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**Human impact on the hydroenvironment of Lake Parishan, SW Iran, through the late
Holocene**

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34 **Abstract**

36 A multiproxy record from Lake Parishan, SW Iran, shows human impact on the lake and its
37 catchment over the last 4000 years. The Parishan record provides evidence of changes in lake
38 hydrology, from ostracod, diatom and isotope analyses, that are directly linked to human activity in
39 the catchment; recorded by pollen and charcoal and supported by regional archaeological and
40 historical data. The lake ostracod fauna is particularly sensitive to human induced catchment
41 alterations and allow us to identify changes in catchment hydrology that are due to more than a
42 simple change in precipitation: evaporation state. Oxygen isotope data from endogenic carbonates
43 follow these faunal changes but also displays a longer trend to more positive values through the
44 period, coincident with regional patterns of water balance for the late Holocene in the eastern
45 Mediterranean.

48 **Keywords**

49 Iran, agriculture, late Holocene, lake, pollen, ostracods

51 Introduction

52 There is compelling evidence from many regions on Earth that people have had a substantial
53 impact on the environment for thousands of years (Ruddiman et al., 2015) e.g. via deforestation
54 (Roberts, 2013) or irrigation (Magee, 2005). Less clear, and pertinently for this volume, is
55 determining at which point people made a global impact such that a 'golden spike' marking the
56 beginning of a new Anthropocene era might be identified (e.g. Gale and Hoare, 2012; Smith and
57 Zeder, 2013). Palaeoenvironmental archives such as lake sediments can often only provide a
58 relatively local or regional view of past environmental change. Importantly, however, lacustrine
59 archives can preserve multiple proxies of change, allowing information on climate, environment
60 and their drivers, including human activity, to be compared directly from one sediment record. This
61 avoids the issues of dating error or archival systematics that can complicate human-climate-
62 environment comparisons from multiple sites, environmental and/or archaeological (e.g. Jones,
63 2013a). Despite clear evidence of past human activity that could alter the environment (e.g. Smith
64 and Zeder, 2013; Ramsey et al., this volume), multiple proxies are needed to establish clear links
65 between this activity and recorded environmental change. Moving past correlation to causation is
66 difficult in the palaeosciences, irrespective of whether the environment is affecting people or the
67 other way round, but is important for a robustly defined Anthropocene.

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69 Here we present a new record of environmental change from Lake Parishan in the Fars Province
70 of South West Iran through the last 4,000 years. We use multiproxy data from a single core to
71 establish the dominant forcing factors of the environment in and around the lake. South West Iran
72 has a long history of human occupation with cereal agriculture and animal domestication dating
73 back to around 10,000 years BP (Weeks, 2013a; Riehl et al., 2013). The Zagros, and their 'Hilly
74 Flanks' have long been of interest to debates about the origins of agriculture (Braidwood and
75 Braidwood, 1949) and people's interaction with their environment in general (e.g. Miller, 2013).
76 Despite this, continuous palaeoenvironmental records from Iran, and especially from the south and
77 east, are scarce. Current palaeoenvironmental and palaeoclimatic understanding for the Holocene
78 of Iran is drawn largely from the records of Lakes Urmia, Zeribar and Mirabad (e.g. Djamali et al.,
79 2008; Stevens et al., 2006), hundreds of kilometres to the north of Parishan (Fig. 1). These sites sit
80 in a different climate regime to Parishan today (Jones, 2013b) and therefore may not reflect past
81 change further south. Our new data provide a more local reconstruction of environmental change
82 to compare directly with local archaeological investigations (e.g. Potts et al., 2009). A pollen record
83 of the past 5000 years does exist from Fars, from Lake Maharlou (Fig. 1) (Djamali et al., 2009), but
84 largely provides information on human-induced landscape change in the catchment, and not
85 hydroclimatic changes of the lake itself.

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87 The Lake Parishan multiproxy data set allows us to narrow down the possible explanations for the
88 changes seen in the Parishan record, with data describing environmental changes within the lake

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2 89 (ostracods, isotopes, diatoms, pollen) and within the catchment (pollen, charcoal, in-wash proxies
3 90 such as magnetic susceptibility). The record provides new palaeoenvironmental information from
4 91 the region as a whole, and shows that people have had a significant impact on lake hydrology at
5 92 various times during the last 4000 years.
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94 **Site Description**

95 Lake Parishan (29.5°N 51.8°E, 820 masl) lies in a fault-bounded basin 15km southeast of Kazerun,
96 in the Fars region of SW Iran (Fig. 1). As one of only a few recently extant lakes in the region Lake
97 Parishan is an important freshwater site and is Ramsar listed (No. 37; 23/06/1975). Over the last
98 25 years, lake area has fluctuated between 0 – 52 km², with a corresponding maximum depth of 0
99 to 5m (UNDP/GEF, 2010). The surface catchment is 270 km². The lake has no surface outflow and
100 given local average annual precipitation (450 mm) and evaporation (2400 to 3100 mm/yr) regimes
101 (Lotfi and Moser, 2010) the lake is liable to drying. The lake dried out completely in 1987
102 (UNDP/GEF, 2010) and again since our fieldwork of 2007. Lake waters had a pH of between 8.5
103 and 9 in 2001/2002 with variable conductivity values between 3500 and 8900 µS/cm (Lotfi and
104 Moser, 2010). Na or Mg and Cl are the dominant ions in the lake. Spring samples show more
105 variability, with water more likely to be Ca – HCO₃ dominated (Shirini Feshan, 2000).
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108 The lake and its catchment are an important agricultural centre with over 800 wells (UNDP/GEF,
109 2010) exploiting its important groundwater resource. Given recent drops in lake level there is
110 significant local effort being put into understanding the lake system (e.g. Lotfi and Moser, 2010;
111 UNDP/GEF, 2010). Lake Parishan's clear sensitivity to changes in the catchment water balance in
112 recent years makes it a potentially useful site for observing past changes in environment.

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113 **Methods**

114 *Fieldwork*

115 Core LP111 was taken in February 2007 using a Livingstone corer (Livingstone, 1955) from the
116 south-central part of the lake (29.514°N, 51.800°E) in 2.1m water depth. Four drives retrieved a
117 ~2.5m core sequence with some small gaps between drives (Fig. 2). In addition, water samples
118 were taken in leak-proof plastic bottles, at arm's length below the surface, for the analysis of the
119 oxygen and hydrogen isotope composition of lake water to help constrain the interpretation of the
120 palaeo-isotope data recorded in core carbonates.
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122 *Sedimentology and Geochemical proxies*

123 Before the cores were sampled magnetic susceptibility was measured every 2cm on a Bartington
124 MS2C Core Logging Sensor with a 60 cm loop, with data corrected for core diameter and drift
125 using the accompanying Multisus software. Loss on Ignition (LOI) at 550°C and 925°C (e.g. Hierl

