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A 700-year multiproxy reconstruction on the Argentinian Pampas inferred from the sediments of Laguna Blanca Grande

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Manuscript: A 700-year multiproxy reconstruction on the Argentinian Pampas inferred from the sediments of Laguna Blanca Grande

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1 **A 700-year multiproxy reconstruction on the Argentinian Pampas inferred from**  
2 **the sediments of Laguna Blanca Grande**

3  
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28

29 **ABSTRACT**

30

31 The Pampean region is a crucial area to obtain sensitive paleoclimatic lacustrine  
32 archives due to the presence of shallow environments in a territory non impacted by  
33 humans until the last centuries. In this study, we provide a paleoecological  
34 reconstruction for the last ca. 700 years based on a multiproxy lacustrine record from  
35 Laguna Blanca Grande, in Olavarría (Buenos Aires, Argentina). Our inferences, which  
36 were based on sedimentary properties, diatom, cladoceran and ostracod assemblages,  
37 offered interesting information about hydroclimatic variability and nutrient increase.  
38 Changes in relative abundances on diatoms, specifically on *Aulacoseira granulata* and  
39 *Aulacoseira granulata* var. *angustissima* and fragilarioids, were used to infer shifts in  
40 nutrient conditions. The remainder proxies together indicated small lake level changes.  
41 Reconstructed hydroclimatic conditions in Laguna Blanca Grande are consistent with  
42 previous paleoecological inferences indicating a humid phase around ca. AD 1450 and  
43 progressive drier conditions ca. AD 1530-1900. A flood gate construction and an  
44 increase of nutrients in the lake revealed a higher human pressure due to population  
45 increase and land-use changes during the last century. Further studies on taxonomy and  
46 autecology of microcrustaceans are needed to effectively unlock the information  
47 contained in biological proxies from Sudamerican records.

48

49 **Keywords:** nutrient enrichment, hydroclimatic reconstruction, lake level changes,  
50 diatoms, cladocerans, ostracods.

51

52 **1. Introduction**

53 The Pampean plains are fertile and vast lowlands that cover more than 750,000  
54 km<sup>2</sup>, which include some regions in Argentina, Uruguay, and the southernmost states of  
55 Brazil (Politis, 2008; Viglizzo et al., 2001; Zarate, 2003). The interannual rainfall  
56 variability in this region is related to the sea surface temperature over the western South  
57 Atlantic, the intensity and position of the South Atlantic Convergence Zone (SACZ) and  
58 the South American summer monsoon (SASM) (Barros et al., 2000; Garreaud et al.,  
59 2009), which explains the migration of moisture and precipitation patterns in

60 subtropical plains producing rainy conditions during the austral summer. In addition to  
61 the intrannual variability, the region is strongly influenced by interannual phenomena  
62 such as the El Niño Southern Oscillation (ENSO). As a result of these different modes  
63 of climatic variability, the Pampean lakes have shown important lakes level changes  
64 during the Holocene, as demonstrated in Laguna Mar Chiquita (Cuña-Rodríguez, 2018;  
65 Coianiz et al., 2014; Piovano et al., 2014; 2009; 2004; 2002), Lagunas Encadenadas del  
66 Oeste de Buenos Aires (Córdoba, 2012; 2014), Lake Nahuel Rucá (Stutz et al., 2010),  
67 Lake Lonkoy (Stutz et al., 2012), Lake La Barrancosa (Plastani et al., 2019), Lake  
68 Adela (Dangavs and Mormeneo, 2012), Laguna del Monte (Dangavs and Pierrard,  
69 2013), Lake La Brava (Irurzun et al., 2014; Laprida et al., 2014) and Lake Melincué  
70 (Guerra et al., 2015; 2017) (Fig. 1). Despite the existence of some paleohydrological  
71 reconstructions, multiproxy inferences in the southern Pampas (classification according  
72 to Iriondo, 1994) are still needed to provide additional information on the timing and  
73 relationship of local hydrovariability with large-scale climatic events (Guerra et al.,  
74 2017; Lüning et al., 2019). Moreover, understanding the interaction of past climatic  
75 changes with limnological features is also crucial to face future challenges such as the  
76 synergic effect of climate warming and eutrophication on local/endemic biodiversity  
77 (Kopprio et al., 2010). This is particularly meaningful when taking into account the  
78 socioeconomical implications and ecosystem services of these lakes, considering that  
79 they play an important role as reservoirs of endemic biodiversity, flood control,  
80 recreation, tourism and climate change mitigation (Iwan et al., 2017).

81 In this study we used a lacustrine record based on diatoms, cladocerans and  
82 ostracods from Laguna Blanca Grande (Buenos Aires, Argentina) to provide new  
83 paleoecological information from southern Pampas (Fig. 1) and also to help to  
84 understand the temporal-spatial features of hydrological changes and the role of human  
85 activities in the last centuries. We hypothesize that this sequence may contain a  
86 relatively good signal of past natural conditions and climate variability since this region  
87 was not highly impacted by humans for a longtime. Documentary and historical sources  
88 (*e.g.*, Djenderedjian, 2012; Mayo, 2000; Pedrotta et al., 2012) indicated that only sparse  
89 native populations inhabit La Pampa until the end of the 19th century. Migration and  
90 farming conversion did not take place until the beginning of the 20th century.  
91 Hydroclimatic variability as well as the recent human impact might have been recorded  
92 in our sediment record.

93

94 **2. Site Description**

95

96 The Argentinian Pampean region comprises an extensive plain (~500,000 km<sup>2</sup>) covering  
97 the center-east of the country. The geological setting is characterized by loess-like  
98 deposits covered by Quaternary aeolian sediments, composed of massive to poorly  
99 stratified sandy silt, partially reworked by fluvial action and with common carbonate  
100 accumulations of unknown origin ( Muhs, 2013; Rubio et al., 2019; Zarate, 2003; Zarate  
101 and Tripaldi, 2012). The climate of southern Pampas is subtropical semiarid,  
102 characterized by mean annual temperature of 14.4°C, with mean values of 2°C in the  
103 coldest months (June-July) and 29°C in the warmest (August). Mean annual rainfall is  
104 about 901 mm (mean values 1988-2010, National Meteorological Service from the  
105 Olavarría meteorological station). The geomorphology of this region -low slope and  
106 absence of geographical features- allows the development of wetlands and shallow lakes  
107 in the most depressed areas of the plain. Most of the lakes are small (< 500 ha), shallow  
108 (~ 4 m), eutrophic and polymictic (i.e., without thermal stratification) (Geraldini et al.,  
109 2011; Laprida, 2008; Quirós and Drago, 1999).

110 Laguna Blanca Grande (Fig. 1; 36°29'12, 99"S, 60°53'45,91"W) is located  
111 between Olavarría and San Carlos de Bolívar, Buenos Aires Province, Argentina. It is  
112 an almost round lake with a 450 ha basin, an average depth of 80 cm and a maximum of  
113 180 cm. It receives water from the Brandsen Creek, in a permanent regime, and from  
114 three temporary watercourses. Water flows from the lake into the Arroyo Las Flores,  
115 where there is a sluiceway to regulate the level. This gate was probably built around the  
116 1950s in a context of water management actions around Lagunas Encadenadas del Oeste  
117 to prevent the impacts of floods and droughts in the region (Monachesi and Albaladejo,  
118 1997).

119 Lake shows alkaline waters (pH= 8.7) and total nitrogen values ca. 81.3 mg/l  
120 (Hassan, 2011). Conductivity is ca. 0.6 mS/cm and hardness is ca. 227 mg/l (Hassan et  
121 al., 2011). The order of major ion concentration is HCO<sub>3</sub><sup>-</sup> >> SO<sub>4</sub><sup>2-</sup> > Cl<sup>-</sup> and Na<sup>+</sup> >>  
122 Mg<sup>++</sup> > K<sup>+</sup> > Ca<sup>++</sup> (Colautti and Remes-Lenicov, 2003). Modern diatom assemblage is  
123 composed of *Aulacoseira granulata*, *Staurosira longirostris*, *Cyclotella meneghiniana*,  
124 *Hippodonta hungarica*, *Pseudostaurosira brevistriata* (Hassan et al., 2011; Hassan and

125 De Francesco, 2018). Zooplankton is composed of the rotifers *Keratella tropica*,  
126 *Notholca* sp., *Asplanchna girodi* and *Polyarthra vulgaris*, the cladocerans *Bosmina* sp.  
127 and *Macrotix laticornis* and the copepod *Notodiaptomus incompositus*. This is the main  
128 source of food of the Argentinian silverside (*Odontesthes bonariensis*), which also  
129 inhabits this lake (Colautti and Remes-Lenicov, 2003). Today, recreational fishing and  
130 tourism are some of the main activities around the lake. However, at regional scale,  
131 Olavarría has a diversified economy with important contribution of agriculture,  
132 stockbreeding and industry related to mining activities (Olavarría, 2010).

