvial fan systems commonly developed, eventually discharging into 33 coastal and shallow marine areas.

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Keywords: Cameros Basin, coastal wetlands, carbon and oxygen isotopes, water-rock
 interaction

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38 Resumen

39 La Formación Leza es una unidad esencialmente carbonática del Cretácico Inferior 40 depositada en el borde norte de la cuenca de Cameros (N de España) en un contexto tectónico 41 extensional. Está formada por facies carbonáticas de agua dulce, con influencia marina, 42 marinas marginales e hipersalinas, intercaladas con cantidades variables de depósitos 43 aluviales, procedentes de la erosión del sustrato Jurásico de la cuenca. Se han obtenido datos 44 de ⁸⁷Sr/⁸⁶Sr, δ^{18} O y δ^{13} C de las facies carbonáticas en las zonas Oriental y Occidental de la 45 cuenca. Los valores de δ^{18} O siguen la tendencia esperable en ambas zonas: son más 46 negativos (hasta-7.9‰) en los carbonatos de agua dulce y más positivos (hasta +2.8‰) en las 47 facies marinas marginales e hipersalinas. Sin embargo, independientemente de la influencia marina o de agua dulce, los valores de ⁸⁷Sr/86Sr de la zona Occidental (0.707373-0.707801) 48 49 son significativamente inferiores y más próximos a los valores publicados para los carbonatos 50 marinos del Cretácico Inferior, que los de la zona Oriental (0.707988-0.709033), donde la 51 influencia marina fue, en general, relativamente mayor y el aporte aluvial menor. Estos 52 resultados indican que las relaciones de ⁸⁷Sr/86Sr estuvieron controladas principalmente por las 53 del agua dulce fluvial que llegaba a la zonas costeras y marinas tras la meteorización y erosión 54 del sustrato Jurásico de la cuenca, carbonático en el Sector Occidental y siliciclástico en el Oriental, y sugieren que, para la interpretación de las relaciones de ⁸⁷Sr/⁸⁶Sr en carbonatos 55 56 costeros y marinos someros, sobre todo de aquéllos depositados en contextos tectónicamente 57 activos, se debería evaluar sistemáticamente la influencia del agua dulce.

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59 Palabras clave: Cuenca de Cameros, humedales costeros, isótopos de carbono y oxígeno,
60 interacción agua-roca

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62 1. Introduction

63 Strontium isotopes are commonly employed for dating and correlating marine carbonates and fossil shells based on the assumptions that the seawater ⁸⁷Sr/⁸⁶Sr ratios of the 64 65 world's oceans have changed through time, but they are, and have been, uniform for a given 66 time (because its residence time is much greater than the time required for marine currents to 67 mix waters) and that, in contrast to trace elements and C and O isotopes, there is no 68 measurable fractionation of the Sr isotopes during precipitation of carbonates and fossil shells 69 (Veizer and Compston 1974; Brass 1976; Burke et al. 1982; Banner 1995; Steuber and Veizer 70 2002; McArthur et al. 2012). Accordingly, many curves reflecting the ⁸⁷Sr/⁸⁶Sr variation of 71 seawater through time have been published since the 70's, and they have been progressively 72 more detailed and based on a larger number of samples for a given period of time (e.g. Veizer 73 and Compston, 1974; Brass 1976; Burke et al. 1982; DePaolo and Ingram 1985; Koepnick et al. 74 1988; Jones et al. 1994; McArthur 1994; McArthur et al. 1994; 2001; 2007; 2012; Veizer et al. 75 1997; 1999; Jenkyns et al. 2002; Prokoph et al. 2008; Roveri et al. 2014; Korte and Ullmann 76 2016; Reghizzi et al. 2017; Wierzbowski et al. 2017). Based on those curves, the strontium 77 isotope stratigraphy (SIS) has become a powerful tool, used for dating and correlating marine 78 carbonate sequences (including deep to shallow and normal to restricted environments), as well 79 as for estimating the duration of stratigraphic gaps, biozones or Stages, for constraining the age 80 of condensed levels and as climatic indicator (e.g. DePaolo and Ingram 1985; Miller et al. 1988; 81 Hess et al. 1989; McArthur et al. 1992; 1993, 2000; Brasier et al. 1996; Bralower et al. 1997; 82 Barbieri et al. 1998; Weedon and Jenkyns 1999; Azmy et al. 1999; Scasso et al. 2001; Ebneth 83 et al. 2001; Steuber 2001; Price and Grocke 2002; Nieto et al. 2008; Frijia and Parente 2008; 84 Bodin et al. 2009; Lugli et al. 2010; Boix et al. 2011; Wehmiller et al. 2012; Williamson et al. 85 2012, Steuber and Schlüter 2012; Bonilla-Rodríguez et al. 2014, Frijia et al. 2015; Bover-Arnal 86 et al. 2016; Caus et al. 2016; Zuo et al. 2018; Frau et al. 2018; Fan et al. 2020, among many others). 87

However, ⁸⁷Sr/⁸⁶Sr values of the riverine and lacustrine freshwater may vary significantly
because they depend on the Sr isotopic values of the rocks that are being weathered and
eroded and on their ⁸⁷Rb content, commonly high in silicate rocks, which, after its decay to ⁸⁷Sr,
will significantly increase the ⁸⁷Sr/⁸⁶Sr ratio (Faure 1977; Stueber et al. 1984; Banner 1995).

92 Because of that, some authors have attempted to distinguish between marine and non-marine 93 environments based on the Sr isotopes (Schmitz et al. 1991; Hofer et al. 2013; Gierlowski-94 Kordesch and Cassle, 2015), and many other authors have studied and analysed the 95 importance of fresh- and seawater mixing and its relevance on the ⁸⁷Sr/⁸⁶Sr values in coastal to 96 shallow marine areas, close to the mainland (Müller et al. 1990; Müller and Mueller 1991; 97 Ingram and Sloan 1992; Andersson et al. 1992; Bryant et al. 1995; Banner 1995; Flecker et al. 98 2002; Flecker and Ellam 2006; Lugli et al. 2010; Topper et al. 2011; Sessa et al. 2012; Topper 99 and Meijer 2013; Roveri et al. 2014; Quijada et al. 2016a; Manzi et al. 2018; Meknassi et al., 100 2018; Roveri et al. 2019; Madhavaraju et al. 2020). In this sense, it has been argued by some 101 authors that the seawater ⁸⁷Sr/⁸⁶Sr ratio is rarely altered by freshwater inputs if salinities are 102 maintained above 10 to 20% (Ingram and Sloan 1992; Bryant et al. 1995; McArthur et al. 103 2012), although it has also been highlighted, even by the same authors, that small inputs of 104 riverine waters may have significant effects on the ⁸⁷Sr/⁸⁶Sr values of carbonate rocks and 105 shells, and that it would be wise to evaluate whether the local riverine inputs have altered the 106 seawater signals (Bryant et al. 1995; Banner 1995; McArthur et al. 2012; Meknassi et al., 2018) 107 when analysing Sr isotopic compositions in coastal and shallow marine carbonates and shells.

