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1	At the edge of the desert: The evolution of the Holocene environment in Lake Rkiz, Senegal Biver Valley
2	River valley
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12	
13	
14	Abstract
15	The analysis of faunal remains, pollen grains and other palynomorphs from a 4 m long core
16	recovered at Lake Rkiz, Mauritania (16.911420 N, -15.172668 W, 0 m asl) documents the
17	major phases of marine intrusion in the Senegal River system during the Holocene. Lake Rkiz
18	was exposed to marine waters from 10 to 4 ka: a brackish lagoon developed at 10 ka,
19	followed by a mangrove between 8.7 and 7 ka both in a calm environment. Rhizophora
20	populations were never developed, however, either because they did not reach the core site at
21	the northern extremity of the lake, or because they were restricted to the Senegal River bed.
22	After 7 ka, the environment became increasingly unstable and the mangrove was replaced by
23	a more clearly marine environment. During the last two millennia, a new, discrete phase of
24	marine intrusion is recorded at Lake Rkiz. Contemporary to that already noted in the Ferlo
25	River valley which feeds the Senegal River through Lake Guiers to the south, it also testifies
26	to a positive oscillation of the sea level. The evolution of the regional vegetation of Lake Rkiz
27	follows that of North tropical Africa with the expansion of a gallery forest of tropical humid
28	affinity dominated by <i>Celtis</i> during the AHP. As elsewhere in the Sahara and Sahel, tropical
29	plant types co-occurred with Saharan ones attesting to a mosaic-like landscape.
30	
31	Keywords: Senegal River; Holocene mangrove; Marine intrusion upstream; Sea level;
32	Pollen: Microfauna & mollusks

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35 **1.** Introduction

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37 The recent paleoenvironemental evolution of the Senegal River delta has been impacted by a 38 combination of two distinct factors: The massive increase in humidity during the "African 39 Humid Period" (AHP) that led to the wide dispersion of tropical plants across the Sahel and the Sahara (Watrin et al., 2009; Hély et al., 2014) and the post-glacial sea level rise from 40 41 about -130 m during the Last Glacial Maximum (LGM) to the present level about 7000 years 42 ago (Fleming et al., 1998) responsible for the penetration of the sea in low-lying coastal areas of western Africa (Faure et al., 1980). Tropical trees (e.g., Alchornea, Macaranga, Elaeis 43 44 guineensis, Tetracera, Celtis, Ficus, Syzygium) formed dense gallery forests as soon as 10.7 ka in the interdunal fens of the Niayes area of Senegal located behind the Atlantic coastal 45 strand between 15° and 16°N. Humid tropical trees were also present in the catchment area of 46 Lake Guiers (Lézine, 1988) and in the Senegal River bed (Michel and Assémien, 1969; 47 Médus, 1984). At the end of the AHP between 3000 and 2500 cal yr BP (labelled as 3 and 2.5 48 49 ka in the text), the degradation of environmental conditions has led to the destruction of these 50 tropical gallery forests (Lézine, 2022). They still persist locally as relict vegetation formations

in the Niayes owing to the presence of a water table near the surface (Trochain, 1940), but
they disappeared from the banks of the Senegal River and its tributaries during the last few
centuries (Fofana et al., 2020; Lézine et al., 2023).

54 The increase in humidity during the AHP was also responsible for the massive development of tropical humid Rhizophora mangroves on the West African coast during phases of 55 56 stabilization of the post-glacial sea level rise (Lézine, 1997). The morphology of the coastal 57 areas played a major role in the wide expansion of these mangroves, particularly in the Senegal River area (Faure et al., 1980). In its lower valley, from the coast to approximately 58 59 225 km to the east, the bottom of the Senegal River bed is about 5 m below the present-day 60 sea level (Michel, 1973). This geomorphological feature facilitated the penetration of marine waters eastward several hundred kilometers from the present coast and the development of 61 62 mangroves during periods of sea level rise in the Holocene (Trochain, 1940; Michel and Assémien, 1969; Médus, 1975; Médus and Monteillet, 1981; Lézine et al., 1988). However, 63 64 our understanding of the Holocene evolution of the West African mangrove is severely 65 hampered by the lack of continuous records. Mangrove levels (most of the time in the form of a peat layer a few centimeters thick) (in Lézine, 1997 and references therein) and shell 66 accumulations forming marine terraces in the interior were used to reconstruct the coastline 67 68 and its variations over the last millennia (e.g., Faure et al., 1980). The irregular and often 69 truncated nature of the sedimentary archives does not permit a continuous reconstruction of the littoral evolution throughout the Holocene. Possible shore level tilt along the Senegal 70 71 River delta during the post glacial sea level rise may have introduced an additional difficulty 72 in our understanding of paleoenvironmental changes in this area (Faure et al., 1980).

In this paper we present a micro-faunal and palynological record from a continuous sedimentary sequence from lake Rkiz (16.911420 N, -15.172668 W, 0 m asl) in order to identify the timing and amplitude of the successive phases of marine intrusion upstream in the Senegal River system from 10 ka to the present.