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2 126 et al., 2001) was undertaken at a 4cm resolution, providing data on changes in organic and
3 127 inorganic carbon content in the core.
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5 128 1cm (sediment length) samples from every 4cm down the core were prepared for isotope analysis
6 129 of sedimentary, endogenic, carbonates at the NERC Isotope Geosciences Facility (NIGF) following
7 130 standard laboratory protocols (e.g. Leng, 2005). Samples were left in 5% sodium hypochlorite
8 131 overnight and then sieved at 90µm in deionised water to remove shell carbonates i.e. from
9 132 ostracods (see below) and snails. Dried samples were reacted in acid at 25°C under vacuum and
10 133 the resulting CO₂ collected for analysis of oxygen and carbon isotope ratios (¹⁸O/¹⁶O, ¹³C/¹²C) on
11 134 an Optima dual-inlet mass spectrometer. δ¹⁸O and δ¹³C values, reported in standard delta units as
12 135 part per thousand deviations from the VPDB standard, have analytical reproducibility of 0.1‰.
13 136 Following preparation for isotope analysis, selected carbonate samples were analysed by XRD in
14 137 the Faculty of Engineering, University of Nottingham to record the carbonate mineralogy. Finely
15 138 ground samples were analysed in cavity mounts (Hardy and Tucker, 1988) on a Siemens D500 X-
16 139 Ray diffractometer. The scanning range was 5-65° 2θ and the scan rate was 2° 2θ per minute with
17 140 a step size of 0.05.
18 141 Bulk organic components of the sediment were analysed for %C, %N and δ¹³C_{organic} at the same
19 142 resolution as the LOI and carbonate isotope data, by combustion in a Costech ECS4010 on-line to
20 143 a VG TripleTrap and Optima dual-inlet mass spectrometer at the NIGF. Samples were first reacted
21 144 in 5% HCl overnight and then thoroughly washed, to disaggregate the sediment and remove
22 145 carbonates. The isotope data are reported in the standard delta units (δ¹³C) as parts per thousand
23 146 deviations from the VPDB standard. Analytical reproducibility is <0.1‰ for δ¹³C and 0.2 for C/N for
24 147 the standard and sample material.
25 148

149 *Pollen*

150 Twenty six samples were treated for pollen analysis following the classical extraction technique
151 described in Moore et al. (1991). An outline diagram at 8cm resolution was first produced, with
152 additional samples then counted between 104 and 132 cm to increase this section to 4cm
153 resolution. Pollen grains were identified using the pollen reference collection developed for Iranian
154 flora at the Institut Méditerranéen de Biodiversité et d'Ecologie (Aix-en-Provence, France) but also
155 the available pollen bibliography on the Mediterranean region, Europe and the Middle East (e.g.
156 van Zeist and Bottema, 1977; Reille, 1992, 1995, 1998; Beug 2004). Pollen typification followed
157 Beug (2004). About 300 pollen grains excluding the pollen of aquatic plants were counted from
158 each sample layer and the percentage values were calculated and plotted using Tilia and TGView
159 software (Grimm, 2004/2005).
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161 The full pollen record from core LPIII will be discussed elsewhere (Djamali et al., in press) and here
162 we present selected taxa in order to reconstruct the environment in and around the lake through
163 the time period of study, in particular the extent of human impact on the catchment and the

164 hydrological variations of the lake. We use the presence of *Olea* (olive) and *Platanus* (plane tree)
165 pollen as indicators of agricultural practice (following Djamali et al., 2009, 2011). We also present
166 the *Quercus* (oak) pollen curve as a measure of the natural vegetation regime. In addition,
167 *Sporormiella* (fungi associated with animal dung (van Geel, 2001)) is used as evidence of pastoral
168 activities. *Sparganium*-type pollen and *Riella* spores provide information on lake level changes
169 (Djamali et al., 2008), while charcoal is a measure of burning in the region, often associated with
170 agricultural practices in the later Holocene (e.g. England et al., 2008), but also potentially of natural
171 origin (e.g. Turner et al., 2007).

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173 *Ostracods*

174 Ostracod faunal assemblages were determined from 40 levels within the core (at a 4cm resolution)
175 in order to characterize the hydrological and hydrochemical evolution of the lake, following
176 methods in Griffiths and Holmes (2000). At each level, ostracod valves were picked from the
177 weighed coarse (>90µm) fraction of samples that had been processed for endogenic carbonate
178 isotope analyses. Ostracod specimens were picked under a low-power binocular microscope and
179 stored in micropalaeontological slides. All of the adult and later-moult-stage valves were recovered
180 from each sample: although the occurrence and preservation of the smallest instars were noted,
181 these moult stages were neither picked nor counted owing to difficulty in identification.
182 Identifications followed Meisch (2000). Valves that were insufficiently complete to allow
183 identification were not picked although the approximate abundance of broken valves in each
184 sample was noted. Ostracod species occurrences were expressed as number of valves per gram
185 of dry sediment, and as percentages.

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187 *Diatoms*

188 A preliminary set of samples for diatom analysis were prepared from 15 levels; approximately
189 every 16cm through the core sequence. These were treated to remove organic matter and
190 carbonates, using standard procedures. The weight of the dried sample was recorded to allow for
191 an estimation of valves per gram. The prepared samples were mounted onto coverslips using
192 Naphrax resin and studied at 1000 x magnification using either an Olympus CX41 or a Zeiss
193 Axioskop2 Plus microscope. Selected levels were examined using a JEOL 6400 Scanning
194 Electron Microscope in the Faculty of Engineering, University of Nottingham. Identifications were
195 made using standard diatom floras including Krammer and Lange-Bertalot (1986 and 1988a and b),
196 Patrick and Reimer (1966 and 1975), and the regional study of Witowski et al. (2008). Ecological
197 interpretations were based mainly on Gasse (1986); Reed (1998); Reed et al., (2012) and
198 information in the European Diatom Database (<http://craticula.ncl.ac.uk/Eddi/jsp/index.jsp>).

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200 **Chronology**

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2 201 Four age estimates, 3 from radiocarbon analysis of bulk organic material (Table 1) and one from U-
3 202 Th analysis (Table 2), have been carried out on the 250cm core. Radiocarbon analyses were
4 203 undertaken at the Poznan Radiocarbon Laboratory. The U-Th analysis was undertaken at the
5 204 Open University, UK. Following acid digestion of bulk sediment samples U and Th were purified
6 205 using ion exchange chromatography before being run on a Nu Instruments Multi-collector ICP-MS
7 206 in a solution of 3% HNO₃, using a sample - standard bracketing technique. Based on these age
8 207 estimates, the core spans from present day (the lake was extant at the time of coring) to around
9 208 4,000 cal BP.

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15 210 The paired set of radiocarbon and U-series age estimates from 225cm were analysed to help
16 211 establish if there was a significant old carbon effect on the radiocarbon ages, because Lake
17 212 Parishan is in a carbonate catchment. Although there were also issues of contamination with the
18 213 U-Series age estimate (Table 2), the use of a standard Open University laboratory correction for
19 214 lake sediments of this type allowed the calculation of an old carbon error of approximately 360
20 215 years for the sequence.

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26 217 The clear shift in the pollen diagram at 125cm, with the establishment of olive (*Olea*) and
27 218 deforestation of oak (*Quercus*), likely marks the start of the Achaemenid Persian Empire 2500 cal
28 219 BP (e.g. Djamali, 2009), which had a significant centre at Persepolis, only 110 km from Parishan
29 220 (see further detail in the Discussion below). This pollen stratigraphic marker confirms the estimate
30 221 of old carbon error for core LP111 (Fig. 2).

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35 223 Our final age model (Fig. 2) is therefore based on the radiocarbon age estimates and a 360 year
36 224 old carbon correction. The dating control on this record from Parishan is important when comparing
37 225 the data with other sites, and the old carbon issues identified here may have implications for the
38 226 age models for other sites in the Zagros such as Zeribar and Mirabad (Stevens et al., 2001, 2006;
39 227 Wasylkova et al., 2008). However, for this paper it is the relationship between proxies from the
40 228 same core that is vital to the discussion, and the temporal relationships between them are
41 229 internally robust.