108 In this study, we have attempted to evaluate this issue by analysing carbonates of the 109 Leza Fm (Late Barremian-Early Aptian, Cameros Basin, N Spain), a coastal and essentially 110 carbonate unit, deposited in the context of the Mesozoic Iberian Extensional System (Fig. 1). 111 This unit is composed of freshwater, marine-influenced and marginal-marine carbonates, 112 restricted marine carbonates and evaporites, and alluvial detrital deposits that mainly derive 113 from the erosion of the uplifted Upper Jurassic substrate, which is essentially carbonate or 114 essentially siliciclastic, depending of the sector of the basin (Durántez et al. 1982; Alonso and 115 Mas 1990; Hernández-Samaniego et al. 1990; Ramirez Merino et al. 1990; Suarez-Gonzalez et 116 al. 2013; 2015). Thus, the main aim of this work is to evaluate the influence of the riverine 117 waters on the ⁸⁷Sr/⁸⁶Sr ratios of carbonates deposited in the different depositional environments 118 (from freshwater to marine and hypersaline) of the Leza Fm, and to evaluate the significance of 119 the ⁸⁷Sr/⁸⁶Sr ratios of the Jurassic substrate, weathered and eroded during deposition, in affecting the seawater ⁸⁷Sr/⁸⁶Sr ratios. 120

121 **2. Geological setting and stratigraphic framework**

122 The Cameros Basin is the northernmost basin of the Mesozoic Iberian Extensional 123 System (Fig. 1A-B). It was developed from the Tithonian to the Early Albian and records up to 124 6,500 m of vertical thickness of sediments (Mas et al. 2011; Omodeo-Salé et al. 2014; 2015; 125 Mas et al. in Martin-Chivelet et al. 2019). During the Eocene to Early Miocene, the Cameros 126 Basin was tectonically inverted, leading to the development of the thrusts that now limit the 127 Cameros structural unit to the north and south (Fig. 1B; Platt 1990; Casas-Sainz and Simón-128 Gómez 1992; Salas and Casas, 1993; Mas et al. 1993; 2011; Guimerà et al. 1995; Salas et a., 129 2011; Mas et al. in Martin-Chivelet et al., 2019).

130 The sedimentary record of the Cameros Basin comprises eight depositional sequences 131 (DS) (Mas et al. 2002; 2011; Arribas et al. 2003; Fig. 1C), and is characterised by the overall 132 and progressive migration of the depocenters of the successive DS to the N, which results in an 133 onlapping geometry observed between the successive units and the underlying Jurassic and, 134 locally Upper Triassic, substrate (Fig. 2A; Guiraud 1983; Mas et al. 1993; Suarez-Gonzalez et 135 al. 2013; 2016a; Omodeo-Salé et al. 2014; Mas et al. in Martin-Chivelet et al. 2019). In general, 136 the sedimentary record of the Cameros Basin is gently folded, and it is apparently unaffected by 137 relevant internal deformation (Guiraud 1983; Guiraud and Seguret 1985; Mas et al. 1993; 138 Omodeo-Salé et al. 2015); however, to the N and NE margins of the basin, where this study has 139 been carried out, it has been interpreted that syn-sedimentary normal faults controlled the 140 generation of the accommodation space and the thickness of some DS (Guiraud, 1983; Gómez-141 Fernández 1992; Quijada et al. 2013a; Suarez-Gonzalez et al. 2013; 2015; 2016a; Omodeo-142 Salé et al. 2014). Specifically, in the studied N and NE Cameros Basin, Suarez-Gonzalez et al. 143 (2016a) have interpreted that small fault-limited tectonic depressions were formed during the 144 Late Barremian-Early Aptian due to the interaction of the general extensional direction (mainly 145 N-S) with the late-Variscan structure of the basement. Additionally, the irregular distribution of 146 the Triassic evaporites (Keuper Facies), and its association with the onlap geometry of DS 1-7 147 onto the Jurassic substrate and even onto the Keuper Facies (Fig. 2A), led Suárez-González et 148 al. (2016a) to interpret a probable role of salt-tectonics on the sedimentation at the northern part 149 of the basin.

The sedimentary infill of the basin includes continental and coastal siliciclastic,
carbonate and evaporite deposits (Fig. 1C). In general, facies distribution along the basin shows

a proximal-distal trend: proximal continental and detrital facies are located to the W and SW; the
distal facies, comprising coastal carbonates, mixed carbonate-siliciclastic or mixed carbonateevaporite deposits, are located to the E and NE of the basin (Mas et al. 1993; 2002; 2011; 2019;
Arribas et al. 2003; Quijada et al. 2013a, 2013b; 2014; 2016b; 2020; Suarez-Gonzalez et al.
2013; 2014; 2015; Sacristán-Horcajada et al. 2015; 2016).

157 The syn-extensional record in the studied area, located at the northern and north-158 eastern margin of the Cameros Basin (Figs. 1B; 2A), comprises deposits of the Urbión and 159 Enciso Groups (DS7; Late Barremian-Early Aptian in age) and of the Oliván Group (DS8; Late 160 Aptian-Early Albian in age) (Figs. 1B-C; 2A; Mas et al. 1993; 2002; 2011;; Suarez-Gonzalez et 161 al 2013; 2015; 2016b; Mas et al., in Martin-Chivelet et al. 2019). The Urbión Group is 162 represented in the studied area by deposits of the Jubera Fm (Figs. 1C; 2A-B), which is 163 composed of reddish conglomerate, sandstone and siliciclastic mudstone (sensu Friedman et 164 al., 1992) and is interpreted as deposited in alluvial fan systems (Alonso and Mas 1993; Mas et 165 al. 2002; 2011; Ochoa 2006; Suárez-González et al. 2013). The Enciso Group overlies and 166 passes laterally to the Urbión Group (Fig. 1C). In the studied area, the lower part of the Enciso 167 Group is represented by the Leza Fm, which overlies and passes laterally to the Jubera Fm, 168 and is the focus of this study (Figs. 1C; 2A-B). The Leza Fm is essentially composed of 169 carbonates, with variable marine influence and content of detrital deposits, and is interpreted as 170 deposited in a coastal-wetland system (Suarez-Gonzalez et al. 2013; 2015). The rest of the Enciso Group, which overlie and pass laterally to deposits of the Leza Fm (Figs. 1C; 2A-B), is 171 172 composed of mixed carbonate-siliciclastic deposits interpreted as deposited in siliciclastic-173 influenced lacustrine and palustrine systems (Mas et al. 1993; 2002; 2011). The Oliván Group is 174 composed of reddish to greenish sandstone and siliciclastic mudstone deposited in fluvial and 175 coastal systems (Mas et al. 2011; Mas et al., in Martin-Chivelet et al. 2019).