78 2. Environmental setting

80 Lake Rkiz occupies a large interdunal depression (Aftout ech Cherguil) north of the Senegal 81 River, about 110 km east of the present coastline (Fig. 1). The NE-SW orientation of the depression is governed by that of the sand dune alignments established during the LGM over 82 83 much of southern Mauritania and northern Senegal (Michel, 1973).). Lake Rkiz is about 5 km 84 wide and 30 km long. It is a shallow lake (maximum depth 2.5 m), hydrologically connected 85 to the Senegal River through a network of multiple channels. The hydrological regime of the lake reflects that of the river in its lower valley. During the rainy season (from June to 86 87 October) Lake Rkiz is supplied by the floodwaters of the Senegal River and its level can rise 88 by 3.5 m. During the dry season, when the inflow from the Senegal River ceases, the lake may 89 dry up completely.

The regional climate is characterized by ca. 185 mm annual rainfall during the rainy season from June to October (maximum 79 mm in August) and maximum temperatures between 33 and 40°C. As a consequence, the vegetation is of Sahelo-Saharan steppic type with a discontinuous herbaceous cover mainly composed of *Cenchrus biflorus*. Trees (*Acacia raddiana, A. scorpioides, Salvadora persica, Balanites aegyptiaca...*) are scarce (Adam, 1960; 1965).

97 **3. Material and Methods**

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99 Core Rkiz-1 was collected in 1985 at the mouth of Wadi Mrea at the NE end of the lake 100 (16.911420 N, -15.172668 W, 0 m asl) using a Russian manual corer (Fig. 1). The

101 sedimentary sequence is 4m long (Fig. 2). Three main units can be distinguished: from the 102 base to 349 cm (Unit I), the sediment consists of a homogeneous light-yellow sand likely 103 derived from to the dune sands deposited during the LGM. It is overlaid by a greenish clay 104 layer containing abundant malacological, crustacean and fish bones fragments, witnesses to 105 the Holocene marine transgression (Unit II: 349-190 cm). The upper unit (Unit III: 190-0 cm) 106 is composed of sandy clay with fine sand and gravel intercalations. This testifies for a less 107 homogeneous environment with runoff deposits from the lake edges (such as the ferruginous gravels, Michel, 1973), lake and mangrove organic material and eolian sands. Thirty-six 108 109 samples were taken at 10 cm intervals for faunal and vegetation analysis.

110

111 **3.1 Dating control**

112

The core chronology was constructed using ¹⁴C accelerator mass spectrometry (AMS) dates 113 114 on seven samples of marine shells (Pachymelania fusca) and bulk sediment (Fig. 2). 115 Radiocarbon dates on shells and shell fragments were calibrated using the Marine 20 116 calibration curve (Heaton et al., 2000) from Calib 8.2 software (Stuiver and Reimer, 1993) taking into consideration the reservoir-age of 511 ± 176 years, calculated for the sector of the 117 118 West African coast closest to the site, facing the Senegal River delta (Ndeye, 2008). The bulk organic sample at 95 cm was calibrated using the IntCal 20 calibration curve (Reimer et al., 119 120 2020). Dates are given in calibrated 1000 years (ka) BP. The age-depth model was then derived from a linear interpolation between two calibrated ¹⁴C ages using CLAM (Blaauw, 121 122 2010). We assume a constant reservoir-age throughout the Holocene time.

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124 **3.2 Faunal remains**

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The faunal samples were sieved under water on a 0.5 mm mesh and then sorted under a binocular microscope. Only 18 samples between 370 and 190 cm depth yielded identifiable remains and the uppermost samples were found to be sterile (Fig. 3).

Anatomical and taxonomic identifications were conducted using a comparative reference
collection from AASPE (library of skeletons and tests), literature (Dautzenberg 1910, 1912,
Gregory 1933, Bouchet 1977, Pointier & Lamy 1998, Humann & DeLoach 1993, Grouard
2001, Serrand 2002, Ardovini & Cossignani 2004, Ahmedou Salem et al. 2014, Carpenter &

- 133 De Angelis 2016, Diedhiou et al. 2018, von Cosel & Gofas 2019, Taylor & Glover 2021), and 134 internet resources (MolluscaBase 2022, World Register of Marine Species 2022). The 135 nomenaleture uses undeted from the WoRMS site
- 135 nomenclature was updated from the WoRMS site.

136 Quantification of faunal remains used multiple measures often used in zooarchaeological 137 analyses (NRd: Number of Determined Remains and MNI: Minimal Number of Individuals). 138 Bivalves were classified into three groups (intact valve, broken valve but with umbo, and fragments (Claassen, 1998; Giovas, 2009; Manca, 2017)). The Bivalve MNI was estimated 139 140 from the lateralized complete valves, lateralized hinges and hooks, edge and complementary 141 non-repetitive elements which allow, according to the cases, the maximum estimate. The 142 Gastropod fragmentation degrees were classified in several groups (complete, sub-complete, 143 complete without apex, complete without opening, margin and lip, lip, apex, apex and spire, 144 first spire, fragment of spire, parietal margin, spine, fragment of columella, fragment of opening or base, umbilicus, fragment of base with umbilicus, columella and opening, 145 operculum. The Gastropod MNI was calculated based on complete shells, number of apexes 146 147 and complementary non-repetitive elements which allows, on a case-by-case basis, the 148 maximum estimate of complete labials (Serrand, 2002; Giovas et al., 2013; Tomadini, 2018).