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47 231 **Results**

48 232 *Sedimentology and Geochemical proxies*

49 233 The data show a number of trends and events through the core (Fig. 3). There is a peak in
50 234 magnetic susceptibility between 200 and 170cm, also marked by an increase in non-combusted,
51 235 residual material from the LOI analyses. The amount of this residual material generally decreases
52 236 up through the core, with a particular step change to lower values between 130 and 110cm. This
53 237 depth interval is also marked by a reduction in magnetic susceptibility, to negligible values, and an
54 238 increase in both organic and inorganic carbon. This step change is part of a long term trend to

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2 239 higher values, from 200cm towards the top of the core, in the amount of organic carbon. C/N ratios
3 240 show a marked shift from values around 14 to lower values, of around 10, at 110cm. These data all
4 241 suggest a general trend to reduced amounts of catchment material, both in terms of inorganic in-
5 242 wash (reduction in residual LOI material and magnetic susceptibility) and non-aquatic organic
6 243 matter, C/N values of 10 are typical of aquatic algae (e.g. Leng et al., 2010), through the core.
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11 245 Carbonate mineralogy is variable down core with samples at 128 and 224cm dominated by calcite,
12 246 samples at 80 and 112cm having a mixture of calcite and aragonite, and samples from 16 and
13 247 64cm dominated by aragonite. The carbonate isotope values also show an up core trend to more
14 248 positive values with a particularly marked step in $\delta^{13}\text{C}$ values at 125cm from $\sim -3\text{‰}$ to $\sim +2\text{‰}$, in
15 249 part explained by the shift in mineralogy; aragonite is 1.9‰ (0.6‰) more positive in $\delta^{13}\text{C}$ ($\delta^{18}\text{O}$)
16 250 compared to calcite precipitated in waters with the same temperature and isotopic composition
17 251 (Grossman, 1984; Grossman and Ku, 1986). Oxygen and hydrogen isotope ratios of the lake
18 252 waters collected in 2007 show a clear evaporative signal (Supplementary Figure 1), and $\delta^{18}\text{O}$
19 253 trends to more positive values likely represents a shift to more evaporative dominant conditions
20 254 given the associated change from calcite to aragonite (e.g. Jones et al., 2006). However, there is
21 255 no co-variation between $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ from the carbonates in LP III (Supplementary Table 1) that
22 256 is typically seen in evaporative lake systems (e.g. Li and Ku, 1997) suggesting an alternative
23 257 control, other than evaporation, on the $\delta^{13}\text{C}$ system. The isotope value of bulk organic matter is
24 258 similar (between -21 and -23‰) until the top 20cm of the core, where it shifts to lower values, and
25 259 it remains unclear what would cause the changes in $\delta^{13}\text{C}_{\text{carbonate}}$, other than changes in carbonate
26 260 mineralogy as discussed above, without impacting on the total dissolved inorganic carbon pool
27 261 also used by the organic material.
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38 263 *Pollen*

39 264 Variations of oak (*Quercus*) and pistachio (*Pistacia*) pollen indicate that the regional forest
40 265 vegetation did not significantly change from the beginning of the sequence (pollen zones LP III-A
41 266 and LP III-B) up to about 120cm (pollen zone LP III-C) where oak pollen decreases while there is a
42 267 significant increase (from 0-1% to 6% of the terrestrial pollen sum) in pollen of cultivated trees
43 268 (*Olea* and *Platanus*) and steppe plants (upland herbs) (Fig. 4). The increasing percentages of the
44 269 pollen of upland herbs in LP III-C is matched by increasing values of the Cerealia-type and
45 270 *Plantago lanceolata*-type pollen (anthropogenic herbs in Fig. 4) as well as dung-associated
46 271 *Sporormiella* (Non-Pollen Palynomorphs). The pollen zone of LP III-C, which represents intensified
47 272 agro-sylvo-pastoral activities in the catchment, is also coincident with a sudden increase of pollen
48 273 of aquatic plants such as *Sparganium*-type. Later, in LP III-D, pastoral activities likely reach their
49 274 maximum, between approximately 70cm to 40cm depth, as this is the period of highest values of
50 275 *Plantago lanceolata*-type pollen and *Sporormiella* spores. Microcharcoal also increases from
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2 276 around 120cm depth, peaking at 60 and 30cm, with occasional peaks lower in the sequence e.g. at
3 277 212cm. Pollen is not well preserved in the uppermost 30cm of the core.

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6 279 *Ostracods*

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8 280 Ostracod shells are found in all of the samples examined, although abundance varies markedly
9 281 from 15 valves (244cm) to 746 valves (228cm). All samples contain adults and a range of juvenile
10 282 instars: adults and juvenile moults are generally very well preserved, even the earliest instars. Most
11 283 of the adult specimens and the vast majority of the juvenile instars are present as disarticulated
12 284 valves, with only a small proportion of carapaces preserved. In total eight ostracod taxa were
13 285 recovered from LPIII (Fig. 5). The presence of adults and a range of juvenile moult stages suggest
14 286 that the assemblages are largely *in situ* with signs of minimal *post mortem* transport.

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17 288 In general, the lower part of the core, below about 100cm, is characterised by a higher abundance
18 289 of shells than above. Overall, the ostracod assemblages are consistent with a slightly saline lake.

19 290 *Cyprideis torosa* is the most common species, followed by *Limnocythere inopinata*, indeterminate
20 291 species 1, *Darwinula stevensoni*, *Candona* sp. juveniles and *Heterocypris salina*. The remaining
21 292 two taxa, indeterminate species 2 and 3, occur sporadically. Based on the stratigraphical
22 293 distribution of taxa, the most striking difference is between the lower part of the core, below
23 294 ~132cm, which is dominated by *C. torosa*, and the upper part dominated by *L. inopinata*. In the
24 295 lowest parts of LPIII, significant numbers of *L. inopinata*, *H. salina* and indeterminate species 1 and
25 296 lesser numbers of *Candona* sp. (probably *Candona* cf. *neglecta*) accompany *C. torosa*. An
26 297 exception to this pattern is at 244 cm, where *L. inopinata* dominates, although total specimen
27 298 numbers are small. In the upper part of the core, *C. torosa* is absent from most levels, although *H.*
28 299 *salina*, *D. stevensoni* and indeterminate species 1 occur in many levels. At 104 to 108cm, a
29 300 reversal to this pattern is found, with *C. torosa* dominant much like in the lower part of the core.

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32 302 *Diatoms*

33 303 Diatoms are only preserved in some sections of the Parishan core (Fig. 6), with most taxa
34 304 indicative of brackish and often shallow water conditions. Counts are low (ca. 100 valves) in spite
35 305 of counting multiple slides. Results from only 8 of the 15 samples were considered adequate to
36 306 interpret and the impact of differential preservation must be taken into account. The sequence can
37 307 be divided into roughly four sections: from the base to 144-145cm; the sample from 128-129cm; a
38 308 section of core where preservation was too poor to allow for counting (112-113 to 32-33cm) and
39 309 the top two samples (9-10 and 16-17cm). The lower part of the core was dominated by *Amphora*
40 310 spp. which appeared to be wrapped in silica (confirmed under the SEM). Some valves could be
41 311 identified as *A. coffeaformis* (*Halamphora coffeaformis*), a benthic, high salinity alkalibiont found
42 312 in both saline lakes and marine environments. Other diatoms present included benthic species
43 313 indicative of high salinity/brackish conditions such as *Campylodiscus clypeus*, *Mastogloia braunii*