133

### 134 **3. Materials and methods**

135

#### 136 **3.1. Coring, sampling and chronology**

137 Sediment cores were recovered with a Livingston piston corer from the deepest  
138 part of the lake (0.6 m) in December 2013. Two parallel and overlapping cores (LBG-A  
139 and LBG-B) were taken from the lake at a distance of 3 m from one another. Core LBG-  
140 A was 77 cm in length while core LBG-B was 102.5 cm in length. Core LBG-A  
141 extended from the mud-water interface to a depth of 77 cm. After the correlation of the  
142 cores, it was decided not to analyze the upper part of the core LBG-B, from the mud-  
143 water interface to 37.5cm, and perform the analysis only from 37.5 cm to the bottom of  
144 the core (65 cm length). This strategy ensured an overlap of ca. 40 cm between the two  
145 cores. Sediment cores were wrapped in plastic film, placed in PVC tubes and stored in a  
146 cool room at 4°C until further processing. Core description was carried out following  
147 the methodology described in Schnurrenberger et al. (2003) and the Munsell color chart  
148 (Munsell Colour Company, 1975). The stratigraphic column was build using the  
149 lithological patterns suggested by the US Geological Survey (2006). Color and main  
150 physical properties such as composition, structure, or degree of humification were the  
151 initial basis to establish a correlation between both cores. Then, the correlation was  
152 confirmed by analytical measurements and biological proxies. Based on this correlation,  
153 four bulk sediment samples along the whole length of the composite record were  
154 selected for AMS  $^{14}\text{C}$  analysis at the CHRONO Center laboratory at Queen's University  
155 Belfast, UK. Radiocarbon dates were calibrated using the ShCal13 database (Hogg et

156 al., 2013) and the 95.4% distribution (2s probability interval) was considered to build  
157 the age-depth model.

158 As one of the samples did not accomplish the principle of superposition, an age-  
159 depth Bayesian statistic model was built using only three samples. The age-depth model  
160 was built using the package Bacon (Blaauw and Christen, 2011) in R 3.4.2 software (R  
161 Development Core Team, 2019). The curve was adjusted with a Gaussian model and  
162 included the starting condition that the surface was -63 as a  $\pm 0$  cal. years BP. Using  
163 these settings, over 7,000 iterations were run using a Markov chain Monte Carlo  
164 (MCMC) method to estimate the unknown parameters in the age-depth model.  
165 Although carbonate accumulations of variable morphology and genesis are common in  
166 the loess sequences (Muhs, 2007), Fontana (2007) showed that reservoir effect was  
167 negligible in a Pampean shallow lake in the same geological setting. Therefore, no  
168 correction for reservoir effect was applied in the model.

169

### 170 **3.2. Analytical and biological methods**

171 Volumetric subsamples of  $1 \text{ cm}^3$  were taken at 5 cm intervals for organic matter,  
172 granulometry and biological analysis (diatoms, cladocerans and ostracods). For  
173 granulometric analysis, samples were dispersed in distilled water after organic matter  
174 and carbonates were dissolved with  $\text{H}_2\text{O}_2$  and HCl, respectively, and analyzed by laser  
175 diffraction (Mastersizer Malvern, 2000). The samples with granulometry  $> 250 \mu\text{m}$   
176 were sieved. Results were integrated into the GRADISTAT V 4.0 program (Blott and  
177 Pye, 2001). The granulometric analysis was performed at the Marine Geology  
178 Laboratory at the Instituto Argentino de Oceanografía, Argentina. Organic matter was  
179 determined by weight loss on ignition (LOI) at  $550 \text{ }^\circ\text{C}$  for four hours. Subsequently, the  
180  $\text{CO}_2$  mass evolved from carbonate was determined by LOI at  $950^\circ\text{C}$  for two hours, and  
181 the carbonate content was calculated by multiplying the weight loss by 1.36 (Heiri and  
182 Lotter, 2001). LOI analysis was carried out at the Centro de Investigaciones en Ciencias  
183 de la Tierra (CICTERRA - CONICET - UNC).

184 For diatom and chrysophyte cyst analysis, samples were treated with sodium  
185 pyrophosphate ( $\text{Na}_2\text{P}_2\text{O}_7$ ) to deflocculate the sediment and to remove the clays. Then,  
186 15 mL of HCl (35%) were added to wet samples to remove carbonates. Finally, samples  
187 were heated to a water bath for 2 hours with  $\text{H}_2\text{O}_2$  to remove organic matter (Metzeltin

188 and García-Rodríguez, 2003). They were washed with distilled water successively until  
189 reaching a neutral pH in between the three treatments mentioned above. Once cleaned,  
190 permanent slides were made using an Entellan® (refractive Index: 1.54) mounting  
191 medium. At least 400 diatoms valves per sample were identified in an optical  
192 microscope at 1250× magnifications with oil immersion. Relative abundances of taxa  
193 (included chrysophyte cysts) were calculated by dividing the number of valves and  
194 cysts from each species by the total count on each slide/sample. These analyses were  
195 carried out at the Geoscience laboratory from CURE-Rocha, Uruguay. Diatom species  
196 were identified using the appropriate keys (Frenguelli, 1941; Gómez and Bauer, 2000;  
197 Hasle and Syvertsen, 1997; Krammer and Lange-Bertalot, 1991a, 1991b, 1988, 1986;  
198 Metzeltin et al., 2005; Metzeltin and García-Rodríguez, 2003; Theriot et al., 1992).  
199 Ecological information on diatom taxa preferences was extracted from Denys (1991);  
200 Theriot et al. (1992), Van Dam et al. (1994), Gómez and Bauer (2000a), Kociolek and  
201 Spaulding (2003), Rühland et al. (2003). Hassan et al. (2009) and Solak et al. (2012).

202 For cladoceran analysis, 1 cm<sup>3</sup> of sample was heated and stirred in 10%  
203 potassium hydroxide (KOH) for 30 min. Then, the sediment samples were sieved  
204 through a 40 µm mesh, following the procedure described by Szeroczyńska and  
205 Sarmaja-Korjonen (2007). The sieve residue was carefully transferred to a beaker. One  
206 tablet of *Lycopodium* spores was placed on a watch glass and a few drops of 10% HCl  
207 were added to dissolve the tablet. Afterward, the solution was poured off into the beaker  
208 and mixed well. Then, the sample was transferred into a test tube and centrifuged for 10  
209 min at 3500 rpm. After centrifugation, the water was discarded, and a small quantity of  
210 ethanol was added and mixed. A slide was placed on a hot plate and liquefied glycerol  
211 jelly with some safranine drops were added. Then, a few drops of sample with ethanol  
212 was added and spread over the coverslip area. Finally, a coverslip was placed on the  
213 slide and pressed gently. Samples were identified under a LOMO/LUMAN fluorescence  
214 microscope at 20-100× magnification. Cladoceran identifications and ecological  
215 characteristics were obtained from the literature on south American cladocerans and  
216 from López-Blanco and Sinev, (2016), Paggi (1998a), (1995) and Smirnov (1971). This  
217 analysis was performed at the Escuela Politécnica Nacional, Ecuador.

218 Sample preparation for ostracod analysis was carried out using the methodology  
219 proposed by Holmes (2001) and Danielopol et al. (2002). Samples were washed and  
220 sieved through a 63 µm mesh. The freeze-cooling technique was used for sample



221 disintegration in fine-grained sediment, by adding sodium hexametaphosphate before  
222 freezing. Then, samples were dried at room temperature for 12 hours, and the valves  
223 were picked out with a fine brush on a Nikon stereoscope (SMZ645) at 50×  
224 magnification. Identifications were based on taxonomic keys and specialized diagnosis  
225 (D'Ambrosio et al., 2015; Laprida, 2006; Ramón-Mercau et al., 2014). Absolute  
226 abundances were calculated from the number of adult valves of each species in 10 mL  
227 of sample. Additionally, charophyte oogonia, fish remains, gastropods and testate  
228 amoebae were also identified. Ostracod analysis was carried out at the GIBPC  
229 laboratory from Universidad Pedagógica y Tecnológica de Colombia.