149 In total twenty-eight taxa of Bivalvia, Gastropoda, Cnidaria, Crustacea and Teleostei were 150 identified out of 2610 remains. They were grouped according to their ecological affinity in 4 groups (marine, estuarine, mangrove, freshwater) according to World Register of MarineSpecies 2022 (S-table 1; S-fig. 1).

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154 **3.3 Pollen and NPP**

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156 Pollen samples were processed using the standard HF method described by Faegri and Iversen 157 (1977). After sieving in a 5 µm mesh, the residue was stained with safranin and mounted in glycerine. Pollen identification was based on the reference slide and photo collections of the 158 159 African Pollen Database (APD) (https://africanpollendatabase.ipsl.fr). Pollen nomenclature, 160 which follows the standards defined by the APD (Vincens et al., 2007), was based on Lebrun 161 and Stork (1991-2015). The pollen microflora was composed of 136 taxa (S-table 2). Pollen 162 taxa were grouped according to the phytogeographical and ecological affinity of the parent species according to Trochain (1940) and Lebrun (1973), among others, with distinction of 163 164 Saharan, Sahelian, Sudanian and Guineo-Congolian pollen types, mangrove, wet indicators 165 and undifferentiated or common to all the phytogeographical entities mentioned above 166 (including Poaceae). Pollen percentages were calculated against a sum excluding mangrove 167 pollen types, wet pollen types (including Cyperaceae) and ferns, the percentages of which 168 were calculated separately against a sum including all the pollen grains and fern spores 169 counted. Non-Pollen Palynomorphs (NPP) include algae (mainly Pediastrum and 170 Botryococcus), undifferentiated Foraminifera, Dinocysts and Fungi. Their percentages were 171 calculated against the total NPP.

172

The Shannon Diversity index (Pielou, 1966) which accounts for both abundance and evenness
of the taxa present was used to evaluate the diversity of pollen and faunal remains using Past
4.03 software (https://www.nhm.uio.no). Both pollen and faunal data are presented under the
form of percentage diagrams using Tilia software (Grimm, 1991).

177178 4. Results

179180 **4.1 Faunal remains**

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182 **4.1.1 Diversity**

The MNI calculated is 1312. Gastropoda largely dominates this collection with an MNI of 684 calculated from 886 identified remains (NRd), immediately followed by Crustacea (MNI= 426; NRd=848), and Bivalvia (MNI=185, NRd762). Cnidaria and Teleostei are poorly represented with MNI= 13 (NRd=122) and MNI=4 (NRd=4), respectively (Fig. 3).

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Maximum of Gastropoda and Crustacea is present at the 8 ka level. These two groups are then replaced by Bivalvia which reach peaks at two distinct levels respectively dated at 7.3 then 6.3 ka. Bivalvia are the most diverse with 12 species and 9 families present, largely accounting for the maximum Shannon diversity index reached at 7 ka. They are followed by Gastropoda with 7 species and 7 families present. On the contrary, only one species of Teleostei, 1 Cnidaria undiff., and 2 Crustacea (Cirripedia and Ostracoda) were present.

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195 **4.1.2 The mollusk assemblages**

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197 Figure 4 shows the diversity and MNI of the main mollusk taxa, the preservation of which

allows distinguishing two main zones. From 10 to 7 ka (zone 1), the preservation rate is high (mean=76%), though fluctuating from 100% to 32%. It abruptly falls to values as low as 3%

200 in the overlying zone 2 (mean=13%).

- 201202 Three sub-zones can be distinguished within zone 1:
- Zone 1a: From 10 to 8.7 ka, the presence of *Peringia ulvae* and the highest values of
 the preservation rate indicate indicate a sheltered lagoon, out of the direct influence of
 waves;
- Zone 1b: From 8.7 to 7.5 ka: *Peringia ulvae* was associated with *Pachymelania fusca*,
 both largely dominating the mollusk assemblages, suggesting the progressive
 development of a mangrove at the coring site. Freshwater inputs are indicated by the
 presence of Ampullariidae (*Lanistes ovum*). The lower values of the preservation rate
 compared to zone 1a (75-50% versus 100%) suggest more dynamic environment;
- Zone 1c: From 7.5 to 7 ka: *Peringia ulvae* and *Pachymelania fusca* strongly decreased and the mollusk assemblage greatly diversified with the appearance of marine and estuarine species such as various Veneridae and *Ungulina rubra*. The high amplitude of the variations in the preservation rate suggests the alternation of phases of calm seas and strong swells.
- 216 Zone 2, from 7 to 5.6 ka, is characterized by the disappearance of the mangrove species and 217 their replacement by marine ones, as shown by the high MNI of Veneridae and Lucinidae 218 species. The lowest values of the preservation rate for this zone indicate significant tidal and 219 backwash movements.

4.2 Pollen and NPP

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The pollen diagram (Fig. 5) is characterized by the dominance of the herbaceous taxa Poaceae undiff. and various types of Amaranthaceae whose cumulative percentages reach up to 90% of the pollen sum, indicating the presence of an open environment throughout the Holocene. As usual in tropical Africa, the pollen percentages of trees are rarely higher than 1% with the exception of Combretaceae, *Celtis* and *Rhizophora* which reach up to 9%, 23% and 7% respectively.