A relevant feature of the Jubera and Leza Fms is that their deposits show significant variations in their thickness, as they were deposited in the above-mentioned small tectonic depressions formed during the Late Barremian – Early Aptian by faulting of the basin substrate (Fig. 2A-B; Alonso and Mas 1993; Suarez-Gonzalez et al. 2013; 2015; 2016a). The substrate of the Cameros Basin is mainly composed of Jurassic, and locally Upper Triassic, rocks (Fig. 2A, see below for details), which underlie syn-extensional deposits through an important

182 unconformity (Alonso and Mas 1993; Mas et al. 1993; 2002; 2011; Benito et al., 2001; 2005; 183 Benito and Mas, 2002; 2006). Erosion of the faulted substrate led to deposition of alluvial fan 184 sediments throughout the Jubera Fm; in the Leza Fm, alluvial fan deposits were restricted to the 185 margins of the depressions, and they changed laterally to carbonate coastal wetland sediments 186 (Fig. 3; Suarez-Gonzalez et al. 2013; 2015). This coastal wetland system was composed of a 187 mosaic of diverse and interrelated environments with influence of both fresh- and sea-water, the 188 latter coming both from the North, from the Boreal Realm, and the South, from the Tethys Sea 189 (Fig. 3; Suarez-Gonzalez et al., 2013).

Two main sectors are distinguished in the studied area, the Western and the Eastern
(Figs. 2-3), which have important differences in the sedimentary features of both the Jurassic
substrate and the syn-extensional Jubera and Leza Fms.

193 In the Western Sector, the substrate of the Jubera-Leza Fms comprises Upper Triassic 194 to Upper Jurassic rocks (Fig. 2A-B; Suarez-Gonzalez et al. 2013; 2015; 2016a). The Triassic is 195 composed of the Keuper Facies, which includes evaporites (gypsum), reddish siliciclastic 196 mudstone and minor dolostone (Fig. 2A; Suarez-Gonzalez 2015; Suárez-González et al., 197 2016a). The Keuper facies are largely deformed, and their thickness varies significantly along 198 the northern Cameros thrust (Fig. 2A). The Lower Jurassic is composed of approximately 150-199 200 m-thick marine limestone and/or dolostone, which were deposited in shallow to very 200 shallow carbonate platforms (Mensink 1966; Bulard 1972; Ramírez-Merino et al. 1990; 201 Hernández-Samaniego et al. 1990). The Middle Jurassic comprises approximately 300 m-thick 202 marine limestone, including a thick Bathonian sequence (up to 150 m thick) mainly composed of 203 oolitic deposits with minor siliciclastics, which were deposited in a shallow carbonate platform 204 (Benke 1981; Benke et al. 1981; Wilde 1990; Ramírez-Merino et al. 1990; Hernández-205 Samaniego et al. 1990; García-Frank et al. 2008). The pre-extensional Upper Jurassic record 206 (Oxfordian and Kimmeridgian) is composed of relatively thin shallow marine units (down to less 207 than 50 m thick), which include abundant coral reef deposits of Kimmeridgian age and minor 208 siliciclastic marine deposits (Benke et al. 1981; Alonso and Mas 1988; 1990; Errenst 1990; 209 Benito and Mas 2006). The overlying syn-extensional sedimentary record of the Jubera and 210 Leza Fms in the Western Sector, although highly variable, reaches up to approximately 500 m 211 of thickness (Fig. 2B; Suarez-Gonzalez et al. 2013). In this sector, the detrital facies of both

units are mainly composed of carbonate clasts coming from the erosion of the different units of
the Jurassic, mainly from the Bathonian to the Kimmeridgian units (Fig. 4A-B; Alonso and Mas
1993; Ochoa 2006; Suarez-Gonzalez et al. 2013; 2015); moreover, in the Western Sector the
overall detrital influence in the Leza Fm is higher than in the Eastern Sector (Suarez-Gonzalez
et al., 2015; see bellow and details in Section 4).

217 In the Eastern Sector, the substrate of the Jubera-Leza Fms is composed of Upper 218 Jurassic rocks (Figs. 2A-B; Suarez-Gonzalez et al. 2013; 2015). The Upper Triassic Keuper 219 Facies (which include very scarce and local outcrops of volcanic to subvolcanic rocks, less than 220 0.01 Km²) and the Lower and Middle Jurassic carbonate rocks are also observed in this sector, 221 but they do not directly underlie the syn-extensional sedimentary record (Fig. 2A; Suarez-222 Gonzalez et al., 2016a) and, thus, they probably they did not crop out during sedimentation of 223 the Jubera and Leza Fms. In the Eastern Sector, deposits from the Upper Triassic to the 224 Bathonian are equivalent to those observed in the Western Sector (Mensink 1966; Bulard 1972; 225 Durántez et al. 1982; Hernández-Samaniego et al. 1990). However, and in contrast to what is 226 observed in the Western Sector, in the Eastern Sector the Callovian to Kimmeridgian 227 sedimentary record is mainly composed of marine sandy limestone, sandstone and quartzite 228 conglomerates, which reach up to around 150 m (Benke 1981; Benke et al. 1981; Durántez et 229 al. 1982; Alonso and Mas 1988; 1990; Wilde 1990; Hernández-Samaniego et al. 1990; García-230 Frank et al. 2008). The syn-extensional sedimentary record of the Jubera and Leza Fms in the 231 Eastern Sector, although also variable, is overall thinner (up to 300 m) than in the Western 232 Sector (Fig. 2B; Alonso and Mas 1993; Suarez-Gonzalez at al. 2013; 2015); moreover in the 233 Eastern Sector, detrital deposits of both the Jubera and Leza Fms came from the erosion of the 234 Callovian and the Upper Jurassic units, which are mainly composed of quartzite conglomerate 235 and sandstone, with minor detrital carbonate fraction (Fig. 4C-D; Suarez-Gonzalez et al. 2015). 236 Additionally, the overall detrital content of the Leza Fm deposits is lower in the Eastern Sector 237 than in the Western Sector (see details in Section 4).

238 3. Materials and methods

Two selected stratigraphic sections of the Leza Fm, one from the Eastern Sector of the northern Cameros Basin (Préjano section), and another one from the Western Sector (the Leza River section) have been used for this study (Figs. 2, 5). A total of 341 rock samples from the

242 Leza Fm (214 limestone or marly limestone, 49 dolostone, 57 sandstone or sandy limestone, 21 243 conglomerate) were collected for laboratory studies (see Suarez-Gonzalez, 2015 for details 244 regarding the facies and samples). For each sample, a polished and uncovered thin section was 245 prepared to 30 µm thickness for petrographic analysis utilizing standard petrographic 246 techniques and cathodoluminescence petrography. For 75 samples, a polished 150-200 µm 247 thick section, matching the 30 µm thin section, was prepared for isotopic analyses. 248 Cathodoluminescence (CL) examination was carried out using a Technosyn® cold 249 cathodoluminescence unit operating at 20-25 kV with 300-400 µA beam current. Following 250 examination with CL, thin sections were stained with Alizarin Red S and potassium ferricyanide 251 (Dickson, 1966) for identification of carbonate minerals.