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2 314 and *Anomoeoneis sphaerophora*. *Nitzschia granulata*, more usually a marine diatom, but also
3 315 occasionally recorded in salt lakes (Gasse, 1986) was noted. Resting spores of *Chaetoceros*
4 316 *amanita* were recorded in transect counts. These taxa are generally epipelagic or benthic and occur
5 317 predominantly in Na-Cl dominated waters of moderate to high conductivity (3000 - >30,000 $\mu\text{S cm}^{-1}$
6 318 in different data sets). The low diatom numbers and rather poor preservation indicate a highly
7 319 evaporated water body, inimical to good diatom preservation. A stress response of accreting silica
8 320 around valves has been recorded in a highly evaporated crater lake in Mexico (Metcalf, 1990).
9 321 This accretion of silica probably accounts for their numerical abundance in the fossil material and
10 322 may not reflect the importance of *Amphora* in the life assemblage. Overall, the conditions
11 323 indicated at the bottom of the core are consistent with recent water samples (see Site Description).
12 324

13 325 The sample from 128-129cm was distinctive because of its high diversity and more abundant
14 326 diatoms. *Nitzschia cf gracilis*, *Brachysira apopina* and *Cymbella (Navicymbula) pusilla* were the
15 327 most abundant taxa (although silica 'wrapped' *Amphora* spp. were still present). The increased
16 328 abundance of diatoms and their better preservation indicate that the lake may have become less
17 329 saline/alkaline, at least intermittently. *N. gracilis* may be planktonic and has a lower reconstructed
18 330 EC optimum than the other taxa found here (around 3000 $\mu\text{S cm}^{-1}$ according to Reed et al., 2012).
19 331 It may also indicate higher nutrient levels (Patrick and Reimer, 1975). High variability in salinity
20 332 may be indicated by the presence of *B. apopina* which has very high reconstructed EC optima (>
21 333 50,000 $\mu\text{S cm}^{-1}$) based on work in Spanish and Turkish lakes (Reed, 1998, Reed et al., 2012). An
22 334 increasing proportion of epiphytic taxa at this depth may indicate more aquatic vegetation near the
23 335 coring site. It is interesting to note that this sample has high percentages of both *B. apopina* and *C.*
24 336 *pusilla* which are both the only species in their genus that are halophilous.
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26 338 The next five samples in the core preserved very few valves, although the sample from 80-81cm
27 339 was again dominated by the 'wrapped' *Amphora* spp, so it appears that conditions again became
28 340 unfavourable for diatom preservation. Preservation is much better in the top sample (9-10 cm) than
29 341 in the one below, which is again dominated by 'wrapped' *Amphora* spp. with *C. clypeus*. *C.*
30 342 *amanita* resting spores again noted. Both these samples show a significant presence of *Epithemia*
31 343 *smithii* (ca. 20%) which may indicate the availability of more aquatic vegetation. Unfortunately, this
32 344 taxon is not recorded in available salinity reconstruction databases. Overall the diatom
33 345 assemblage indicates shallow, high conductivity water.
34 346

35 347 Discussion

36 348 There are strong and significant correlations between the different variables analysed for this study
37 349 (Supplementary Table 1). Based on these correlations, two end-member environmental states are
38 350 interpreted for the last 4,000 years in and around Lake Parishan and its catchment. These two lake
39 351 states are marked most clearly in the stratigraphic record by the shift between the two dominant

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2 352 ostracod taxa (Figs. 5 and 7) and the interpretation of the changes seen between these two taxa,
3 353 and associated shifts in other proxies, are key to our overall interpretation of the data set.
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6 355 Although *C. torosa* has a higher salinity tolerance than *L. inopinata* (e.g. Holmes, 1992), a
7 356 decrease in salinity midway through the sequence is not consistent with other proxies including the
8 357 positive $\delta^{18}\text{O}$ trend (more evaporation), change in carbonate mineralogy from calcite to aragonite
9 358 (more evaporation) and the disappearance of diatoms from the record. Collectively these latter
10 359 three proxies suggest an increase in evaporative enrichment above about 120cm. *Cyprideis torosa*
11 360 and *L. inopinata* also have differing preferences on a water composition (alkalinity: Ca) gradient
12 361 with *C. torosa* preferring Ca-enriched and alkalinity-deplete waters (alkalinity: Ca <1), and *L.*
13 362 *inopinata* preferring Ca-deplete and alkalinity-enriched conditions (alkalinity: Ca >1) (Forester,
14 363 1983; 1986). These two contrasting water types fall on separate evaporative pathways (e.g. Hardie
15 364 and Eugster, 1970) and so cannot be explained by differences in the degree of evaporative
16 365 evolution.
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18 367 Lake conditions supportive of *L. inopinata* occur in the latter half on the record from LPIII but also
19 368 occur in three distinct phases between ~4,000 and 3,000 cal BP (Fig. 7), with the event at ~3200
20 369 cal BP being marked more noticeably by a decrease in *C. torosa* rather than an increase in *L.*
21 370 *inopinata*. Of note is that the substantial shifts in other proxies for in-lake and catchment conditions
22 371 c. 2,200 cal BP and in the latter half on the record, e.g. reduced oak pollen, increasing aquatic
23 372 macrophyte vegetation (*Sparganium*-type pollen), increased charcoal and increased *Sporomiella*,
24 373 also occur in these three events during the 4th Millennium BP.
25 374

26 375 Without the addition of human activity it is difficult to develop a scenario that would lead to the
27 376 combination of changes in the environmental proxies observed. Tectonic activity may impact upon
28 377 groundwater flow, potentially changing both the amount and source of surface- and/or ground-
29 378 water to the lake. However, the Parishan system seems to shift, relatively frequently, between two
30 379 clear states. It seems unlikely that successive tectonic events would firstly change water flow
31 380 conditions and then reset to the original condition at the next tectonic event, and for this to happen
32 381 repeatedly. We have discussed above why a change in evaporation amount, as part of a climatic
33 382 precipitation – evaporation lake level control, cannot fully explain the shifts in all variables recorded.
34 383

35 384 Combined, these multi-proxy data therefore suggest that during periods of increased catchment
36 385 agriculture (increased burning, increased deforestation, increased animal dung), surface (reduced
37 386 in-wash) and groundwater flow were reduced to the lake. With an increase in precipitation relative
38 387 to surface and groundwater inflow from the carbonate catchment the amount of Ca in lake waters
39 388 was relatively reduced, leading to an increase in the alkalinity/Ca ratio and a shift in the ostracod
40 389 assemblage. As relatively less water entered the lake, lake levels fell and the waters become more
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2 390 evaporatively enriched (as the ratio of evaporation to total input to the lake, precipitation plus
3 391 surface-and ground-water inflow, increased) leading to higher $\delta^{18}\text{O}$ values and aragonite
4 392 deposition (rather than calcite). This fall in lake levels also led to more shallow water areas suitable
5 393 for *Sparganium* type reeds to grow. Increased lake productivity, as marked by increased organic
6 394 material in the sediments fits a hypothesis of increased nutrient in-wash into the lake during
7 395 periods of increased agriculture. The data therefore suggest that increased agriculture in the
8 396 catchment had a significant impact on lake hydrology and biogeochemistry. *Limnocythere*
9 397 *inopinata* and associated end members of other proxy ranges therefore mark periods when the
10 398 lake was impacted by human activity whereas *C. torosa* marks a more natural lake and catchment
11 399 scenario (Table 3).
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14 401 The impacted period immediately after 2400 cal BP, most likely associated with the beginning of
15 402 the Achaemenid Empire (see below), is very marked in the record and shows a different set of
16 403 inter-proxy relationships compared to the full record. The diatom flora at this point in the core
17 404 suggest a freshening of the system, as do lower $\delta^{18}\text{O}$ data. Other proxies do not show anything in
18 405 particular of note here that is not consistent with the overall patterns described above. This level in
19 406 the core (~ 130 cm) represents the onset of the most intensive agricultural phase in the LPIII proxy
20 407 record, marked especially by the rise in olive pollen. Given that diatoms disappear almost entirely
21 408 from the core in the levels above this it appears there may have been a very short freshening of
22 409 the system, perhaps associated with the initial catchment deforestation, at the onset of the
23 410 Achaemenid Empire. This increase in water flow into the lake during anthropogenic catchment
24 411 disturbance may, arguably, be a more typical response to human activity in a lake catchment (e.g.
25 412 Rosenmeier et al. 2002) but is only short-lived at Parishan, with longer term proxy shifts
26 413 suggesting the long term impact of agriculture is reduced amounts of water entering the lake.
27 414 Conditions through this phase are marked by the lowest oak pollen percentages in the record,
28 415 suggesting a possible threshold in catchment vegetation below which runoff is increased.
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31 417 **Lake Parishan and regional archaeological evidence**