229

230 Three main zones can be distinguished from the pollen grain, spores and NPP diagram:

231

Zone 1 (10-8.7 ka) is typified by exceptionally high percentages of *Celtis* (maximum: 23%).
Among the herbaceous plants, taxa from the lake margin such as *Typha* and Cyperaceae, are
present. *Botryococcus* dominates the NPPs suggesting shallow and nutrient poor fresh to
brackish water. The presence of *Ephedra* pollen grains, even in percentages lower than 1%
suggest a Saharan (desert) influence. (This pollen zone corresponds to Zone 1a of the mollusk
assemblages).

238

239 Zone 2 (8.7-4.5 ka) records the decline of *Celtis*. The other tropical tree taxa, notably Alchornea and various types of Combretaceae remain present though in small percentages. 240 241 Among the herbaceous plants, the abundance of Amaranthaceae and in particular 242 Amaranthus/Chenopodium-type and Suaeda allows identification of sub-zone 2a (8.7-7.1 ka) 243 (= Zones 1b and 1c of the Mollusk assemblages). The presence of Suaeda suggests saline 244 conditions while the presence of Fungi confirms the, partial or seasonal, desiccation of Lake 245 Rkiz at the core site. The replacement of *Botryococcus* by dinoflagellates within the NPP 246 indicates that a marine influence was present, which is confirmed by the occurrence of 247 Rhizophora mangrove pollen grains. Pollen types present in subzone 2a continue into the overlying subzone 2b (7.1-4.5 ka) (= Zone 2 of the Mollusk assemblages) though in smaller 248 249 percentages, suggesting drier conditions. The NPPs suggest an unstable environment with alternating saline (Dinoflagellate cysts in particular at 6.2 ka) and freshwater to brackish
 (*Botryococcus*) conditions.

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253 Zone 3 (4.5 ka-modern) records a change in the environment with the disappearance of 254 mangrove taxa. In sub-zone 3a (4.5-2.4 ka) Celtis and Alchornea disappear but the presence 255 of e.g., Lannea suggests that the tropical influence was still present, although significantly 256 reduced. After 2.4 ka (sub-zone 3b (2.4-1.6ka)) the Sahelian taxa Acacia and Commiphora 257 which are associated with the Fabaceae family increase, indicating the transition to the 258 modern environment. Rare tropical plants of Sudanian or more humid types (e.g., Lannea, 259 Macaranga, Bridelia...) are occassionally present but in percentages lower than 1%, and the 260 reappearance of the Saharan taxa (Ephedra) indicates the return to dry conditions. The 261 lacustrine environment is dominated by Botryococcus. The populations of the lake edge develop progressively with Typha in sub-area 3b then Cyperaceae in sub-zone 3c. The 262 263 predominance of fresh-water during sub-zone 3c (1.6-0.83 ka) and 3d (0.83-modern) 264 indicated by the dominance of *Pediatrum* in the NPP assemblages is interrupted around 1.3 ka by the discrete reappearance of mangrove taxa and dinoflagellates, testifying for the brief 265 266 return of marine influence.

268 **5.** Discussion

270 **5.1** The Holocene sea level variations and their impacts in the Lake Rkiz area

5.1.1 The *Rhizophora* mangrove273

274 The Senegal River delta is the northernmost limit of the major West African mangrove 275 species, Rhizophora racemosa (Trochain, 1940; Adam, 1965; Spalding et al., 1997). Today 276 this tropical humid species is present only as small populations or isolated individuals along 277 the creeks closest to the coast in the Senegal River delta near Saint Louis (Senegal) or on the 278 Thiong island (Mauritania) at roughly 16°N. As the sea level rose after the LGM, and thanks 279 to the intensification of the monsoon rains, it spread widely along the West African coasts 280 during the AHP (Lézine, 1997) as far north as 21.25°N (core GIK16017-2, Hooghiemstra 281 1988). In the Senegal River area, *Rhizophora* was present on the continental shelf opposite the 282 present-day delta at 13.8 ka (core Rosilda N 110-Z ; Lézine, 1987). It penetrated eastward in 283 the rear delta after 11.4 ka (Médus and Monteillet, 1981) and in the Potou interdunal 284 depression at the southern limit of the delta (Lézine, 1987). In this context, and given the 285 topography of the Senegal River bed, the intrusion of the coastal and marine influence up to 286 Lake Rkiz from 10 to 4.7 ka is not unlikely. Between 10 and 8.7 ka, the low faunal diversity, 287 the only occasional presence of dinoflagellates and the very low percentages of *Rhizophora* 288 pollen, show that the mangrove was only slightly developed and that *Rhizophora* populations 289 did not reach the core site, i.e., the most distant sector from the channels connecting Lake 290 Rkiz to the Senegal River. Excellent faunal preservation eliminates the potential long-distance 291 transport and suggests the presence of a lagoon protected from the direct influence of swells 292 and strong water currents. The time interval between 8.7 and 4.7 ka during which the 293 mangrove developed at Rkiz encompasses the transgressive episode of the "Nouakchottien" 294 (Elouard, 1968). During this episode centered around 6-5 ka, the sea level was higher by 1 to 295 2 m compared to its current position (Faure et al., 1980). Sea water intrusion inland, in the 296 rear delta, and along the Senegal River and its tributaries, is attested by numerous accumulations of marine shells forming terraces (e.g., Faure et al., 1980 and references 297 298 therein) (Fig. 6 and 7A; S-table 3), palaeobotanical (Rhizophora pneumatophores) and 299 botanical (relict species of mangrove forests) observations (Trochain, 1940), and pollen