After petrographic examination and staining, well-preserved micritic or dolomicritic samples (samples displaying mudstone to wackestone texture *sensu* Dunham, 1962), lacking detrital input and petrographic evidence of diagenetic alteration, were selected for isotope analyses and microsampled directly from thick sections using a microscope-mounted drilling system, using 0.3 mm in diameter dental burs. When possible, approximately 20 mg of powdered samples were obtained, from which 100-150 µg were used for C and O isotopic analyses, and the rest for ⁸⁷Sr/⁸⁶Sr determinations.

259 62 analyses for δ^{13} C and δ^{18} O determinations were performed in the Stable Isotope 260 Laboratory at the University of Michigan. Sample powders reacted at 73 °C during 6 and 12 261 minutes (for calcite and dolomite, respectively) in an automated carbonate reaction system 262 (CarboKiel-IV) coupled directly to the inlet of a Thermo MAT 253 gas ratio mass spectrometer. 263 Isotopic ratios were corrected for ¹⁷O contribution and are reported in per mil notation relative to 264 the VPDB standard. Values were calibrated utilizing NBS 19 as the primary standard, and 265 analytical precision was monitored by daily analysis of NBS powdered carbonate standards. 266 Precision was better than 0.1 ‰ for both δ^{13} C and δ^{18} O measurements.

267 58 ⁸⁷Sr/⁸⁶Sr ratios were determined on an automated multicollector TIMS-Phoenix®
268 mass spectrometer at the Geochronology and Isotope Geochemistry Centre of the
269 Complutense University of Madrid. Carbonate powder was dissolved in 5 ml of 0.5M acetic acid.
270 Once dried, 1 ml of 3M HNO₃ (Merk-Suparapur[™]) was added to the sample and dried again.

271 3ml of 3M HNO₃ (Merk-Suparapur[™]) were then added to the samples, which were 272 subsequently centrifuged at 4000 r.p.m. during 10 minutes, in order to eliminate the solid 273 residue (clay minerals, quartz, etc). For the Sr chromatographic separation, an extraction resin 274 SrResinTM (Trisken International) was employed. The Sr was recovered with 0.05M HNO3 as 275 eluent. The fraction in which Sr was concentrated was recovered and dried for analysing in the 276 mass spectrometer. Sr analyses have been corrected for possible interference of ⁸⁷Rb and they 277 have been normalized to the value ⁸⁸Sr/⁸⁶Sr=0.1194. Analytical precision was monitored by 278 analysis of the NBS 987 standard (the mean value obtained for 7 samples were 0.710247; 2σ = 279 0.000008). Analytical uncertainties (referred to 2σ) were 0.01% for ⁸⁷Sr/⁸⁶Sr ratios. Blanks of Sr 280 preparations were lower than 0.05 ng/ml. The standard error for each sample was equal to or 281 lower than 3.

282 4. Results: The sedimentary record and isotopes of the Leza Fm.

283 The Leza Fm is a predominantly carbonate coastal unit, which has variable content of 284 detrital sediment and also shows a general trend of upwards increase in marine influence in 285 both the Eastern and Western sectors (Fig. 5; see detailed facies descriptions and 286 interpretations in Suarez-Gonzalez et al. 2013; 2014; 2015; 2016b; 2019). According to these 287 authors, tectonic activity caused significant variations in subsidence and thickness the unit 288 (which ranges from 20 m to 280 m) and in the lateral distribution of its facies, which include a 289 wide variety of interrelated deposits (Fig. 3), grouped into five facies associations (FA, Figs. 5-290 6):

291 The alluvial fan FA is characterised by conglomerates and sandstones. As pointed out 292 above (Section 2), in both the Eastern and the Western sectors, the main components of these 293 facies are lithoclasts of the Middle and Upper Jurassic (Bathonian to Kimmeridgian) rocks of the 294 basin substrate, which are mainly marine sandstone and quartzite conglomerates with minor 295 limestone in the Eastern Sector, and marine limestone in the Western Sector (Fig. 4). Deposits 296 of the alluvial fan FA are interbedded with freshwater to shallow marine carbonate deposits and, 297 in the Eastern Sector, also with hypersaline marine dolostone, which are described below (Figs. 298 5-6).

299 The freshwater and the marine-influenced limestone FAs are both characterised by 300 black and fetid limestones and, less commonly, by marls, which are widespread in both sectors 301 (Figs. 5; 6A-D). Abundant biota is observed in both FAs, including charophytes (in the 302 freshwater FA; Fig. 6A-B), dasycladales (in the marine-influenced FA; Fig. 6C-D), ostracods, 303 gastropods and vertebrate remains. Microbialites (oncoids, skeletal stromatolites and 304 thrombolites) are also observed (Suarez-Gonzalez et al., 2019). Both freshwater and marine-305 influenced limestone FAs are generally arranged in thickening-upwards sequences up to 4 m 306 thick, with abundant desiccation and edaphic features at their top (Figs. 5, 6A, C), and are 307 interpreted as deposited in shallow water-bodies with diverse salinities, from fresh to near-308 marine, which underwent periods of desiccation and inundation, and were surrounded by 309 vegetated areas (Suarez-Gonzalez et al. 2015). Thus, both FAs are sedimentologically 310 equivalent, differing only in the influence of seawater and their palaeontological content. 311 Lithoclasts of Jurassic rocks may occur within these limestone beds, indicating lateral 312 association with the alluvial fan palaeoenvironments and suggesting that the source of 313 freshwater to the water bodies was related to the weathering and erosion of the Jurassic 314 substrate of the basin (Suarez-Gonzalez et al. 2013; 2015).

Regarding isotopic compositions, <u>freshwater limestone</u> yields δ^{18} O values of -7.9 to -5.8‰ in the Eastern Sector and of -6.5 to -4.3 ‰ in the Western Sector; δ^{13} C values range between -5.2 and -0.4‰ in the Eastern Sector and between -8.0 and -2.6‰ in the Western Sector; ⁸⁷Sr/⁸⁶Sr ratios ranges between 0.708785 and 0.707988 in the Eastern Sector and between 0.707373 and 0.707719 in the Western Sector (Figs. 5, 7; Table 1 of Supplementary material).

321 <u>Marine-influenced limestone</u> yields δ^{18} O values of 5.4 to -5.6‰ in the Eastern Sector 322 and of -2.1 to -4.8‰ in the Western Sector; δ^{13} C values range between -1.7 and -5.9‰ in the 323 Eastern Sector and between -3.5 and -4.9‰ in the Western Sector; ⁸⁷Sr/⁸⁶Sr ratios in the 324 Western Sector range between 0.707597 and 0.707801 (Figs. 5, 7; Table 1 of Supplementary 325 material). It was not possible to obtain enough micritic carbonate powder for performing Sr 326 isotopes in thick sections of this FA in the Eastern Sector, because of the abundance of calcite 327 cement filling intra- and interparticle primary porosity, which could have altered the Sr isotopic 328 values of depositional (micritic) carbonate.