230 Stratigraphic diagrams showing the relative abundances of each proxy were  
231 performed using the *Psimpoll* 4.27 software (Bennett, 2009). The broken stick model  
232 was applied to determine the number of significant stratigraphic zones (Bennett, 1996).  
233 Major zones were identified using the optimal division information content for  
234 ostracods and the binary splitting by sums-of-squares for cladocerans and diatoms.  
235 Before performing the zonation analyses, ostracod data were transformed using square  
236 root to stabilize the variances and increase the importance of rare species. Diatom data  
237 were  $\log(x+1)$  transformed to give less weight to dominant and/or abundant taxa. Only  
238 the most abundant species were used for diatom analysis; those whose abundance was  
239 greater than 2% in at least 3 samples and considering that the percentage removed was  
240 less than 10% of the total abundance of each sample (Karst and Smol, 2000).

241 Detrended correspondence analysis (DCA) were applied to diatom data to  
242 estimate the degree of species turnover (Hill and Gauch, 1980). DCA has yielded good  
243 results for diatom analysis, allowing interpretation of records with different temporal  
244 scales and environmental gradients (e.g., Correa-Metrio et al., 2014; Hassan et al.,  
245 2012). The meaning of DCA axis 1 was inferred in terms of *a priori* knowledge of their  
246 distribution in modern environmental gradients (Bicudo et al., 2016; Hassan et al.,  
247 2009). Then, the axis scores DCA1 was plotted stratigraphically to provide a simplified  
248 picture of changes through time. DCA analysis was performed using “*vegan*” package  
249 (Oksanen et al., 2019) in software R 3.6.3 (R Development Core Team, 2020).

250

#### 251 **4. Results**

252



#### 253 4.1. Sedimentology and physical characteristics

254 Physical properties and sedimentology analysis in cores LBG-A and LBG-B  
255 resulted in an overlap of ca. 40 cm. 66.5 cm in core LBG-A overlapped with 27.5 cm in  
256 core LBG-B, producing a composite sequence of 104 cm. This composite record was  
257 composed of dark-brownish sandy silt sediments with an increasing proportion of sand  
258 from top to the bottom part of the core. Eleven facies and six sedimentary units were  
259 distinguished, their colors varied from brown (10 YR 4/3) to dark brown (10 YR 4/29)  
260 (Table 1; Fig. 2). The mean values of organic matter throughout the first 75 cm of the  
261 sediment core were ca. 7% with a decreasing trend toward the top. A sharp shifting was  
262 observed ca. 75 cm (contact zone of Facies 8 and 9), where the organic matter content  
263 decreased towards the bottom of the core. Carbonate content was low, with mean values  
264 around 0.05% and increasing values in Unit 1.

265

#### 266 4.2. Age-depth model

267 The chronological model based on three  $^{14}\text{C}$  radiocarbon dates (Fig. 3) yields an  
268 average sedimentation rate (SR) of 1.47 mm/yr for Laguna Blanca Grande sediment  
269 core. However, the model showed distinct sedimentation rates depending on the age  
270 (Fig. 3). From 99.5 to 40 cm, the SR was relatively higher but it slowed down from 40  
271 cm to the top of the sediment core.

272

#### 273 4.3. Diatoms

274 A total of 55 diatom taxa representing 34 genera were identified in the Laguna  
275 Blanca Grande sediment core. The most representative taxa belonged to seven genera  
276 (*Aulacoseira*, *Cyclotella*, *Thalassiosira*, *Nitzschia*, *Amphora*, *Staurosira*, and *Surirella*).  
277 *Aulacoseira granulata* (Ehrenberg) Simonsen and *Aulacoseira granulata* var.  
278 *angustissima* (O.Müller) Simonsen dominated the sediment record (Fig. 4). Other  
279 abundant species were also *Aulacoseira ambigua* (Grunow) Simonsen, *Aulacoseira*  
280 *muzzanensis* (Meister) Krammer and *Cyclotella meneghiniana* (Kützing). Zonation  
281 analysis indicated the presence of four zones (DT-1, DT-2, DT-3 and DT-4). DT-1 zone  
282 (99.5-72 cm) was characterized by higher relative abundances of *A. granulata* and *C.*  
283 *meneghiniana*. In DT-2 (72-25 cm), the percentages of *A. granulata* decreased although

284 they were still high. The two upper zones (DT-3 and DT-4; 24-10 cm and 9-0 cm,  
285 respectively) were associated with a relative increase of *A. granulata* var. *angustissima*  
286 in detriment of *A. granulata*. Besides, in the uppermost zone (DT-4), *Staurosira*  
287 *longirostris* (Frenguelli) Metzeltin, Lange-Bertalot & García-Rodríguez, *Staurosira*  
288 *construens* Ehrenberg and *Surirella* *rorata* Frenguelli increased their relative  
289 abundances.

290 DCA axes 1 and 2 of diatom samples were 1.52 and 1.16 standard deviations  
291 (SD) of species turnover in length. *A. granulata* var. *angustissima*, *S. longirostris* and *S.*  
292 *construens* had the highest scores on DCA Axis 1, while *A. granulata* and *A.*  
293 *muzzanensis* and chrysophytes, were associated with the lowest scores (Fig. 4).

294

#### 295 4.4. Cladocera

296 A total of six Cladocera taxa were identified in Laguna Blanca Grande (Fig. 5);  
297 *Bosmina* Baird, 1846, which usually prefers pelagic environments and five littoral taxa  
298 *Chydorus sphaericus*-group, *Leydigia* sp. Kurz, 1875, small *Alona* sp., *Alona* cf. *affinis*  
299 (Leydig, 1860) and *Pleuroxus* Baird, 1843 (Fig. 5). *Bosmina* was the dominant taxa  
300 throughout the sediment core with relative values ca. 70 - 90%. In the CL-1 zone (99.5 -  
301 75 cm), *Bosmina* showed its maximum relative abundances. CL-2 (75-10 cm) zone was  
302 marked by higher relative abundances of *Chydorus sphaericus*-group, which together  
303 with the benthic *Leydigia* had a more continuous presence throughout the zone.  
304 However, both groups of cladocerans showed several changes along the whole zone,  
305 especially around 32 cm, where *Chydorus* increased at the expense of *Bosmina*. CL-3  
306 (10 - 0 cm) was characterized by higher values of *Bosmina* and decreasing values of the  
307 benthic species.

#### 308 4.5. Ostracods

309 Four species of ostracods were recorded in the Laguna Blanca Grande sediment  
310 record (Fig. 6): *Limnocythere cusminskyae* Ramón-Mercau, 2014, *Cyprideis salebrosa*  
311 Bold, 1963, *Heterocypris incongruens* Ramdohr, 1808 and *Ilyocypris ramirezi*  
312 Cusminsky and Whatley, 1996. The number of individuals per sample never exceeds  
313 400 and the maximum abundance was 40 ind /mL (Fig. 6). *Limnocythere cusminskyae*  
314 and *H. incongruens* were the main taxa of the assemblage. *L. cusminskyae* dominated

315 over 90% of the assemblages and represented 85% of individuals throughout the core.  
316 In fact, six out of the 23 samples contained monospecific populations of *L.*  
317 *cusminskyae*, which were dominated by females. *Ilyocypris ramirezi* was only found in  
318 two samples, while only juvenile valves from *C. salebrosa* were found in five depths.  
319 The preservation of the valves throughout the composite sequence was heterogeneous,  
320 with both well preserved and broken specimens.