300 analyses (Médus, 1975; Médus and Monteillet, 1981; Lézine, 1988). According to Michel and 301 Assémien (1969), Rhizophora populations could have spread up to Bogué, 250 km east of the present coast in the bed of the Senegal River during this episode (Fig. 6). At Rkiz, the 302 303 mangrove was progressively exposed to increased marine influence after 7 ka and a more 304 dynamic environment subject to waves, marine and freshwater inputs was established as 305 shown by the important presence of Bivalvia, Veneridae (notably Pelecyora isocardia) and 306 Lucinidae in the fauna, the strong fragmentation of faunal remains and the alternation of 307 dinoflagellates and freshwater algae within the NPP assemblage. Increased currents and wave 308 turbulence led to the complete destruction of the faunal remains from the sediment at 5.6 ka. 309 However, *Rhizophora* pollen grains show that the mangrove influence persisted up to 4.2 ka. 310 These results agree with previous studies showing that the decline of *Rhizophora* mangroves 311 along the northern coast of Senegal occurred, according to the sites, between 5.6 ka at 14.9°N (Thiave, Lézine et al., 1985) and 3.6 ka off the Senegal River delta at 16.0°N (Core GeoB 312 313 9503-5, Bouhimetarhan et al., 2009). Evidence of marine influence disappeared from the 314 pollen and NPP assemblages between 4.2 and 2.1 ka that closely corresponds to the 315 "Tafolian" sea level regression (Hébrard, 1972). Then, the renewed presence of a few 316 *Rhizophora* pollen grains between 2.1 and 0.6 ka shows that the last, discrete episode of saline 317 intrusion into the Senegal and Ferlo Rivers during the last millennium (Faure et al., 1980; Monteillet et al., 1981; Fofana et al., 2020) also reached Lake Rkiz. The absence of faunal 318 319 remains in the sediment and the very low Rhizophora pollen percentages confirm that this 320 event was minor.

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322 **5.1.2** Vertical movements in the Rkiz Lake area during the Holocene?

323 324 The elevation of mangrove levels and shell accumulations have been used to reconstruct the 325 long-term evolution of sea level rise off Senegal: The peat layer in the Rosilda N 110 Z core 326 located at -60 m below sea level is dated 13.8 ka (Lézine, 1987). In the rear delta of the 327 Senegal River Médus and Monteillet (1981) identified a Rhizophora mangrove level at -33.6 328 m dated at 11.4 ka in line with the dating of the Eudoxie and Lea long cores by Faure et al. 329 (1980) (9.9 ka and 9.4 ka at -23 and -20.5 m, respectively). All these dates are consistent with 330 the general trend of the post-glacial sea level rise (Fleming et al., 1998). The most surprising 331 result of our study is the depth at which the early Holocene mangrove development has been 332 identified at Rkiz. There is a difference of about 20 m with the depth of the contemporary 333 levels identified in the rear delta of the Senegal River (Fig. 7B). Early Holocene vertical 334 movements are therefore suggested by our data in the sector of Lake Rkiz located in a region known to have also been affected by significant deformations during the Quaternary, such as 335 336 the subsidence of the Senegal River delta, and the faulting of the "Dôme de Guiers" (Michel, 337 1973; Barusseau et al., 2009).

338

339 5.2 The regional vegetation340

341 During the AHP, lakes and wetlands expanded in the present day Sahara and Sahel 342 (Lézine et al., 2011) while large paleodrainages were formed or re-activated (Drake et al., 343 2011), in response to the increase in monsoon rainfall. As a result, tropical plants widely spread into the Sahel and the Sahara (Watrin et al., 2009). Due to the scarcity of data in these 344 345 (semi-) arid environments where lacustrine archives are poorly preserved, the major difficulty 346 in understanding this profound environmental change was to identify how the tropical humid plants could have migrated north and where they developed. Instead of a general displacement 347 348 of the vegetation zones towards the north and therefore the development of a more or less 349 wooded savannah in the Sahara and the Sahel (Hoelzmann et al., 1998), pollen data show the 350 presence of gallery forests of tropical humid type (Sudanian to Guineo-Congolian) in a drier 351 regional environment. It has been suggested that tropical plants have used the banks of the 352 rivers where they benefited from the moisture of the soil as migration routes to enter the Sahel 353 and Sahara (Watrin et al., 2009; Hély et al., 2014). At Rkiz the regional landscape has always 354 been extremely open as shown by the percentages of herbaceous taxa, which may be equal to 355 or higher than 90% in total. Tropical plants (Guinean-Congolian, Sudanian) have co-existed 356 with typically Sahelian species (e.g., Acacia, Commiphora) and even Saharan species 357 (Ephedra, Artemisia). This mixture of plants, which today can only be found in distinct 358 vegetation zones, explains the peak in diversity observed between 8.8 and 8.3 ka, which is 359 consistent with the Holocene forest optimum in tropical North Africa (Lézine, 1989) (Fig. 8). 360 Tropical influence at Lake Rkiz persisted until 2.4 ka which is consistent with what we know 361 from the end of the AHP in Western Africa (Lézine, 2021). In the nearby Niaye region along the Atlantic coast between Dakar and St Louis for instance, the dense gallery forests, which 362 363 widely developed during the AHP dramatically collapsed around 2.5 ka. At Rkiz, the decrease 364 of tropical taxa was gradual and an arid phase emerged as early as 4 ka. The discrete reappearance of tropical taxa after 2 ka suggests the forest galleries that extended along the 365 366 Senegal River during the Holocene had not completely disappeared over the last millennia. 367 They were still present in the Saint Louis area as late as 0.35 ka (Fofana et al., 2020).