329 The marginal-marine carbonate FA (Fig. 6E-F) is characterised by grey limestone or 330 dolostone, which are slightly different in the Eastern and in the Western sectors. In the Eastern 331 Sector, these deposits are characterised by alternation of oolitic and micritic dolostone 332 containing abundant ostracods, miliolid foraminifers and agglutinated stromatolites (Suarez-333 Gonzalez et al., 2014; 2016b; 2019), which displays flaser, wavy and lenticular bedding and 334 subaerial exposure features. In the Western Sector, this FA consists of grey limestone, either 335 bioclastic (containing ostracods, miliolid foraminifers and gastropods) or micritic, displaying 336 commonly fenestral porosity and desiccation features. Facies in both sectors are interpreted as 337 deposited in shallow marginal-marine areas affected by tides. This FA was dominated by 338 seawater, with salinities mostly higher than in the marine-influenced FA, but still probably 339 variable, temporarily decreasing due to freshwater influence (mainly in the Western Sector) or 340 increasing, due to evaporation (in the Eastern Sector, see next FA), which in turn, favored early 341 dolomite precipitation (Suarez-Gonzalez et al. 2014; 2015; 2016b; 2019), as observed in recent 342 coastal carbonate settings where syn-depositional dolomite commonly precipitates (e.g. Tucker 343 and Wright, 1990; Warren, 2016 and references therein).

In the Eastern Sector, marginal-marine dolostone of this FA yields δ^{18} O values of -1.3 to +0.6‰, δ^{13} C values of -6.7 to -1.8‰ and 87 Sr/ 86 Sr ratios of 0.708302 to 0.708978. In the Western Sector, marginal-marine limestone of this FA yields δ^{18} O values of -4.0 to -1.1‰, δ^{13} C values of -4.8 to -1.7‰ and 87 Sr/ 86 Sr ratios of 0.707679 to 0.707761 (Figs. 5, 7; Table 1 of Supplementary material).

<u>The hypersaline dolomite-evaporite FA</u> (Fig. 6 G-H) is only observed in the Eastern Sector, where it is laterally associated with deposits of the marginal-marine FA (Figs. 3; 5). This FA is characterised by laminated dolomicritic deposits, containing rare ostracods and foraminifers, abundant pseudomorphs after gypsum and anhydrite, desiccation features and, locally, micritic-evaporitic stromatolites. The dolomite-evaporite FA is interpreted as deposited in very shallow and restricted hypersaline marine areas, which reached high salinity through evaporation (Suarez-Gonzalez et al. 2013; 2015; 2019).

356 Dolostone of this FA yields δ^{18} O values of -1.0 to +2.8‰, δ^{13} C values of -8.2 to -1.3‰ 357 and ⁸⁷Sr/⁸⁶Sr ratios of 0.708304 to 0.709033 (Figs. 5, 7; Table 1 of Supplementary material).

358 **5. Discussion: Constraints of applying strontium isotope stratigraphy in coastal and**

359 shallow marine carbonates

Isotopic data obtained from carbonates of the different FA of the Leza Fm in both studied sectors are apparently contradictory. As discussed below, the δ^{18} O values reflect the sedimentary palaeoenvironment and the nature of the waters from which they precipitated, however, the ⁸⁷Sr/⁸⁶Sr values reflect the strong influence of the riverine waters, which may have had a wide range of Sr isotope compositions, depending on the nature of rocks that were being weathered and eroded during deposition.

366 δ^{18} O values obtained from carbonates of the different FA, in both the Eastern and the 367 Western sectors, are consistent with carbonate precipitation from fresh- to marine and even 368 hypersaline waters. Freshwater limestone in both sectors shows the lowest δ^{18} O values. δ^{18} O 369 values progressively increase in marine-influenced limestone, in marginal-marine limestone (in 370 the Western Sector) and dolostone (in the Eastern Sector) and in hypersaline dolostone (Figs. 371 5, 7A-B). According to Prokoph et al. (2008), the oxygen isotopic composition of the Upper 372 Barremian-Lower Aptian marine limestone and calcitic fossils deposited in low latitude (as that 373 of the Iberian Plate for that time, e.g. Masse et al. 2000) ranged approximately between --2.8 374 and -0.4‰ (Figs. 5, 7A-B). In the case of the Leza Fm, δ^{18} O values of marginal-marine 375 limestone of the Western Sector range between -4.0 and -1.1‰, which are close to or in the 376 range of the published marine values by Prokoph et al. (2008) (Figs. 5, 7A-B). In the Eastern 377 Sector, δ^{18} O values of marginal-marine dolostone range between -1.3 to +0.6‰, being heavier 378 than those of the marginal-marine limestone of the Western Sector, but also mostly in the range 379 of the published marine values (Fig. 7A-B). The heavier δ^{18} O values in dolostone (the mean δ^{18} O value of marginal marine dolostone is -0.31‰ whereas the mean δ^{18} O value of marginal 380 381 marine limestone is -2.95‰; Table 1 of supplementary material) are consistent with the 382 difference between the water-calcite and the water-dolomite fractionation factors for a given 383 temperature, which results in dolomite δ^{18} O values of ~ 3‰ heavier than those of the calcite precipitating from the same water and at the same temperature (Fritz and Smith 1970; 384 385 Friedmann and O'Neil 1977; McKenzie 1981; Tucker and Wright 1990; Arenas et al., 1997). 386 Freshwater limestone, as expected, shows more negative δ^{18} O values, which are in the range of

387 carbonates precipitated from meteoric freshwater at low latitudes (e.g. Hudson 1977; Allan and 388 Matthews 1982; Lohmann 1987; James and Choquette 1990; Tucker and Wright 1990). Marine-389 influenced limestone has δ^{18} O values ranging between the marine and freshwater limestones, 390 which would be derived from the mixing of sea- and freshwater at different proportions (e.g. Frank and Lohmann 1995). Dolostone of hypersaline facies has the heaviest δ^{18} O values, 391 392 ranging from -1.0 to +2.8‰. These values are in the range of the δ^{18} O values obtained from 393 dolomites precipitated in Cenozoic and recent coastal sabkhas undergoing intense evaporation 394 (e.g. Tucker and Wright 1990; Arenas et al., 1997; Warren 2016, and references therein).

395 Regarding the C isotopic compositions, Prokoph et al. (2008) estimated δ^{13} C values 396 ranging approximately between +0.8 and +3.4‰ for marine limestone and calcitic fossils 397 deposited in low latitude during the Late Barremian-Early Aptian. δ^{13} C values of all analysed 398 carbonates of the Leza Fm, however, are largely variable and more negative than those 399 published for marine carbonates (Fig. 7A). This shift to lighter values would have been caused 400 by the incorporation of ¹²C into calcite and dolomite, which could have derived from the 401 oxidation of organic matter, commonly soil-derived (Allan and Matthews 1977; 1982; Lohmann 402 1987; Tucker and Wright 1990; Leng and Marshall, 2004), and/or from bacterial sulphate 403 reduction, as observed in some modern coastal sabkhas (Tucker and Wright 1990; Warren 404 2016 and references therein). In fact, the combination of relatively invariant δ^{18} O values with 405 highly variable and negative δ^{13} C values is a common trend in meteoric or in transitional marine-406 to-meteoric systems (Lohmann, 1987; James and Choquette, 1990).