32 418 Given the apparent impact of people on Lake Parishan and its catchment we review these new
33 419 palaeoenvironmental results against the regional archaeological framework.
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36 422 Northern Fars has a long history of agro-pastoral production, stretching back to the early Holocene
37 423 (Weeks, 2013b). The earliest pollen samples from the LPIII core, which cover the period from c.
38 424 4000 cal BP, include peaks in micro-charcoal at c. 3800 and 3500 cal BP (Fig. 7) and correlate
39 425 with increasing human activities in the landscape. These peaks equate to the archaeological
40 426 periods known as the Middle and Late Kaftari, which witnessed a dramatic increase in known
41 427 human settlement in Fars (Potts et al. 2009). However, the LPIII core does not show the evidence
42 428 for the early adoption of arboriculture in the Kaftari period that was seen clearly in the Maharlou
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2 428 core (Djamali et al. 2009, 2011). Following the Kaftari period, between c. 3500-2500 years BP,
3 429 archaeological surveys in Fars indicate a relatively widespread and significant reduction in the
4 430 number of known settlements (Potts et al. 2009). Although the site of Tall-e Jidun in the Kazerun
5 431 plain close to Lake Parishan appears from surface collection survey to have been occupied from c.
6 432 3500 years BP (Nobari et al. 2009), the corresponding section of the LPIII core shows few
7 433 indicators of human agro-pastoral activities or landscape modification, especially in the micro-
8 434 charcoal record.
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14 436 The spike in pollen from cultivated trees (primarily *Olea* and *Platanus*) in the LPIII core at 120cm
15 437 provides informative parallels and contrasts with existing archaeological and historical information.
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17 438 This period coincides with the rise of the Achaemenid empire and a “massive investment of energy
18 439 and organizational power to bring the Persian heartland under cultivation” (Henkelman 2013: 528).
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20 440 This includes archaeological evidence for the construction of various water control mechanisms in
21 441 the vicinity of the Persian capitals Pasargadae and Persepolis, including dams, barrages, and
22 442 irrigation canals, that significantly expanded the area that could be reached by irrigation
23 443 (Boucharlat 2013; Boucharlat et al. 2012; Sumner 1986). Texts from Persepolis attest to large-
24 444 scale agricultural and pastoral production supported by the availability of large numbers of
25 445 dependent labourers to harvest crops and maintain canals, as well as the efforts of local
26 446 independent farmers and herders. In particular, there is historical evidence for extensive cultivation
27 447 of fruit trees during the Achaemenid period and for the foundation of estates that had variable
28 448 functions, including orchards, parks, gardens, and hunting preserves (Boucharlat 2013: 513;
29 449 Uchitel 1997). One single administrative text from Persepolis lists more than 6000 fruit tree
30 450 seedlings to be planted in five estates, including apple, mulberry, pear, quince, date, pomegranate,
31 451 and olive (Henkelman 2013: 528). Interestingly, although historical sources from the Classical
32 452 period describe Fars as a fertile and well-watered region with many farms and gardens with fruit
33 453 trees, heavily wooded hills and river banks densely covered with plane trees and poplars (Sumner
34 454 1986: 17-18), they commonly note the absence or poor quality of olives and olive trees in the
35 455 general region of southern Iran (Djamali et al., in press). The pollen evidence from the LPIII core, in
36 456 particular the evidence for olive cultivation, thus provides a significant new strand of evidence for
37 457 the discussion of arboriculture in Achaemenid and post-Achaemenid Iran.
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49 459 The Kazerun region falls broadly within the area of the Achaemenid heartland (Henkelman 2008)
50 460 and it is likely to have witnessed significant occupation at this time, as can be documented not only
51 461 in the immediate hinterlands of Persepolis and Pasargadae, but also c. 50 km to the north of
52 462 Kazerun in the Mamasani District (Askari Chaverdi et al. 2010; Potts et al. 2009). Unfortunately,
53 463 Achaemenid settlement in the immediate vicinity of Lake Parishan remains poorly understood.
54 464 Achaemenid occupation has been recorded at a number of sites in the Kazerun plain, and there is
55 465 a large Achaemenid settlement recorded at the mound of Tall-e Jidun (Nobari et al. 2009, 2012).
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2 466 However, the region is more famous as the location of the important Sasanian city of Bishapur, and
3 467 increasing human use of the landscape in the Sasanian and early Islamic periods (c. 1700 years
4 468 BP onwards) (Keall 1989; Calmard 2013) is supported by the increasing micro-charcoal and
5 469 *plantago lanceolata*-type pollen counts in the upper sections of LP111, if not by significant evidence
6 470 of tree cultivation (Djamali et al., in press).
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472 **Summary**

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11 473 The multi-proxy sequence from Lake Parishan provides a case study of clear human impact on a
12 474 lake environment, marked in the stratigraphic record. It is not uncommon for lake records from the
13 475 later Holocene to show such impact, especially in the pollen data (e.g. England et al., 2008;
14 476 Djamali et al., 2009). The Parishan record is rare in that changes in lake hydrology can be linked to
15 477 these human impacts over much of the last 4000 years. The ostracod fauna of Parishan seem
16 478 particularly sensitive to these catchment alterations by people, and due to the particular
17 479 characteristics of the two dominant species allow us to differentiate change in catchment hydrology
18 480 with a more complex model than a simple change in evaporative state.
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26 482 The $\delta^{18}\text{O}$ record also follows these changes, but superimposed on a longer trend to more positive
27 483 values through the core that matches regional patterns of lake change for the late Holocene in the
28 484 eastern Mediterranean (e.g. Roberts et al., 2008). We cannot discount some climate control on the
29 485 proxy data from Parishan. However, given the sensitivity of the lake today, and our own
30 486 observations of its desiccation, it is apparent that peoples' exploitation of groundwater and surface
31 487 waters, and management of catchment vegetation do significantly impact lake levels at Parishan.
32 488 Our record of the last 4000 years shows this is not a new problem; further investigation of the
33 489 archaeological record would now be of interest in terms of the potential impact of these proposed
34 490 human-induced hydrological changes on the populations that caused them.
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41 492 As ever, the discussion of the Parishan record highlights the need for caution in the interpretation
42 493 of any late Holocene palaeoenvironmental record from substantially inhabited parts of the world
43 494 solely in terms of climate. Human impact on the proxy record is clear. However we define and
44 495 discuss the Anthropocene (c.f. Ruddiman et al., 2015) these data provide evidence of human
45 496 impact on the local environment including the lake, an important natural resource, in direct
46 497 (agriculture) and, arguably, indirect (catchment disturbance and hydrology) ways at various points
47 498 through the last 4,000 years. Via our multi proxy approach we present one case study where
48 499 uncertainty in the link between environmental correlation and human causation is greatly reduced.
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For Peer Review