321 OST-1 (104 -75 cm) was characterized by the presence of gastropods, fish  
322 scales, charophytes, *L. cusminskyae* (< 25 ind/mL), *H. incongruens* and juveniles from  
323 *C. salebrosa* ). In OST-2 (75 - 63 cm) and OST-4 (51 - 39 cm), ostracods were absent.  
324 The top of OST-3 (63-51cm) was mainly composed of articulated shells of *L.*  
325 *cusminskyae* (< 40 ind/mL) with some valves of *H. incongruens*. In OST-5 (39 - 7 cm),  
326 gastropod remains, fish scales and testate amoebae were recorded, while ostracods were  
327 again abundant (< 30 ind/mL). *Limnocythere cusminskyae* was the dominant species,  
328 representing 85% of the assemblage, followed by *H. incongruens* (13%), while *I.*  
329 *ramirezi* and *C. salebrosa* represented the remaining 2%. In OST-6 (7 - 0 cm), ostracods  
330 were less abundant than in the previous zone. Only specimens of *L. cusminskyae* with  
331 ruptured and disarticulated valves together with testate amoebae and fish remains were  
332 recovered (Fig. 6).

333

## 334 5. Discussion

335 The biological assemblage in Laguna Blanca Grande is typical of shallow, eutrophic  
336 and alkaline freshwater systems from the Pampean region (Hassan, 2011; Laprida,  
337 2006; Paggi, 1998b; Plastani et al., 2019; Smol, 1985). Biological proxies coupled with  
338 sediment properties' changes suggested a shift in nutrients and hydroclimatic conditions  
339 during the last centuries. Diatom assemblage was marked by high relative abundances  
340 of *A. granulata*, which is replaced by the variety *angustissima* in the upper part of the  
341 record. Given the different ecological preferences of these varieties for nutrient  
342 conditions (Bicudo et al., 2016; Kilham and Kilham, 1975; Stoermer et al., 1985;  
343 Turkia, 1999), this is interpreted as changes in trophic state. Bicudo et al. (2016) and  
344 Turkia and Lepisto (1999) found that *A. granulata* has a lower weighted average  
345 optimum for nitrogen and phosphorous than the variety *angustissima*. Moreover, the  
346 increase of *A. granulata* var. *angustissima* in our sequence also occurs together with an

347 increase of *C. meneghiniana*, which is favored by high organic and turbid waters with  
348 limited light penetration (Hassan, 2013). These inferred changes in nutrient conditions  
349 are further evidenced by the DCA axis 1, which reflected the turnover of *A. granulata*  
350 by the variety *angustissima* in the upper part of the record and showed negative scores  
351 dominating before 1750 AD (Fig. 6). This inferred change in nutrient availability was  
352 not followed by an increase in organic matter, as shown in Figure 2. In many lakes, a  
353 large fraction of organic matter is decomposed under conditions high concentrations of  
354 oxygen and resuspension of sediments generated by wind (Meyers and Ishiwatari,  
355 1995). These conditions are more frequent in polymictic lakes, like Pampean lakes,  
356 where the entire water column is mixed over the year or even daily. Mixing would  
357 produce an oxic environment at the bottom, increasing the rates of decomposition of  
358 organic matter by microorganisms in the upper part of the record. Furthermore, the  
359 nutrient increase might have triggered the observed shift in species composition but not  
360 a biovolume increase, as also noted in other paleoecological studies (e.g. López-Blanco  
361 et al., 2011).

362         However, the remaining biological and sedimentological proxies did not register  
363 a distinct change in the nutrient state, but they may indicate small oscillations in lake  
364 levels. The interpretation of cladoceran assemblages has limitations derived from the  
365 restricted knowledge of taxonomical and autecological characteristics in this part of the  
366 world. The assemblages showed the dominance of *Bosmina* sp., which generally has  
367 pelagic preferences. Regarding nutrient availability, *Bosmina* has eurioic preferences  
368 since it has been recorded both in oligotrophic (López-Blanco et al., 2020, 2011) and  
369 eutrophic environments (George, 1974; Lotter et al., 1998; Solis et al., 2018). The main  
370 components of the cladoceran littoral-benthic community might have been favored  
371 either under lower lake levels and/or under an increase in the trophic conditions. *C.*  
372 *sphaericus* and *Alona* are ubiquitous taxa with a great capacity for colonization, they  
373 can benefit from both nutrient enrichment and shallower conditions (Alonso, 1996;  
374 Smirnov, 1971). However, when plotting planktonic/benthonic ratio from cladoceran  
375 assemblages and comparing with ostracods, sedimentological data and other regional  
376 reconstructions, all together were concordant with small changes in hydrovariability  
377 (Fig. 7). Ostracods show a response to conductivity changes and thus, to small lake  
378 level changes. Four zones (OST-1, 3, 5 and 6) were characterized by the dominance of  
379 *L. cusminskyae*, which is highly tolerant to alkaline conditions and oligohaline waters

380 (Laprida, 2006; Marquez et al., 2016) and suggest a higher solute concentration at  
381 shallower levels. These biozones were intercalated by OST-2 and OST-4, where the  
382 absence of ostracods may indicate unstable conditions for ostracod colonization.

383 At the bottom of the sediment core (99.5 - 72 cm; ca. AD 1335-1472) (DT-1;  
384 CL-1; OST-1), less eutrophic conditions are inferred by higher values of *A.granulata*.  
385 The sediment here was composed of dark brown sandy coarse silt (Unit 1), poor in  
386 organic matter. The highest proportion of sand fraction in this unit, as well as the  
387 highest sedimentation rate, suggest important fluvial input that might have increased the  
388 lake level. The dark brown color associated with low organic matter content indicates  
389 relatively strong reducing conditions and a deposition in a perennial lacustrine  
390 environment (Wu and Li, 2004). The highest sedimentation rate calculated from the  
391 age-depth model in this section (Fig. 3) and the higher values of *Bosmina* and *A.*  
392 *granulata*, which was also related to high river flow conditions (Hötzel and Croome,  
393 1996; Nogueira, 2000; Wang et al., 2009), are also compatible with considerable  
394 sediment input due to the high water inflow in Laguna Blanca Grande. Higher lake level  
395 reconstructed in the lower LIA (ca. AD 1270-1340) in Laguna Blanca Grande agrees  
396 well with the idea of a more humid phase inferred from Botuverá Cave (Bernal et al.,  
397 2016; Fig. 7) and with more humid conditions during the preceding period, the  
398 Medieval Climatic Anomaly (ca. AD 900-1300) (Cioccale, 1999; Iriondo and Kröhling,  
399 1995). At the regional scale, this inference is concordant with the high-level stands  
400 described in Lagunas Encadenadas del Oeste (Laprida et al., 2009), Laguna Mar  
401 Chiquita (Coianiz et al., 2014; Piovano et al., 2002) and Lake Melincué (Guerra et al.,  
402 2015) (Fig. 7).

403 From ca. 72 – 9 cm (ca. AD 1472-1930) (DT-2) decreasing proportions of *A.*  
404 *granulata* in favor of *A.granulata* var. *angustissima* indicated variable conditions in  
405 terms of nutrient enrichment. Documentary sources show that indigenous societies were  
406 already present in the Pampean region 1000 years ago (Mazzanti, 2003). However, most  
407 of the native inhabitants were small and nomadic groups of hunter-gatherers and the  
408 population situated southern to Rio Salado was very scarce until the eighteenth century  
409 (Aldazabal, 2002). In AD 1828, historical documents situated the military fort of Blanca  
410 Grande during the “Previous Seasons to the Desert Conquest” very close to the lake for  
411 a very short time (Crivelli, 2013). Before Laguna Blanca is already mentioned in some  
412 historical documents from AD 1770-1790 (Floury-Dagorn, 2013), but there is historical

413 and documentary evidence of the presence of sparse native population whose main  
414 activity was stockbreeding (Pedrotta et al., 2012). This period of variable nutrient  
415 enrichment was probably combined with small lake level oscillations with a tendency  
416 towards shallower conditions at the end of this period. Laminated dark brown silty fine  
417 sand (Unit 2) that changes to black sandy silt highly bioturbated with higher organic  
418 matter content (Unit 3) suggests a deposition by suspension in a low energy  
419 environment, which is concordant with the absence of ostracods (OST-4, OST-2) and  
420 with higher *Bosmina* abundances at relatively higher levels. The decrease in sand  
421 content indicates limited fluvial input in comparison with the previous period, which is  
422 supported by a lower SD in the chronological model (Fig. 3). The fine clastic lamination  
423 could respond to variations in the water and sediment inputs into the lake. In Unit 5, the  
424 sediment was composed of dark brown sandy coarse silt with iron mottles indicating  
425 intermittent oxidized conditions, which suggests that they were mainly deposited on a  
426 very shallow lake with high mixed conditions or temporary subaerial exposure.  
427 Ostracods also showed the highest total values and a more continuous presence,  
428 supporting the idea of decreasing lake levels. In particular, *L. cusminskyae* dominated  
429 the assemblage, which, together with *H. incrongruens*, might also indicate oscillations  
430 in lake levels and subsequent conductivity variations (Kihn et al., 2017; Laprida and  
431 Valero-Garcés, 2009; Marquez et al., 2016). Lake level fluctuations with a progressive  
432 reduction of level from 1340 AD to 1900 AD are ascribed to drier conditions and  
433 frequent extreme events registered in the Pampean region during the LIA (Córdoba,  
434 2012; Córdoba et al., 2014; Guerra et al., 2017; Laprida et al., 2009; Piovano et al.,  
435 2009).