407 ⁸⁷Sr/⁸⁶Sr ratios, as pointed out in the introduction, are commonly employed for dating 408 and correlating marine carbonates (strontium isotope stratigraphy, SIS; McArthur, 1994; Veizer 409 et al. 1997, 1999; Prokoph et al., 2008; McArthur et al., 2012, among many others) and, 410 according to published data (Jones et al., 1994; McArthur et al., 2001; 2012; Prokoph et al., 411 2008), the ⁸⁷Sr/⁸⁶Sr values of Upper Barremian-Lower Aptian marine carbonates approximately 412 range between 0.7073 and 0.7075 (Figs. 5, 7C). However, and in contrast to what occurs with 413 the O isotopic compositions, ⁸⁷Sr/⁸⁶Sr data of the Leza Fm carbonates do not show the 414 expected trend according to their different depositional palaeoenvironments in none of the studied sectors; moreover, the ⁸⁷Sr/⁸⁶Sr ratios obtained in the Eastern and Western sectors are 415

significantly different (Figs. 5, 7C). In the Eastern Sector, the ⁸⁷Sr/⁸⁶Sr ratios are higher (more 416 417 radiogenic) than those expected for Upper Barremian-Lower Aptian marine carbonates. 418 Additionally, it is also significative that all the analysed carbonates, regardless of their fresh-, 419 marine- or hypersaline-water origin, show similar range of ⁸⁷Sr/⁸⁶Sr values in each sector (more 420 radiogenic in the Eastern Sector and less radiogenic in the Western Sector), despite δ^{18} O 421 values being different according to their depositional palaeoenvironments (Fig. 7B-C). It could 422 be argued that the Sr isotopic compositions, which are similar in all the carbonates of each 423 sector, but different in the Eastern and in the Western sectors (Figs. 5, 7C), could be derived 424 from diagenetic alteration. However, powder samples were carefully obtained from the areas of 425 thick sections of samples with no evidences of diagenetic alteration; furthermore, it is important 426 to note that, if the Sr isotopic ratios would have been altered by diagenetic fluids, changes in the 427 oxygen isotopic compositions would be also expected, because the $\delta^{18}O$ of carbonates depends 428 on the δ^{18} O of the waters from which they precipitate (a diagenetic fluid, if alteration had 429 occurred) and on temperature (e.g. Fritz and Smith 1970; Friedmann and O'Neil 1977; Banner 430 1995). However, the δ^{18} O values are in accordance with the different palaeoenvironmental 431 conditions where carbonates of the different FAs were formed, thus indicating the absence of 432 significant diagenetic modification of the original isotopic signature.

433 Therefore, it is interpreted that the Sr isotopic values of carbonates of the Leza Fm are 434 largely derived from ⁸⁷Sr/⁸⁶Sr compositions of the riverine freshwaters that discharged into the 435 coastal and shallow marine areas after weathering and eroding the faulted and exposed marine 436 Jurassic rocks of the substrate. In the Eastern Sector, the eroded Middle to Upper Jurassic 437 substrate, is mainly siliciclastic (e.g. Durántez et al. 1982; Hernández-Samaniego et al., 1990; 438 see Section 4 for detail). In this regard, riverine waters may have a wide range of ⁸⁷Sr/⁸⁶Sr 439 values, but if they have weathered and eroded Rb-rich silicate rocks (such as granite, gneiss and siliciclastic sedimentary rocks), their 87Sr/86Sr values are commonly higher than those 440 441 estimated for the Phanerozoic seawater (Faure 1977; Stueber et al. 1984; 1987; Banner 1995). 442 In that case, even small inputs of riverine waters into the sea may have significant effects on the ⁸⁷Sr/⁸⁶Sr values of the resulting mixed water, which may increase significantly (Banner 1995, 443 444 Bryant et al. 1995; McArthur et al., 2012; Meknassi et al., 2018). Thus, in the Eastern Sector, 445 where essentially siliciclastic Jurassic rocks were being weathered and eroded, the more

radiogenic ⁸⁷Sr/⁸⁶Sr values of the riverine freshwater would have significantly contributed to
increase the marine ⁸⁷Sr/⁸⁶Sr values of carbonates of the Leza Fm, while the oxygen isotopic
compositions reflect their formation in the different freshwater to marine and hypersaline
depositional palaeoenvironments (Fig. 7B-C).

450 In the Western Sector, however, the ⁸⁷Sr/⁸⁶Sr values of freshwater to marginal-marine 451 limestone are lower and closer to the published data of the Upper Barremian-Lower Aptian 452 marine carbonates than those of the Eastern Sector, even though the overall marine influence 453 in the Western Sector was lower than in the Eastern Sector (Figs. 5, 7C). In fact, in the Western Sector, the freshwater limestone shows ⁸⁷Sr/⁸⁶Sr values closer to the Upper Barremian-Lower 454 455 Aptian marine Sr isotopic values than those of the marginal-marine limestone, although the $\delta^{18}O$ 456 values record the expected trend, consistent with their depositional environment (more negative 457 in freshwater than in marginal marine carbonates) (Figs. 5, 7B-C, see above). These 458 contradictory data may be explained if the following facts are considered: 1) in the Western Sector, the substrate that was being eroded was mainly composed of Lower to Upper Jurassic 459 460 carbonates (e.g. Ramírez-Merino et al. 1990; Hernández-Samaniego et al. 1990; see section 4); 2) the ⁸⁷Sr/⁸⁶Sr ratios of the Jurassic marine carbonates range between ~ 0.7068 and ~0.7077 461 462 (Jones et al. 1994; MacArthur 2001; Prokoph et al. 2008; MacArthur et al. 2012); 3) the range of the Jurassic Sr isotopic values, in turn, includes the range of ⁸⁷Sr/⁸⁶Sr values from the Late 463 464 Barremian to Early Aptian (Figs. 5, 7C). In this sense, and similarly to what occurs when 465 weathering Rb-rich silicate rocks, previous authors have interpreted that meteoric waters and 466 brines that interact with marine carbonates may inherit Sr isotope compositions that are 467 diagnostic of the age of the carbonates (e.g. Müller et al. 1990; Müller and Mueller 1991; Banner 468 et al. 1994; Banner 1995). Thus, in the Western Sector, the riverine freshwater would have had 469 less radiogenic ⁸⁷Sr/⁸⁶Sr values compared to those in the Eastern Sector, because Jurassic 470 carbonates were being weathered and eroded, in contrast to the Eastern Sector, where the 471 Jurassic substrate was mainly siliciclastic. Thus, in the Western Sector, when riverine and 472 seawater mixed, both with similar Sr isotopic compositions, the resulting water would have 473 similar Sr isotope ratio than that of both end members, leading to freshwater and marine 474 carbonates having similar Sr isotopic ratios, but different δ^{18} O values, depending on their

475 depositional environment, and even leading to freshwater carbonates showing ⁸⁷Sr/⁸⁶Sr values
476 within the range of the Late Barremian-Early Aptian values (Figs. 5, 7B-C).