368

369 Thirty-six pollen taxa of humid tropical type were found at Rkiz. This number is surprisingly 370 similar to that observed on the opposite bank of the Senegal River, about 95 km to the south at Guiers 3 (16.28°N) (Lézine, 1988). However, the percentages calculated over the entire 371 372 period covered by the pollen series are on average two and a half times lower at Rkiz (6%) than at Guiers (15.5%). Comparing all pollen sites in Senegal and Mauritania, humid tropical 373 374 taxa are 68 at Diogo (15.2°N) where they account for an average of 25% of the pollen sum. 375 These values drop to 37 taxa and 15.5% at Guiers 3, 36 taxa and 6% at Rkiz and finally 15 376 taxa and 1.5% at Chemchane (20.93°N). The contrast between the percentages of tropical taxa 377 between Guiers and Rkiz only less than a hundred kilometers apart suggests that the Senegal 378 River may have constituted a barrier to the northward migration of tropical plants during the 379 AHP in north west Africa. One exception is *Celtis*, which has reached exceptionally high 380 percentages at Rkiz (23%) and so far unreported in tropical North Africa during the Holocene. 381 The modern distribution of Celtis toka does not exceed 16°N (Fig. 9). During the Holocene, it 382 expanded widely into Mauritania (Monod, 1983) and more widely across the whole Sahara to southern Libya at 24.8°N (Mercuri, 2008). However, Celtis percentages never exceeded 4% at 383 384 any of the Saharan and Sahelian pollen sites, suggesting that, with the exception of Rkiz, this 385 species never expanded much within the gallery forests. 386

387 6. Conclusion

388 389 Our study shows that, despite a substantial distance of more than 180 km from the mouth of 390 the Senegal River, Lake Rkiz was subject to marine water influx during two distinct episodes 391 of sea level rise. The first one, from 10 to 4.5 ka, is a long period during which the 392 environment of Lake Rkiz changed from a calm coastal environment evolving from a lagoon 393 to a mangrove to a more pronounced, unstable, marine environment. Two major features 394 emerge from our study: (1) the elevation of the marine influence reported in core Rkiz-1, is 395 significantly higher than what is known from the global sea level curve, which suggests 396 important vertical movements at Lake Rkiz (2) the "Nouakchottian" transgression, that led to 397 the development of mangroves in Lake Guiers and in the Senegal River bed up to Bogué 398 during the Middle Holocene, resulted in a highly variable environment at Rkiz as evidenced 399 by the increased fragmentation in the faunal remains. A second discrete phase of marine

intrusion is detected during the last 2 millennia, after the regressive phase of the "Tafolian".
Regional vegetation at Rkiz was open, dominated by herbaceous plants throughout the
Holocene. Tropical trees however formed gallery forests along the river and lake beds. These
results are consistent with the general pattern observed in the landscape evolution of West
Africa during the Holocene.

405

406 Author contributions

407 AML and SG designed research; AML performed fieldwork; AML and SG performed
408 research: AML palynology and SG zooarchaeology; AML and SG analyzed data; AML and
409 SG wrote the paper with contributions from both authors.

410

411 Data availability

412 data Pollen deposited in the African Pollen Database (APD) are 413 (https://africanpollendatabase.ipsl.fr: DOI:https://doi.org/10.14768/apd.0bba-rk05). Data for 414 fossil Celtis distribution are from the APD (https://africanpollendatabase.ipsl.fr), and for the 415 modern one from GBIF (https://www.gbif.org) and MNHN (https://www.mnhn.fr) databases 416 and the literature. Microfauna data are presented in the Supplementary Material.

417

418 **Declaration of competing interest**

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

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434 References435

- 436 Adam, J.G., 1960. Contribution à l'étude de la flore et des groupements végétaux de l'Afrique
 437 occidentale. La Mauritanie sahélienne occidentale. Inventaire de l'herbier Sadio Kane de
 438 la région du lac Rkiz (Trarza). *Notes Africaines* 86, 44-8.
- 439 Adam, J.G., 1965. La végétation du delta du Sénégal en Mauritanie. Bulletin de l'Institut
 440 Fondamental d'Afrique Noire, A 27 (1), 121–138.
- 441 African Pollen Database (APD), https://africanpollendatabase.ipsl.fr.
- Ahmedou Salem, M.V., van der Geest, M., Piersma, T., Saud, Y., van Gils, J.A., 2014.
 Seasonal changes in Mollusc abundance in a tropical intertidal ecosystem, Banc
 d'Arguin (Mauritania): testing the 'depletion by shorebirds' hypothesis. *Estuarine*, *Coastal and Shelf Science* 136, 26e34. DOI: 10.1016/j.ecss.2013.11.009
- Ardovini, R., Cossignani, T. 2004. West African seashells (including Azores, Madeira and Canary Is.). L'Informatore Piceno, Ancona Italy. 319 p.
- Bouchet, P., 1977. Distribution des Mollusques dans les mangroves du Sénégal. *Malacologia*16 (1), 67-74.