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For Peer Review

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2 667 **Figures**

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5 669 **Figure 1** Location map for Lake Parishan (P) in Iran, compared to location of lakes Urmia (U),
6 670 Zeribar (Z), Mirabad (Mi) and Maharlou (Ma). Topographic background is from the GTOPO30
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8 671 digital elevation model (Data available from the U.S. Geological Survey).
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11 673 **Figure 2** Age Depth model for core LP111. The final age model (solid line) is produced from the
12 674 radiocarbon age estimates (diamonds; 2σ ranges) corrected by a 360 year old carbon effect
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14 675 (crosses) based on the U-Series age estimate (squares). The triangle marks the pollen
15 676 stratigraphic marker for the beginning of the Achaemenid empire, used as a check on the old
16 677 carbon estimate.
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20 679 **Figure 3** Sedimentary and geochemical data from core LP111 plotted against depth. Organic C data
21 680 here are from the Costech analyses and show the same trends as the LOI data (not shown). The
22 681 location of carbonate samples analysed for mineralogy are shown (A=Aragonite; C=Calcite).
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26 683 **Figure 4** Selected pollen data for the LP111 sequence (from the full sequence from Djamali et al, in
27 684 press) grouped into pollen of trees and shrubs (Trees), herbaceous plants growing on well-drained
28 685 soils outside the wetland (Upland Herbs), aquatic and subaquatic plants growing inside and around
29 686 the wetland (Aquatics/Subaquatics), biological remains of animal and fungal origin (Non-Pollen
30 687 Palynomorphs; NPPs) and the dinoflagellates (Algae). AP/NAP: Arboreal/Non-Arboreal Pollen;
31 688 LPAZ: Local Pollen Assemblage Zones.
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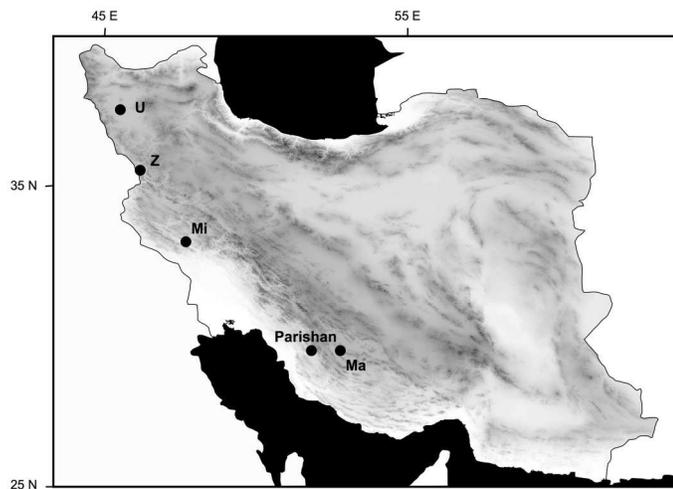
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35 690 **Figure 5** Ostracod fauna from the LP111 core plotted against depth.
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37 692 **Figure 6** Diatom flora (% count) from core LP111 plotted against depth.
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40 694 **Figure 7** Summary figure showing the key variables highlighting lake water and catchment change
41 695 in and around Lake Parishan. Periods of human impact, as defined by *L. inopinata* (Table 3), are
42 696 highlighted.
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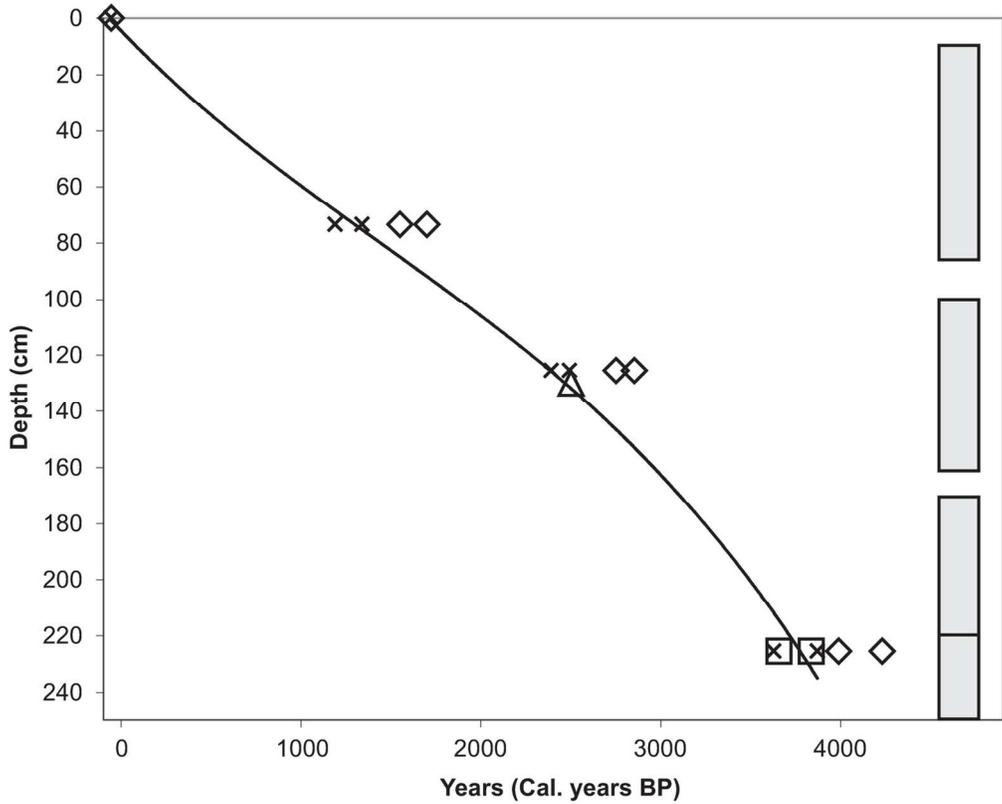
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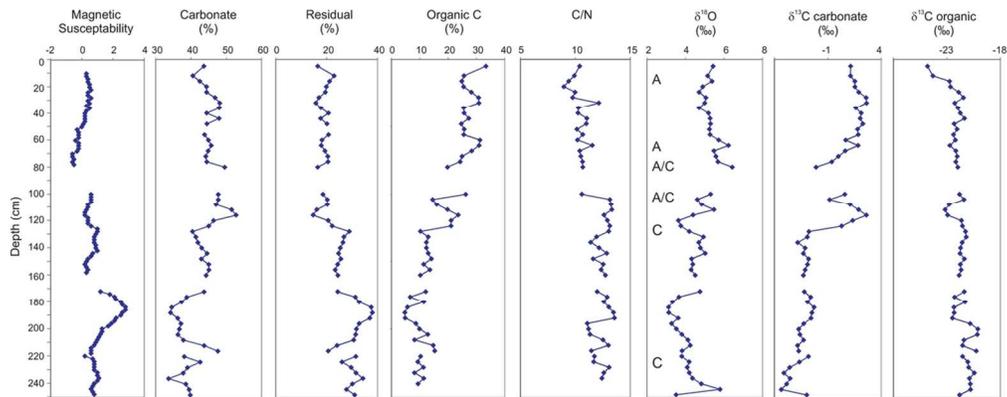
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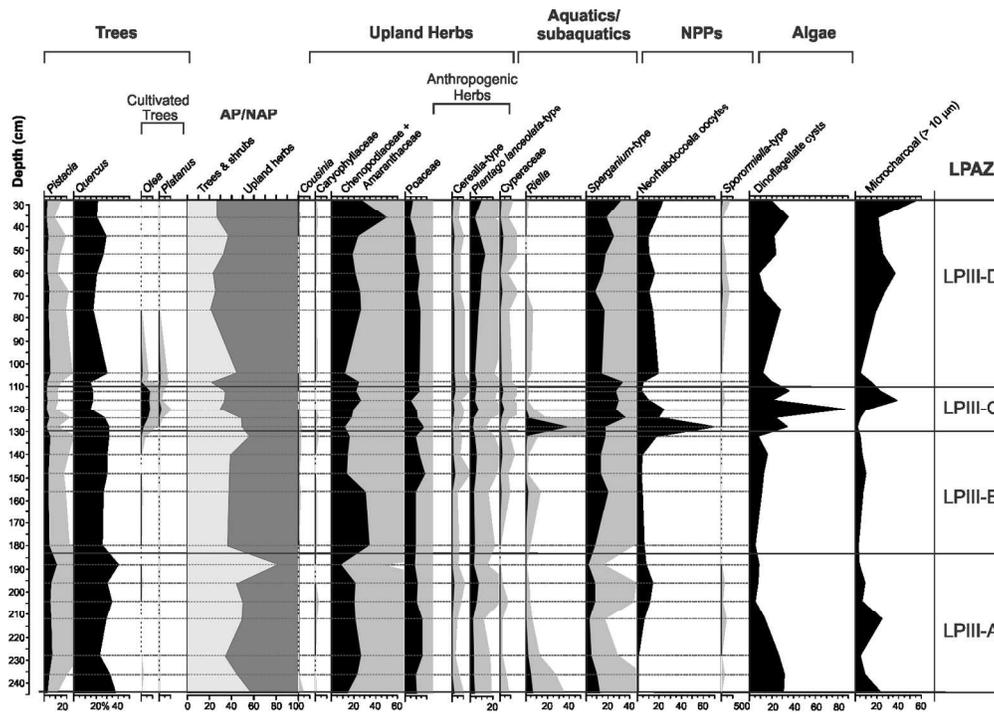