477 In this regard, Bryant et al. (1995) made a two-component mixing model to calculate the 478 ⁸⁷Sr/⁸⁶Sr of waters of different salinities and tested it with analyses performed in molluscs from 479 estuaries of the Mississippi Sound and coastal Florida. On the one hand, these authors found 480 out that both, their model results and analyses, suggested that even in the most marginal-481 marine systems the freshwater flux did not have a measurable influence until the salinity was 482 very low (10 ppt or less), and these findings have been used for interpreting that "when rivers locally lower the salinity of seawater, marine ⁸⁷Sr/⁸⁶Sr is rarely altered at salinities above 20 psu" 483 (McArthur et al. 2012; note that psu is equivalent to g/kg or 0/00). Nevertheless, Bryant et al. 484 485 (1995) also pointed out that carbonates precipitating in estuarine settings not always record the global marine ⁸⁷Sr/⁸⁶Sr value because, once the marine signature is affected by freshwater 486 487 input, 87 Sr/86 Sr seawater values change rapidly, leading McArthur et al. (2012) to highlight that 488 "when dating coastal and shallow water faunas, it is wise to establish that the local riverine 489 inputs did not alter the ⁸⁷Sr/⁸⁶Sr of seawater in the depositional environment.". More recently, Meknassi et al. (2018) have analysed ⁸⁷Sr/⁸⁶Sr values from modern marine carbonate skeletons 490 491 (bivalves, gastropods, cephalopods, chitons, and calcifying algae) collected in coastal settings 492 worldwide and have found out that epibenthic and eurytopic organisms, such as bivalves and 493 gastropods, from coastal domains with water mass restriction, low salinity or strong continental supplies, may display slight to considerable offsets compared to the ⁸⁷Sr/⁸⁶Sr values of modern 494 495 seawater. Based on their data, these authors calculated that "only 10%, 33%, and 52% of the 496 published Phanerozoic ⁸⁷Sr/⁸⁶Sr curve can provide time calibration with respective accuracies of 497 ± 1 , ± 2 and ± 3 m.y., hence obscuring most dating at the scale of the shorter Phanerozoic

498 stages".

In the Leza Fm, our data strongly support that the ⁸⁷Sr/⁸⁶Sr ratios of all carbonates (freshwater, marine-influenced, marginal-marine, and hypersaline) were, not only affected, but mainly controlled by the Sr ratios of the riverine freshwater, which discharged into the coastal and marine areas after weathering and eroding the carbonate or siliciclastic Jurassic substrate of the basin. Interestingly, this strong influence of freshwater Sr isotopic values is observed even without petrographic evidence of detrital input in the analysed carbonate samples, and

505 even though the δ^{18} O values of carbonates are coherent with their formation from fresh- to 506 marine and hypersaline waters. Nevertheless, it is important to highlight that it has been 507 possible to make this interpretation because the Jurassic substrate of both studied sectors is 508 different; if the Jurassic substrate were carbonate and similar in both sectors, as occurs in many 509 areas of the Iberian Basin (Gómez, et al. 2004; 2019), it would not have been possible to accurately evaluate the influence of the riverine water in the ⁸⁷Sr/⁸⁶Sr of carbonates, considering 510 511 that the ⁸⁷Sr/⁸⁶Sr of Barremian-Aptian marine carbonates, as well as most of those of the 512 Cretaceous, are in the range of the Jurassic ⁸⁷Sr/⁸⁶Sr marine values, being even identical for 513 some periods of time (Jones et al. 1994; McArthur 2001; Prokoph et al. 2008; MacArthur et al. 514 2012). Thus, our data indicate that caution should be taken when interpreting Sr isotopic data 515 for performing SIS studies obtained from fossil shells and/or carbonates (even those well-516 preserved and/or without direct evidence of detrital input) deposited in coastal and shallow 517 marine sedimentary environments. This is particularly important for coastal and shallow-marine 518 carbonates deposited in active tectonic settings. In these settings, the uplifted areas are 519 significantly prone to weathering and erosion, leading to the common development of alluvial 520 fan systems that eventually discharge into coastal and shallow marine settings, as occurred in 521 the Iberian Basin during the Late Jurassic-Early Cretaceous extensional phase.

522 6. Conclusions

523 The Lower Cretaceous Leza Fm (Cameros Basin, N Spain) is an essentially carbonate 524 unit, which was deposited in the context of the Mesozoic Iberian Extensional System, and 525 whose deposition and thickness were strongly controlled by faulting of the basin substrate 526 (mainly composed of Jurassic rocks). A coastal wetland system that included carbonates 527 deposited in freshwater, marine-influenced, marginal-marine and hypersaline water bodies, 528 formed at that time in the studied area. This system was laterally related with alluvial fans, 529 whose deposits (conglomerate, sandstone, sandy limestone and marl), sourced in the faulted 530 and exposed basin substrate, are interbedded with carbonates.

Two sectors have been differentiated, Eastern and Western, based on the sedimentary features of the Leza Fm deposits and those of the Jurassic substrate, which are slightly different. In the Eastern Sector, the marine influence is higher with the detrital inputs being lesser than in the Western Sector; additionally, the Middle and Upper Jurassic rocks of the

substrate are essentially siliciclastic in the Eastern Sector but essentially carbonate in theWestern Sector.

537 The δ^{18} O values in both sectors follow the expected trend, in accordance with the 538 different depositional palaeoenvironments: more negative values (down to -7.8‰) are recorded 539 in freshwater carbonates and more positive values (up to +2.8%) in marine to hypersaline marine facies. However, and independently of the marine or freshwater influence, the ⁸⁷Sr/⁸⁶Sr 540 541 ratios of the carbonates in the Western Sector (0.707373-0.707801) are significantly lower and 542 closer to the published Lower Cretaceous marine ⁸⁷Sr/⁸⁶Sr ratios, than those obtained in the 543 Eastern Sector (0.707988-0.709033) although, in this sector, the overall marine influence was 544 higher and the detrital alluvial input lower.

These data strongly support that the ⁸⁷Sr/⁸⁶Sr ratios of all carbonates studied herein (freshwater, marine-influenced, marginal-marine, and hypersaline) were strongly controlled by the Sr ratios of the riverine freshwater, which arrived to the coastal and marine areas after weathering and eroding the carbonate or siliciclastic rocks of the Jurassic substrate of the basin that was faulted and exposed, even if there is no petrographic evidence of detrital influence, and even if the δ^{18} O values of carbonates are coherent with their formation in fresh- to marine and hypersaline waters.

Thus, our data indicate that caution should be taken when interpreting Sr isotopic data for performing SIS studies obtained from fossil shells and/or carbonates (even those wellpreserved) deposited in coastal and shallow marine sedimentary environments, particularly in active tectonic settings, such as those occurring in the Iberian Basin during the Late Jurassic and Early Cretaceous extensional system.