- 450 Bouhimetarhan, I., Dupont, L., Schefuß, E., Mellenhauer, G., Mulitza, S., Zonneveld, K., 451 2009. Palynological evidence for climatic and oceanic variability off NW Africa during 452 the late Holocene. Quaternary Research 72 (2),188-197. DOI: 453 10.1016/j.yqres.2009.05.003
- 454 Carpenter, K.E., De Angelis, N., 2016. FAO species identification guide for fishery purposes
 455 the living marine resources of the Eastern Central Atlantic. Vol. 2. Bivalves,
 456 gastropods, hagfishes, sharks, batoid fishes, and chimaeras. Food and Agriculture
 457 Organization of the United Nations, Roma, pp. 665-1509.
- 458 Claassen, C., 1998. Shells. Cambridge University Press, Cambridge, 266 p.
- Cosel, R. von, Gofas, S., 2019. Marine bivalves of tropical West Africa: from Rio de Oro to
 southern Angola. *Faune et Flore Tropicales* 48, 1-1104. DOI: 10.1111/maec.12622
- 461 Dautzenberg, P., 1910. Contribution à la faune malacologique de l'Afrique occidentale. Actes
 462 *de la Société Linnéenne de Bordeaux* 64, 47-228.
- 463 Dautzenberg, P., 1912. Mission Gruvel sur la côte occidentale d'Afrique (1909-1910).
 464 Mollusques marins. Annales de l'Institut océanographique, Fondation Albert Ier, Prince
 465 de Monaco, 5 (3), 1-111.
- 466 Diedhiou, Y., Youm, C.I., Sow, E.H., Sarr, R. Malou, R., 2018. Les amas coquilliers d'âge
 467 holocène du lac Retba (Sénégal, Afrique de l'Ouest) : inventaire faunistique, lithologie
 468 et paléo-environnement Holocene. *Bulletin de l'Institut Scientifique, Rabat, Section*469 *Sciences de la Terre* 40, 39-52.
- 470 Drake, N.A., Blench, R.M., Armitage, S.J., Bristow, C.S., White, K.H., 2011. Ancient
 471 watercourses and biogeography of the Sahara explain the peopling of the desert.
 472 *Proceedings of the National Academy of Sciences* 108 (2), 458-462. DOI:
 473 10.1073/pnas.1012231108
- 474 Elouard, P., 1968. Le Nouakchottien. Etage du Quaternaire du Sénégal. Annales de la Faculté
 475 des Sciences de Dakar 22, 121-128.
- 476 Faegri, K., Iversen, J., 1977. Textbook of Pollen Analysis. Blackwell, Oxford, 328 p.
- Faure, H., Fontes, J.C., Hébrard, L., Monteillet, J., Pirazzoli, P.A., 1980. Geoidal change and
 shore-level tilt along Holocene estuaries: Senegal River area, West Africa. *Science* 210
 (4468), 421-423. DOI: 10.1126/science.210.4468.421
- Fleming, K., Johnston, P., Zwartz, D., Yokoyama, Y., Lambeck, K., Chappell, J., 1998.
 Refining the eustatic sea-level curve since the Last Glacial Maximum using far-and intermediate-field sites. *Earth and Planetary Science Letters* 163 (1-4), 327-342. DOI: 10.1016/S0012-821X(98)00198-8
- Fofana, C.A.K., Sow, E., Lézine, A.-M., 2020. The Senegal River during the last millennium. *Review of Palaeobotany and Palynology* 275, 104-175. DOI:
 10.1016/j.revpalbo.2020.104175
- 487 GBIF database, https://www.gbif.org
- 488 Giovas, C.M., 2009. The shell game: analytic problems in archaeological mollusc
 489 quantification, *Journal of Archaeological Science* 36, 1557-1564. DOI:
 490 10.1016/j.jas.2009.03.017
- 491 Giovas, C.M., Clark, M., Fitzpatrick, S.M., Stone, J., 2013. Intensifying Collection and Size
 492 Increase of the Tessellated Nerite Snail (Nerita Tessellata) at the Coconut Walk Site,
 493 Nevis, Northern Lesser Antilles, AD 890–1440. *Journal of Archaeological Science* 40
 494 (11), 4024–38. DOI: 10.1016/j.jas.2013.05.008
- 495 Gregory, W.K., 1933. Fish skulls. A study of the evolution of natural mechanisms. *American* 496 *Philosophical Society* 23 (2), 1–481.
- 497 Grimm, E., 1991. *Tilia and Tiliagraph*. Illinois State Museum, Springfield, 52 p.
- 498 Grouard, S., 2001. Subsistance, systèmes techniques et gestion territoriale en milieu insulaire 499 antillais précolombien. Exploitation des Vertébrés et des Crustacés aux époques