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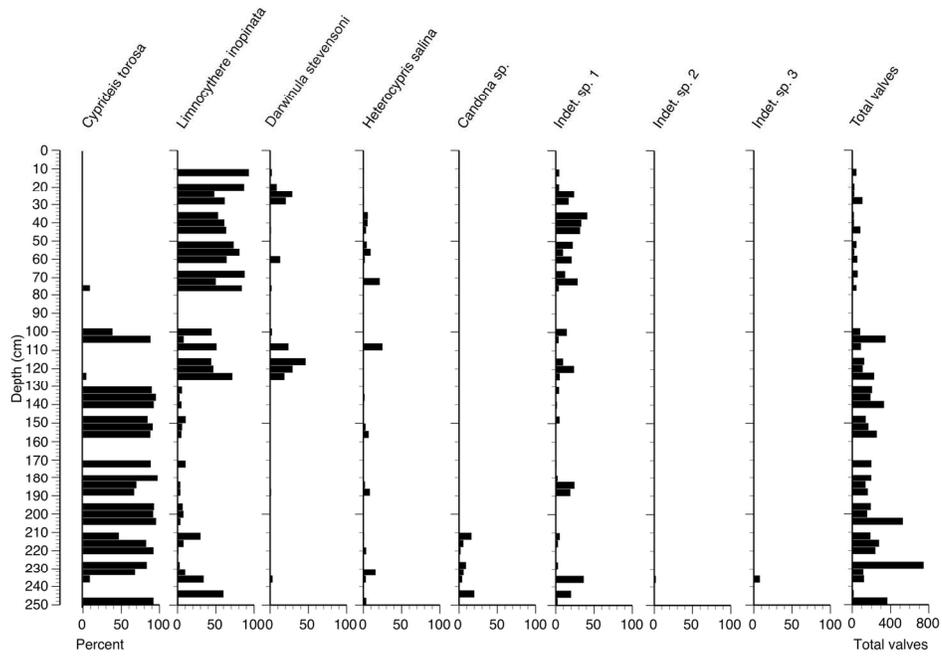
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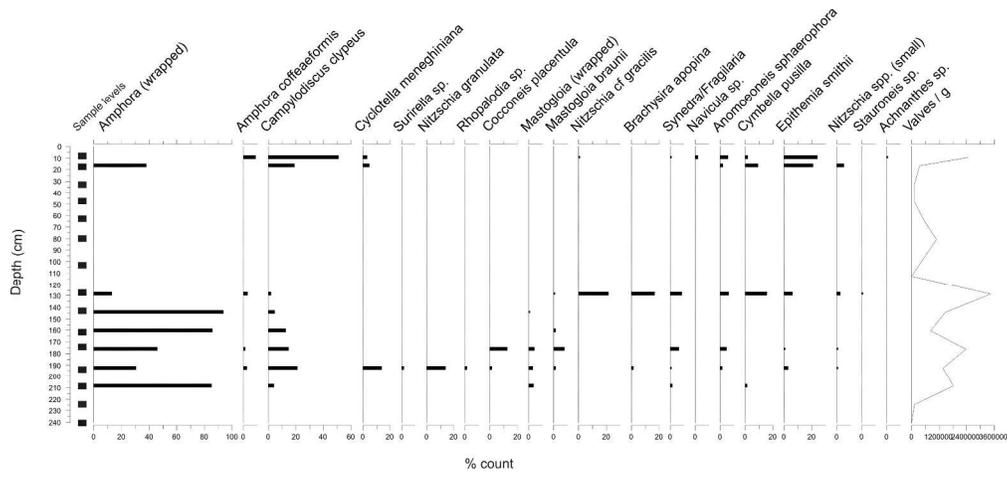
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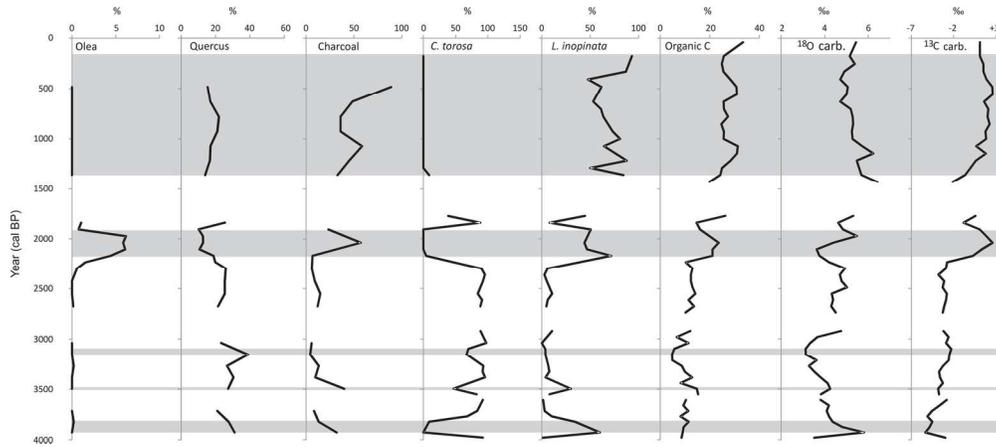
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Table 1 Radiocarbon age estimates from core LP III. Calibration undertaken using Calib 7.0 and IntCal13; 2 sigma ranges are reported.