436 Nutrient enrichment is inferred in the upper part of the sediment record (from 9  
437 cm; AD 1925) by higher relative abundances of the diatoms *A.granulata* var.  
438 *angustissima* and by the establishment of *S. longirostris* and *S. construens* (Dixit et al.,  
439 1992). Hassan et al. (2014) also interpreted a shift from *Aulacoseira* spp. by fragilarioid  
440 taxa as a nutrient increase caused by the development of intensive farming activities.  
441 The Pampean Region experienced large scale deforestation since the end of the 19th  
442 century due to agriculture and railroad construction, which favored soil degradation  
443 (Dussart et al., 2011; Melo, 2004). Locally, colonists from Russia and Germany were  
444 established in Olavarría from AD 1878 within a national strategy to inhabit the



445 Pampean region. Intensification of the farming activities to obtain wheat, potatoes, corn  
446 and vegetables transformed the original landscape (Pedrotta et al., 2012).

447 In the uppermost part of the sequence (7.5cm; from ca. 1940), higher and  
448 relatively steady values of lake level with episodes of high energy are inferred. The  
449 sedimentary properties of Unit 6 and by the presence of disarticulated ostracod valves  
450 indicated events of higher mechanical disturbance. However, this inference in our  
451 record did not agree with regional reconstructions, which recognized changes in  
452 hydrovariability. A regional increase of precipitation in AD 1940 was detected in  
453 instrumental records (Garreaud, 2009; Guerra et al., 2017; Pasquini et al., 2006).  
454 Similar hydroclimatic tendencies were also recognized in different paleolimnological  
455 records from the Pampean plain (Córdoba et al., 2014; Fontana, 2005; Laprida and  
456 Valero-Garcés, 2009; Piovano et al., 2009; Stutz et al., 2014). However, at a local scale,  
457 intensive periods of flooding and droughts in the western part of the Buenos Aires  
458 province led to agriculture losses and hydrological works (Monachesi and Albaladejo,  
459 1997). The sluice gate construction in the 1950s might have effectively controlled lake  
460 levels in Laguna Blanca Grande and might be the origin of Unit 6 and the disarticulated  
461 valves of ostracods.

462 Although our paleoecological reconstruction agrees well with previous  
463 inferences, historical documents and instrumental records, the timing of the  
464 reconstruction in the upper part of the record (from the last  $^{14}\text{C}$  date to the top of the  
465 core) should be considered with caution because the age control points here are sparse.  
466 A higher number of independent  $^{14}\text{C}$  tie points or a  $^{210}\text{Pb}$ - $^{137}\text{Cs}$  chronology in the upper  
467 part of the sediment would provide a more accurate chronology for the recent human  
468 impact.

469

## 470 **6. Conclusions**

471 Overall and despite the low resolution of this sedimentary record, our paper  
472 contributes to increasing the spatial resolution in the Pampean plain in terms of both  
473 humidity and nutrient enrichment, as well as to understand the role and trends of natural  
474 variability versus anthropogenic impact in the last centuries. Human activities started in  
475 this region around ca. AD 1800 with the official foundation of the first towns (Pedrotta  
476 et al., 2012) but anthropogenic impacts were not evident until the twentieth century,

477 when agricultural expansion (Monachesi and Albaladejo, 1997) led to nutrient  
478 enrichment and the establishment of a new diatom assemblage dominated by *A.*  
479 *granulata* var. *angustissima* and by fragilarioid taxa. The main periods of hydrological  
480 variability, notably the humid phase (ca. AD 1450) and progressive drier conditions  
481 mirror previous reconstructions in the region. Recent alterations of the hydrological  
482 cycle (ca. AD 1950) are consistent with further anthropogenic impacts in Laguna  
483 Blanca Grande, already shown by the nutrient enrichment. High-resolution studies  
484 would improve our understanding of complex climatic patterns operating in this zone.  
485 However, further studies on taxonomy and autecology are needed to refine the  
486 paleoecological interpretations based on biological proxies and to effectively unlock the  
487 information contained in its sediment records.

488

489

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845 **Tables**

846 Table 1. Lithological description and facies characterization of the composite core LBG

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<b>Facies</b>	<b>DEPTH (cm)</b>	<b>DESCRIPTION</b>	<b>SEDIMENTARY UNITS</b>
Facies 1	0-7	Massive, very fine sandy coarse silt, dark brown color, saturated in water, rich in organic matter, abundant plant macro remains.	Unit 6
Facies 2	7-11	Very fine sandy very coarse silt, brown color, with massive structure. It has diffuse planar contact with the underlying unit..	
Facies 3	11-25.5	Massive, very fine sandy coarse silt, dark brown, with iron mottles, It has indistinct contact with the underlying unit	Unit 5
Facies 4	25.5-26	Very fine sandy very coarse silt, dark grayish brown color, with massive structure. It has indistinct contact with the underlying unit	
Facies 5	26-31.5	very fine sandy coarse silt, dark brown color, with massive structure. It has indistinct contact with the underlying unit	Unit 4
Facies 6	31.5-52	Very fine sandy very coarse silt, brown color, with massive structure. It has indistinct contact with the underlying unit	
Facies 7	52-67.5	Very fine sandy very coarse silt, black color, with high bioturbation. It has a diffuse planar contact with the overlying unit.	Unit 3
Facies 8	67.5-75.0	Very coarse-silty fine sand, lightly laminated, with thin light brown sandy (1-2 mm) laminae interbed in dark-organic bed, very dark brown color. It has a planar sharp contact with the overlying unit.	Unit 2
Facies 9	75.0-81.5	Very fine sandy very coarse silt. Presence of root remains, and light bioturbation, very dark brown color. It has a planar sharp contact with the overlying unit.	
Facies 10	81.5-93	Very fine sandy very coarse silt with a higher proportion of silt than the overlying unit, brown-black color. It has a planar sharp contact with the overlying unit.	Unit 1
Facies 11	93-104	Very fine sandy very coarse silt, black color. It has a diffuse planar contact with the overlying unit	

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854 **Figure captions**

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856 Figure 1. Location map of the study site. A) Location of the study site in South  
857 America, showing the location of the main paleolimnological studies in the region: 1)  
858 La Brava (Irurzun et al. 2014); 2) Lakes Nahuel Rucá and 3) Lonkoy (Stutz et al., 2010,  
859 2012); 4) Lake La Barrancosa (Plastani et al. 2018); 5) Lagunas Encadenadas del Oeste  
860 (Córdoba et al. 2012); 6) Lake Adela (Dangavs & Mormeneo, 2012); 7) Laguna del  
861 Monte (Dangavs & Pierrad (2013); 8) Lake Melincué (Guerra et al. 2015) and 9)  
862 Laguna Mar Chiquita B) Photograph of the lake and C) Satellite image of the lake  
863 showing the inlets and outlets.

864 Figure 2. Correlation and physical properties of the Laguna Blanca Grande sediment  
865 cores. From left to right: cross-correlation between LBG-A and LBG-B, dark red  
866 rectangles show the depth of radiocarbon samples; sedimentary facies; lithology;  
867 lithological description; granulometry; organic matter (LOI), carbonate content and  
868 sedimentary units of the resulting composite core LBG.