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1008 Figure captions

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Fig. 1. Geological setting of the studied area. A. Map of the Iberian Peninsula, showing the preMesozoic Variscan Massif and the Mesozoic Iberian Extensional System (MIES). Red square
marks the location of the Cameros Basin. B. Geological map of the Cameros Basin showing the
location of the studied area. Red square marks the location of Figs. 2A and 3. C. Chrono- and
lithostratigraphic chart of the eight depositional sequences (DS) of the Cameros Basin
sedimentary infill, modified after Mas et al. (2011). The Leza Fm (highlighted in red) is part of
DS7, Late Barremian – Early Aptian in age.

1016 Fig. 2. A. Detailed geological map of the northernmost margin of the Cameros Basin (modified 1017 from Suarez-Gonzalez et al. 2015). Note the tectonic control and individualization of outcrops of 1018 the Jubera and Leza Fms, deposited mostly on top of the Bathonian to Kimmeridgian marine 1019 Jurassic substrate of the basin and, locally, over the pre-Bathonian marine Jurassic or over the 1020 Triassic Keuper facies. Note that the Jubera plus Leza Fms lithotopes are limited by faults, and 1021 that the colour code of this map is equivalent to that of the stratigraphic chart of Fig. 1C. B. 1022 Correlation panel of the stratigraphic sections showing the thickness of the Jubera and Leza 1023 Fms (the stratigraphic sections analysed in this study, the Leza River section (LZ), in the 1024 Western Sector, and the Préjano section (PR), in the Eastern Sector, are highlighted with red 1025 asterisks). On top, to the right, there is a simplified map showing the location of each section 1026 and the correlation line (in blue). The top of the Leza Fm has been used as datum for 1027 correlation. The thickness of the Jubera Fm has been obtained from Ochoa (2006), Hernández-1028 Samaniego et al. (1990) and our own measurements. Note the tectonic control of the Jubera 1029 and Leza Fms and the lateral facies change between the Jubera and Leza Fms. Modified from 1030 Suarez-Gonzalez et al. 2013.

Fig. 3. A. Palaeogeographic map of NE Iberian Peninsula during deposition of the Leza Fm.
(modified from Suarez-Gonzalez et al. 2013) B. Detailed palaeogeographical scheme showing a
general interpretation of the spatial distribution of depositional environments in NE Cameros
Basin during sedimentation of the Leza Fm coastal wetlands and the complex array of
sedimentary environments that characterizes these multifaceted depositional systems (see
location in A; modified from Suarez-Gonzalez et al., 2015). The map covers approximately the

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same area as the map in Fig. 2A (see names of the main towns in both maps). White areas

represent zones with no outcrops of Lower Cretaceous rocks. These areas may have been
those where the Jurassic substrate of the Cameros Basin cropped out during sedimentation,
being the source of the Leza Fm alluvial sediments.

1041 Fig. 4. A. Field photograph showing conglomerates of the Leza Fm at the Western Sector, 1042 where they are mainly composed of fragments of Jurassic carbonates. **B**. Transmitted light 1043 photomicrograph showing in detail the detrital facies of the Leza Fm at the Western Sector. 1044 Note that many clasts are composed of oolitic limestone (JL) and reworked single ooids (JO) 1045 and echinoderms fragments (blue arrow), which are identical to those of the Upper Jurassic 1046 limestone deposited in the studied area, and other carbonate extraclasts (red arrows) of 1047 unknown origin. Quartz grains (Q), are also present in variable amounts. C-D. Field 1048 photographs of the detrital alluvial sediments of the Leza Fm at the Eastern Sector. C. Very 1049 poorly sorted conglomerate made up of large marine sandstone clasts (yellow arrows), quartzite 1050 pebbles (white), and sandy matrix. Coin is 2.3 cm in diameter. **D**. Poorly-sorted conglomerate 1051 made up mainly of quartzite pebbles (white) in a carbonate matrix.

Fig. 5. Detailed stratigraphic sections used for isotopic analyses: the Leza River section, in the Western Sector, and the Préjano section, in the Eastern Sector (see Fig. 2 for location). In each section the δ^{18} O (pink) and the ⁸⁷/Sr/⁸⁶Sr (black) are shown. The range of published δ^{18} O and ⁸⁷Sr/⁸⁶Sr values for the Barremian-Aptian marine carbonates and, in the case of the Sr isotopes, for the Jurassic are also shown. Blue circles represent the depositional environment where the different carbonate facies were deposited.

1058 Fig. 6. Photographs showing the different studied carbonate facies, which were deposited in

1059 different sedimentary palaeoenvironments. **A-B**. Freshwater facies association. **A**. Field

1060 photograph showing tabular black limestone interbedded with thin beds of greenish marl. Note

1061 root traces at the top of the beds. Coin is 2.3 cm in diameter. **B**. Transmitted light

1062 photomicrograph showing a wackestone of charophyte thalli forming the nucleus of oncoids. C-

- 1063 D. Marine-influenced facies association. C. Field photograph showing tabular marine-influenced
- 1064 black limestone. Note the root traces at the top of the bed. Tip of hammer at the top-right corner
- 1065 for scale. **D**. Transmitted light photomicrograph showing a wackestone of dasycladales green

1066 algae. E-F. Marginal-marine facies association. E. Field photograph of dolostone displaying

1067 wavy bedding with alternation of grainy facies showing small-scale cross-bedding (red arrows)

1068 and muddy facies (blue arrows). Coin is 2.4 cm in diameter. F. Transmitted light

1069 photomicrograph showing a wackestone-packstone of benthic miliolid foraminifera and

1070 ostracods. G-H. Hypersaline facies association. G. Field photograph of hypersaline dolostone

1071 displaying abundant calcite pseudomorphs after gypsum (yellow arrows). H. Transmitted light

1072 photomicrograph showing a dolomudstone displaying calcite pseudomorphs after lenticular

1073 gypsum.

1074 Fig. 7. A. Carbon and oxygen isotope compositions of carbonates of the different facies 1075 associations in the Leza River section (Western Sector) and in the Préjano section (Eastern 1076 Sector). In both sectors, carbonates of each facies association have negative and variable δ^{13} C values and relatively invariant δ^{18} O values (more negative in freshwater limestone and 1077 1078 progressively more positive in marginal-marine and hypersaline carbonates). **B-C**. δ^{18} O and 1079 ⁸⁷Sr/⁸⁶Sr values, respectively, *versus* the depositional environments where carbonate 1080 precipitated in both sectors. Note that in both sectors and, as expected, δ^{18} O values (B) are 1081 more negative in freshwater limestone, becoming progressively heavier in marine-influenced, 1082 marginal-marine and hypersaline carbonates. Note that most of the marine carbonates are in 1083 the range of or close to the published δ^{18} O values of the Barremian-Aptian marine carbonates (see text for explanation). However, 87 Sr/ 86 Sr ratios in each sector are very different, but δ^{18} O 1084 1085 values of carbonates precipitated in the different palaeoenvironments in each sector are similar. 1086 Also note that ⁸⁷Sr/⁸⁶Sr ratios in the Western Sector are similar or in the range of the published 1087 Barremian-Aptian marine carbonates, even those obtained from freshwater carbonates. 1088

1089 Table 1 (Supplementary material). Isotopic data of the analysed samples

1090

Fig. 1













Fig. 4







Fig. 7