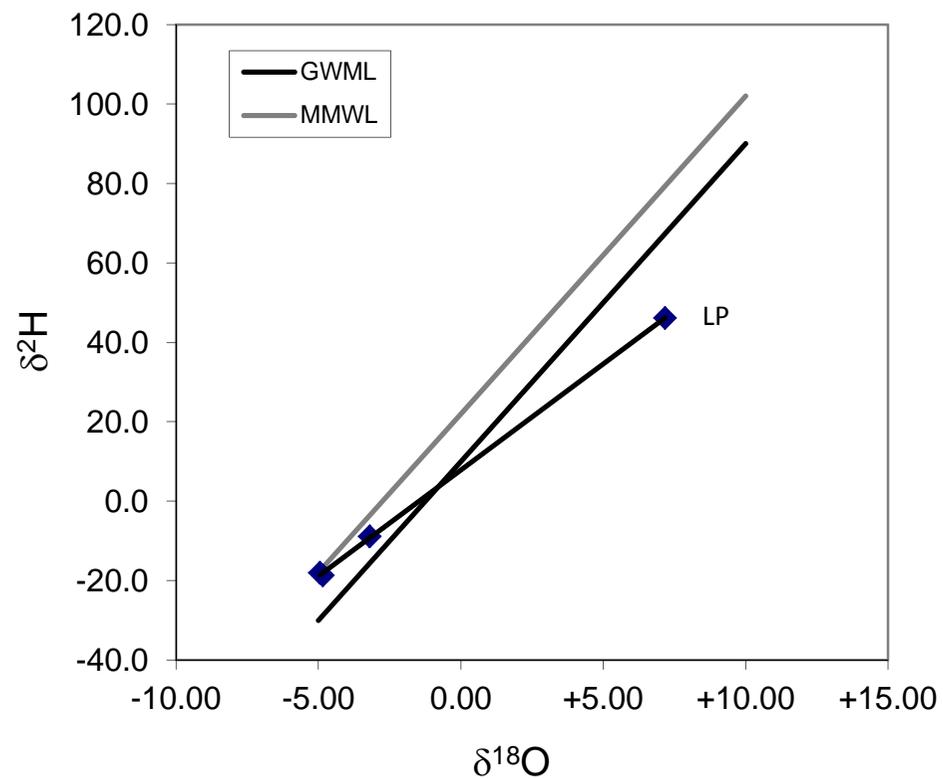
Sample name	Laboratory code	Radiocarbon age (years BP)	Calibrated age (years BP)	
LP III 73-74	Poz-21263	1705 ± 30	1549	1698
LP III 125-126	Poz-45410	2685 ± 35	2750	2851
LP III 225-226	Poz-21264	3750 ± 30	3987	4229

Table 2 U-Series age estimate from core LP III. Age Corrected using ²³²Th/²³⁸U Detrital Molecular Ratio of 3.12.

	²³⁸ U (ppm)	²³⁴ U (ppm)	²³⁰ Th (ppb)	²³² Th (ppb)	²³⁰ Th/ ²³² Th	²³⁰ Th/ ²³⁴ U	²³⁴ U/ ²³⁸ U	Age (years BP)	²³⁰ Th/ ²³⁴ U corr	²³⁴ U/ ²³⁸ U corr	Corrected Age (years BP)
	8.63390	0.00076	0.01427	1254.72	2.12425	0.06226	1.62174	6992	0.03379	1.65294	3745
Uncertainty	0.06816	0.00000	0.00012	226.81	0.08675	0.00051	0.01318	117	0.0004	0.01946	90

Table 3 The two end member environmental states for Lake Parishan through the last 4000 years based on relationships defined in Supplementary Table 1

Impacted lake state	Natural lake state
High <i>L. inopinata</i>	High <i>C. torosa</i>
Alk/Ca > 1	Alk/Ca < 1
Low oak	High oak
High charcoal	Low charcoal
High Sporormiella	Low Sporormiella
High Sparganium	Low Sparganium
High carbonate	Low carbonate
High organic carbon	Low organic carbon
Low $\delta^{13}\text{C}_{\text{org}}$	High $\delta^{13}\text{C}_{\text{org}}$
High $\delta^{18}\text{O}$	Low $\delta^{18}\text{O}$
High $\delta^{13}\text{C}_{\text{carb}}$	Low $\delta^{13}\text{C}_{\text{carb}}$
Low magnetic susceptibility	High magnetic susceptibility



Supplementary Figure Isotopic values of lake waters from Lake Parishan (LP) collected in February 2007 compared to nearby waters from the Bishapur River and local springs. The lake water describes a clear Local Evaporation Line from the Global and Mediterranean Meteoric Water Lines.

Supplementary Table Correlation analysis (r^2 and p values shown) between selected variables from core LP111. Significant relationships are shaded.

	Olea	Quercus	Charcoal	Sporormiella	Sparganium	C_torosa	L_inopinata	Carb	C_org	d13C_org	C:N	d18O_carb	d13C_carb	Mag_susc
Olea	1	-0.464	0.085	-0.179	0.554	-0.308	0.165	0.436	0.157	-0.241	0.392	-0.286	0.410	-0.134
		0.022	0.705	0.401	0.005	0.144	0.440	0.033	0.465	0.256	0.058	0.175	0.046	0.532
Quercus		1	-0.543	-0.259	-0.712	0.567	-0.570	-0.740	-0.691	0.534	0.212	-0.317	-0.735	0.637
			0.009	0.222	0.000	0.004	0.004	0.000	0.000	0.007	0.319	0.131	0.000	0.001
Charcoal			1	0.683	0.374	-0.684	0.639	0.646	0.774	-0.400	-0.624	0.579	0.698	-0.532
				0.000	0.086	0.000	0.001	0.001	0.000	0.065	0.002	0.005	0.000	0.011
Sporormiella				1	-0.053	-0.443	0.595	0.181	0.610	-0.214	-0.578	0.423	0.334	-0.404
					0.807	0.030	0.002	0.398	0.002	0.315	0.003	0.039	0.110	0.050
Sparganium					1	-0.532	0.473	0.709	0.529	-0.356	-0.098	0.230	0.709	-0.459
						0.007	0.020	0.000	0.008	0.088	0.649	0.280	0.000	0.024
C_torosa						1	-0.916	-0.488	-0.704	0.479	0.351	-0.536	-0.659	0.593
							0.000	0.016	0.000	0.018	0.092	0.007	0.000	0.002
L_inopinata							1	0.497	0.792	-0.440	-0.519	0.656	0.637	-0.711
								0.014	0.000	0.031	0.009	0.000	0.001	0.000
Carb								1	0.704	-0.586	-0.193	0.436	0.787	-0.697
									0.000	0.003	0.366	0.033	0.000	0.000
C_org									1	-0.480	-0.662	0.621	0.844	-0.747
										0.018	0.000	0.001	0.000	0.000
d13C_org										1	0.036	-0.337	-0.644	0.415
											0.869	0.107	0.001	0.044
C:N											1	-0.630	-0.363	0.466
												0.001	0.081	0.022
d18O_carb												1	0.334	-0.793
													0.111	0.000
d13C_carb													1	-0.512
														0.011

Light grey shading: Correlation is significant at the 0.05 level.

Dark grey shading: Correlation is significant at the 0.01 level.