869 Figure 3. Chronological model for Laguna Blanca Grande sequence based on three  
870 AMS  $^{14}\text{C}$  dates. The upper panel shows a table with the radiocarbon dates on Laguna  
871 Blanca Grande analyzed at 14CHRONO Centre for Climate, the Environment, and  
872 Chronology School of Geography, Archaeology, and Palaeoecology at the Queen's  
873 University Belfast (UBA code). Calibrated dates showed in the table were obtained in  
874 OxCal applying the ShCal13 calibration curve. Material dated: bulk sediment. The  
875 sample not used in the model is indicated in red. In the bottom panel, the Bayesian age  
876 model showing the calibrated  $^{14}\text{C}$  dates (transparent blue) and the age-depth model  
877 (darker greys indicate more likely calendar ages; grey stippled lines show 95%  
878 confidence intervals; red curve shows single 'best' model based on the weighted mean  
879 age for each depth) (Blaauw and Christen, 2011).

880 Figure 4. In the upper panel, the relative abundance of diatom species and Crysophyte  
881 cysts in LBG sediment core. Diatom zones (DT1-4) are based on the indicated cluster  
882 constrained analysis. In the bottom panel, a DCA ordination of (black circles), showing  
883 presenting the ecological space occupied by samples (red diamonds).

884 Figure 5. Relative abundance of the cladocerans in Laguna Blanca Grande sediment  
885 core. Cladocera zones (CL1-3) are based on the indicated cluster constrained analysis

886 Figure 6. Absolute abundance of ostracods in Laguna Blanca Grande sediment core.  
887 \*Only juveniles were recorded. Zonation was defined using optimal division  
888 information content after applying a square-root transformation on the data set.

889 Figure 7. Comparison of proxies from the Laguna Blanca Grande sequence and selected  
890 local and regional palaeoclimate reconstructions in South America and the Pampean  
891 region. In the upper panel regional and Pampean reconstructions. In the lower panel  
892 compilation of the biological and sedimentological proxies together with  
893 historical/documentary data of anthropogenic activities in Laguna Blanca Grande. From  
894 upper to lower part of this panel: *Bosmina* sp. (%), ostracod biozones, sedimentary  
895 units, sedimentation rate, loadings from diatom DCA axis 1, paleoecological inferences  
896 about hydroclimatic variability/nutrient enrichment and historical data of land  
897 occupation and uses.

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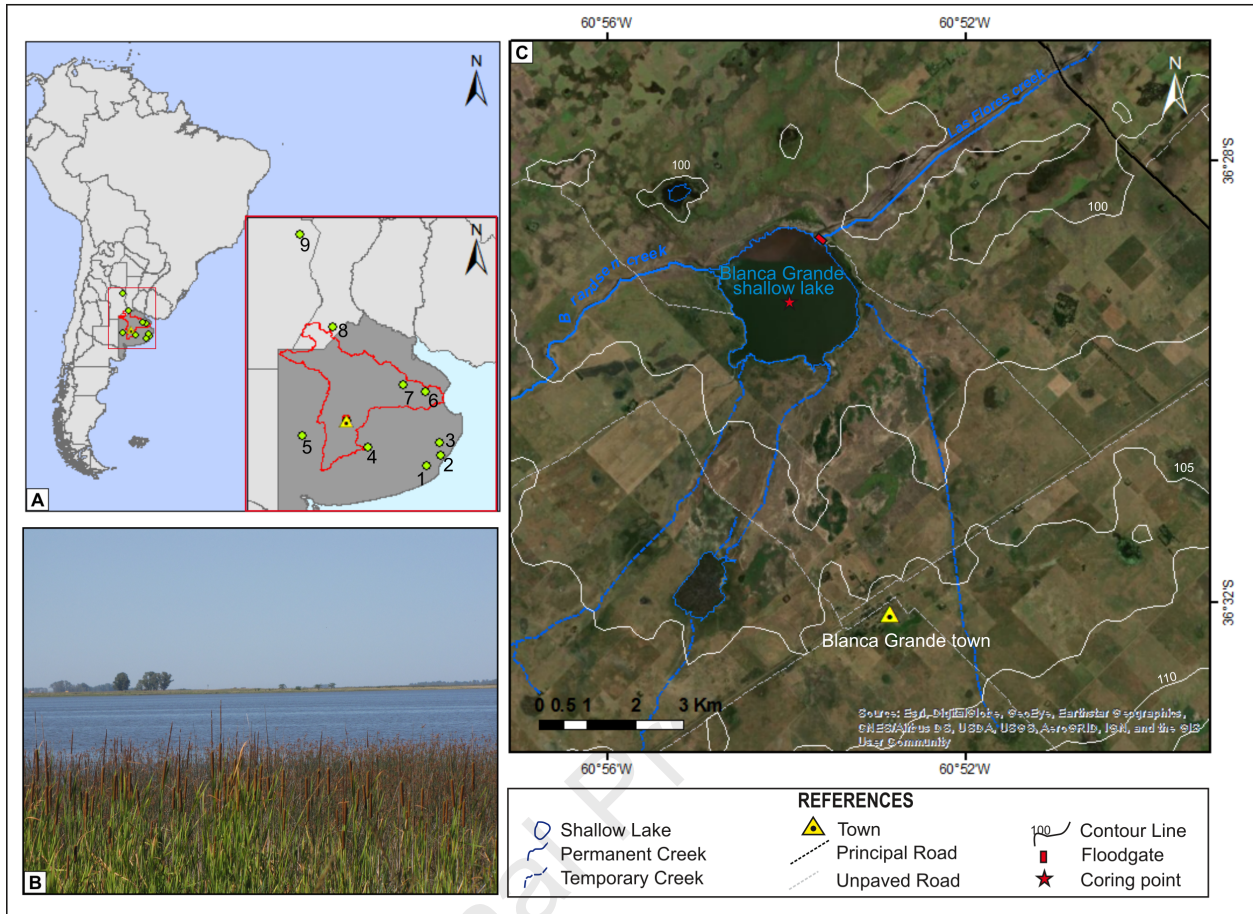
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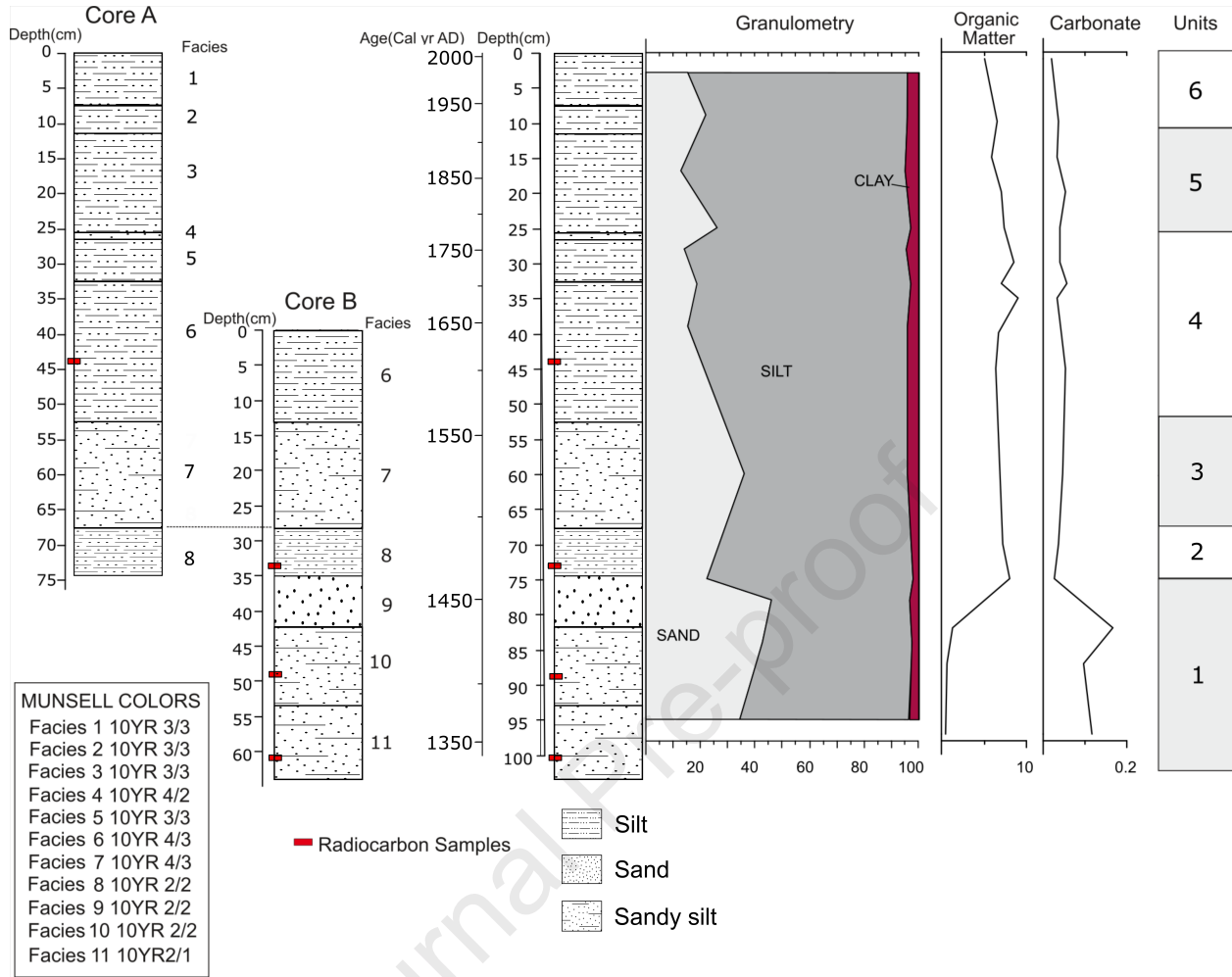
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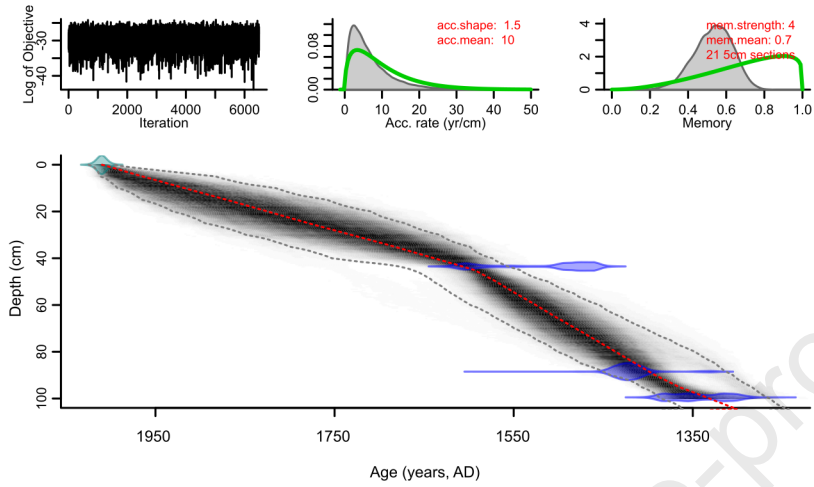
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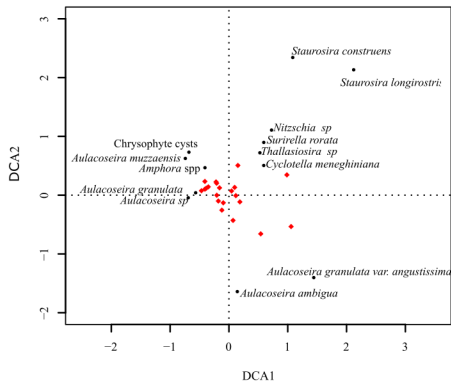
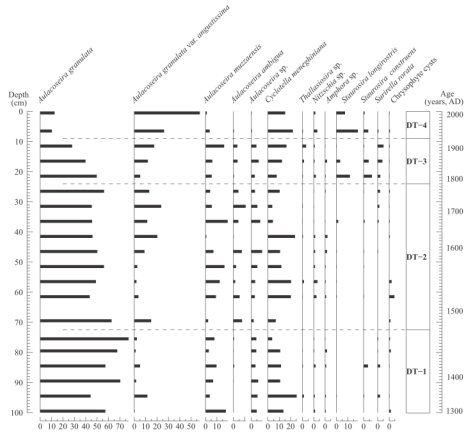


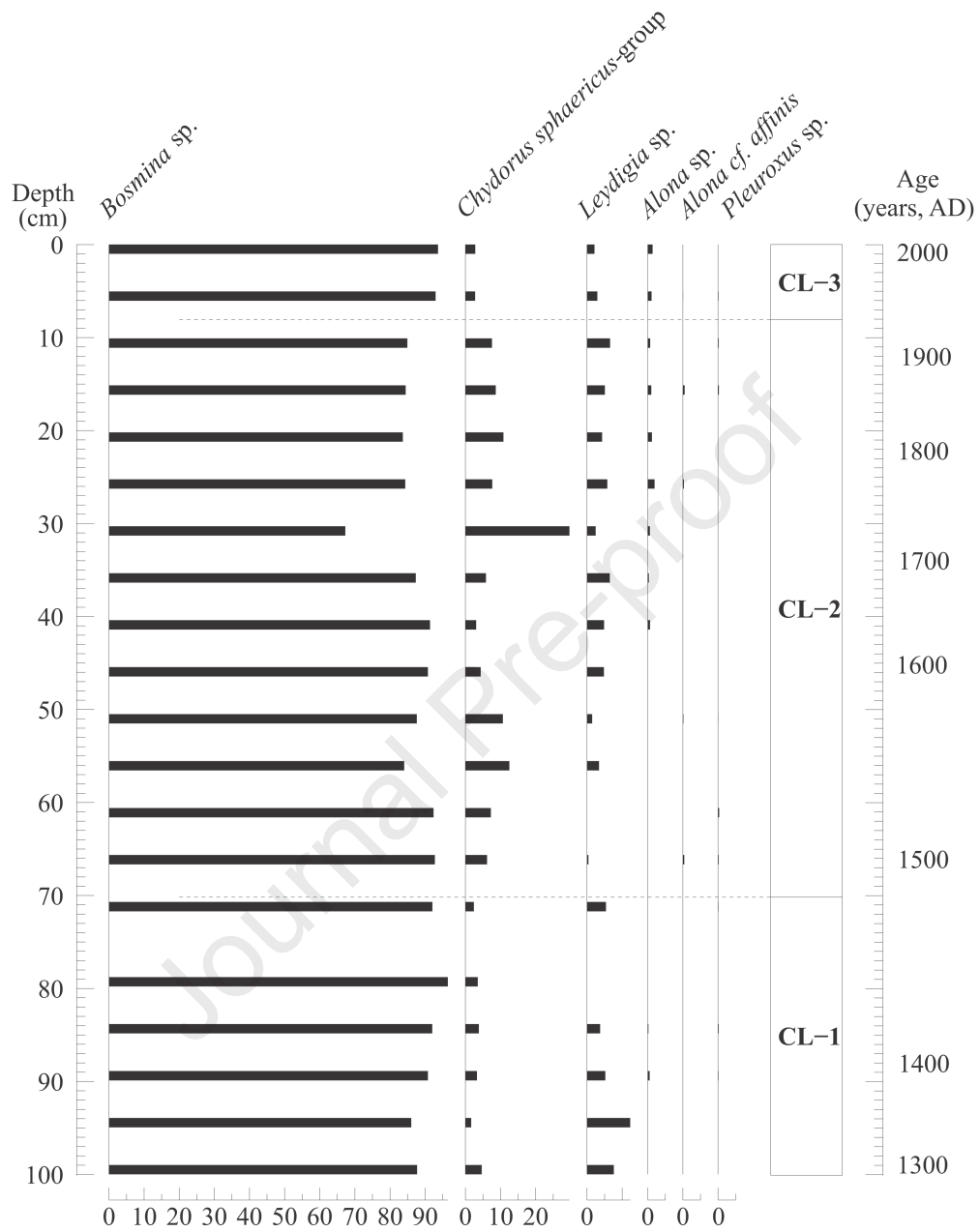


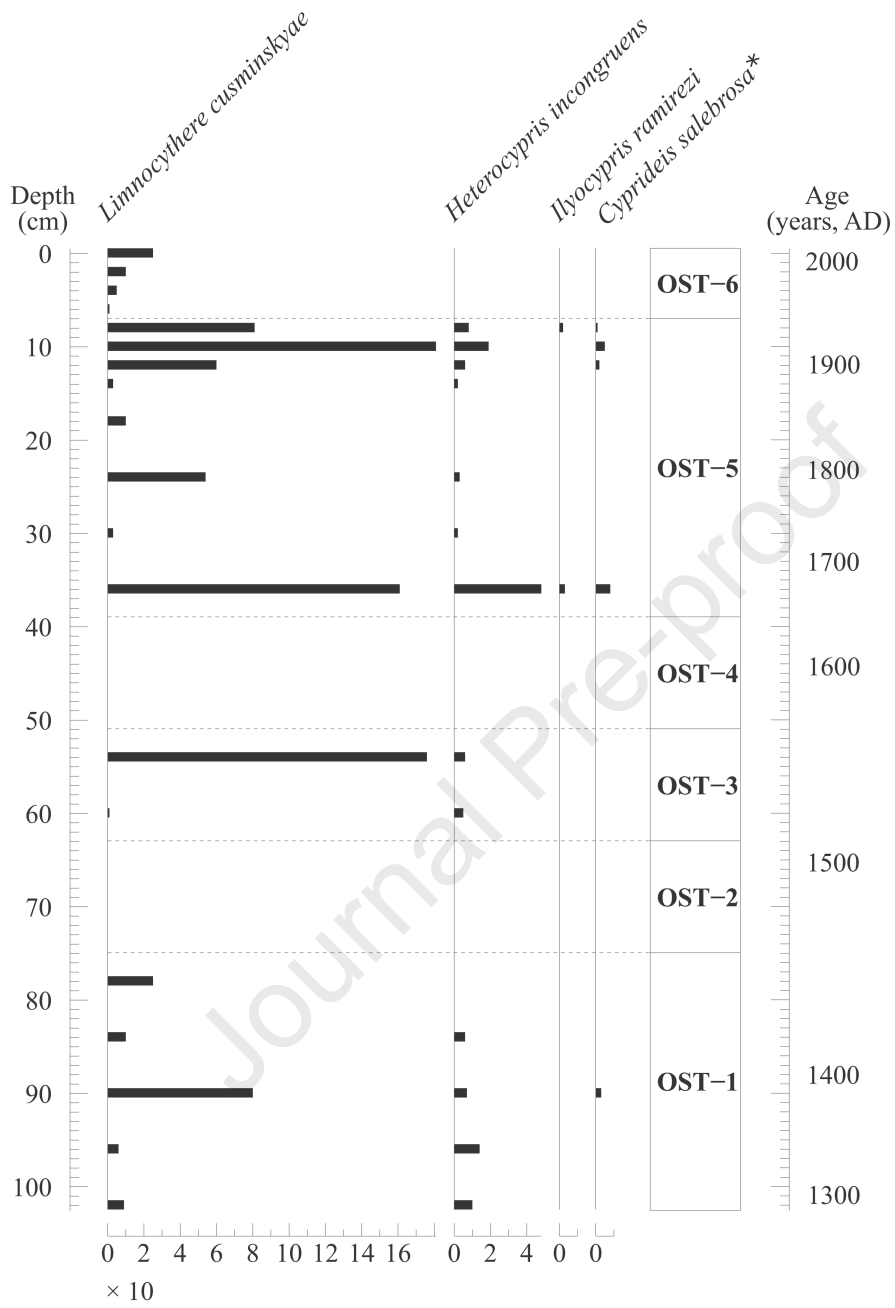


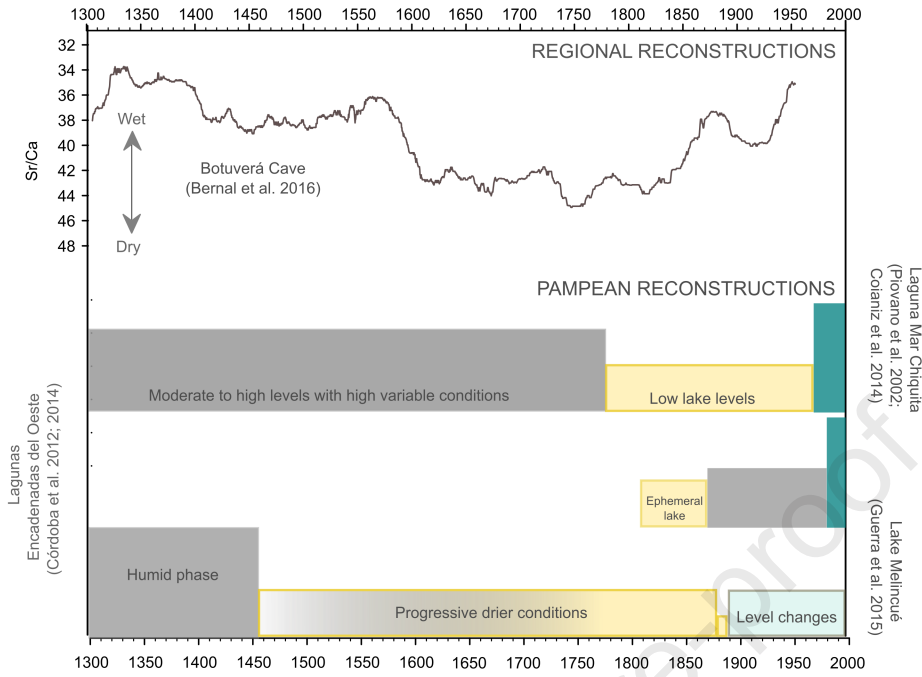
Composite core depth (cm)	Depth LBG-A (cm)	Depth LBG-B (cm)	Lab code	<sup>14</sup> C AMS age (yr BP)
43.25	43-43.5		UBA-30901	425±27
72.5		33-34	UBA-30902	1004±33
88.5		49-50	UBA-39237	549±29
99.5		60-61	UBA-23301	682±32



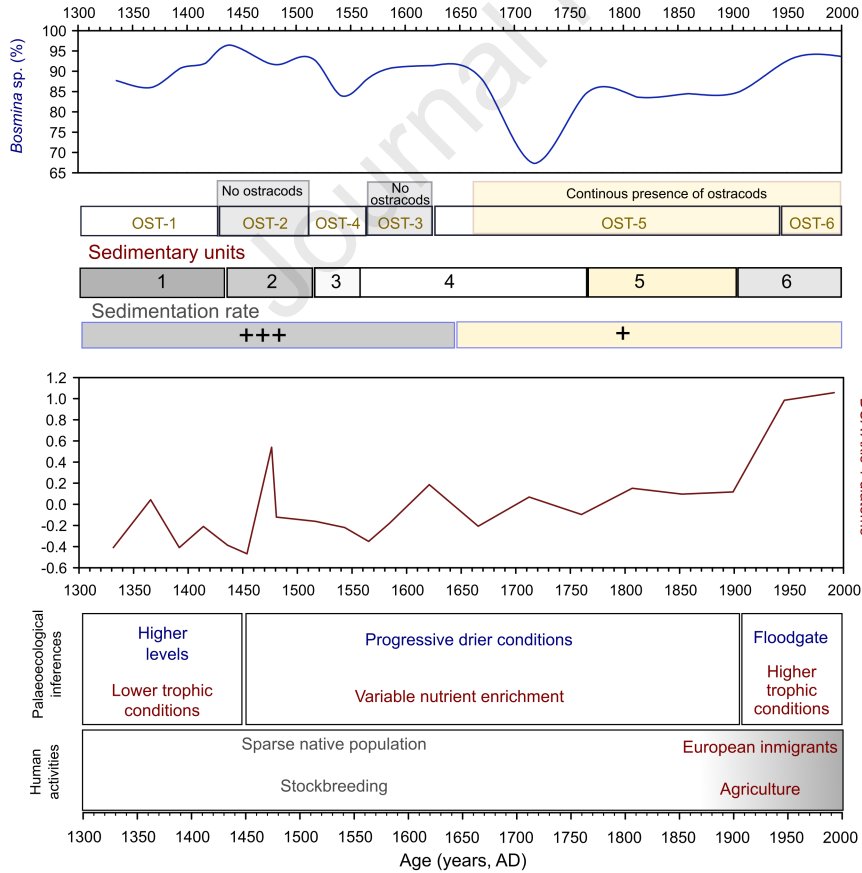








**LAGUNA BLANCA GRANDE (this study)**



## HIGHLIGHTS

- Lakes in the Pampean region contain important archives of past environments.
- Sediment properties and biological proxies were analysed in a 700-years record.
- Lakes level changes mirror past hydroclimatic variability.
- Anthropogenic activities induced a nutrient increase in the last century.

Journal Pre-proof



**Declaration of interests**

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests:

Journal Pre-proof