- Saladoïdes et Troumassoïdes de Guadeloupe (400 av. J.-C. à 1 500 ap. J.C.). PhD,
 Paris X Nanterre University, 1073 p.
- Heaton, T.J., Köhler, P., Butzin, M., Bard, E., Reimer, R.W., Austin, W.E.N., Bronk Ramsey,
 C., Grootes, P.M., Hughen, K.A., Kromer, B., Reimer, P.J., Adkins, J., Burke, A., Cook,
 M.S., Olsen, J., Skinner, L.C. 2020. Marine20-the marine radiocarbon age calibration
 curve (0-55,000 cal BP). Radiocarbon 62. DOI : 10.1017/RDC.2020.68
- Hébrard, L., 1972. Un épisode Quaternaire en Mauritanie (Afrique occidentale) à la fin du
 Nouakchottien: le Tafolien 4000-2000 ans avant le présent. *Bulletin de l'Association sénégalaise pour l'étude du Quaternaire ouest africain*, Dakar 33-34, 5-16.
- Hély, C., Lézine, A.-M., and APD, 2014. Holocene changes in African vegetation: tradeoff
 between climate and water availability. *Climate of the Past* 10 (2), 681-686. DOI:
 10.5194/cp-10-681-2014
- Hoelzmann, P., Jolly, D., Harrison, S. P., Laarif, F., Bonnefille, R., Pachur, H.J. 1998. MidHolocene land-surface conditions in northern Africa and the Arabian Peninsula: A data
 set for the analysis of biogeophysical feedbacks in the climate system. *Global Biogeochemical Cycles 12* (1), 35-51. DOI : 10.1029/97GB02733
- Hooghiemstra, H., 1988. Changes of major wind belts and vegetation zones in NW Africa
 20,000–5000 yr B.P., as deduced from a marine pollen record near cap blanc. Riview of
 Palaeobotany and Palynology 55 (1-3), 101-140. DOI: 10.1016/0034-6667(88)90056-5
- Humann, P. and DeLoach, N., 1993. *Reef Coral identification, Florida, Caribbean, Bahamas.*New World Publications Inc., Jacksonville, Florida, 276 p.
- Hutchinson, J., Dalziel, J.M., 1954-1972. *Flora of West Tropical Africa*. Vols 1-3.
 Whitefriars, London.
- Lebrun, J.-P., 1973. Énumération des plantes vasculaires du Sénégal. GERDAT-IEMVT,
 215 p.
- Lebrun, J.-P., Stork, A.L., 1991-2015. Énumération Des Plantes à Fleurs d'Afrique Tropicale *et Tropical African Flowering Plants: Ecology and Distribution*. Vols. 1-7.
 Conservatoire et Jardin botaniques de la Ville de Genève, 256 p.
- 528 Lézine, A.-M., 1987. Paléoenvironnements végétaux d'Afrique nord-tropicale depuis 12 000
 529 B.P. Analyse polliniques de series sédimentaires continentals (Sénégal-Mauritanie).
 530 PhD Aix-Marseille II University. 2 vols.
- Lézine, A.-M., 1988. New pollen data from the Sahel (Lake Guiers Senegal). *Review of Palaeobotany and Palynology* 55, 141-154. DOI: 10.1016/0034-6667(88)90082-6
- 533 Lézine, A.-M. 1989. Late Quaternary vegetation and climate of the Sahel. *Quaternary* 534 *Research*, 32 (3), 317-334. DOI : 10.1016/0033-5894(89)90098-7
- Lézine, A.-M., 1997. Evolution of the West African mangrove during the late Quaternary: a
 review. *Géographie Physique et Quaternaire* 51, 405-414. DOI : 10.7202/033139ar
- 537 Lézine, A.-M., Bieda, S., Faure, H., Saos, J.-L., 1985. Etude palynologique et
 538 sédimentologique d'un milieu margino-littoral: la tourbière de Thiaye (Sénégal).
 539 Palynological and sedimentological studies of a coastal environment: the peat bog of
 540 Thiaye (Senegal). Sciences Géologiques, bulletins et mémoires 38 (1), 79-89.
- Lézine, A.-M., Hély, C., Grenier, C. Braconnot, P., Krinner, G., 2011. Sahara and Sahel
 vulnerability to climate changes, lessons from Holocene hydrological data. *Quaternary Science Reviews* 30 (21-22), 3001-3012. DOI: 10.1016/j.quascirev.2011.07.006
- Lézine, A.-M., Lemonnier, K., Waller, M.P., Bouimetarhan, I., Dupont, L., 2021. Changes in
 the West African landscape at the end of the African Humid Period. *Palaeoecology of Africa* 35, 65-83. DOI : 10.1201/9781003162766-6
- 547 Manca, L., 2017. La fracturation et la fragmentation des coquilles. Une problématique
 548 partagée entre archéozoologie, taphonomie et technologie. *In* : Christensen, M., Goutas,
 549 N., (eds.) À *coup d'éclats ! La fracturation des matières osseuses en Préhistoire :*

- 550 *discussion autour d'une modalité d'exploitation en apparence simple et pourtant mal* 551 *connue, Séances de la Société préhistorique française* 13, Paris, 43-53.
- Médus, J., 1975. Spectres palynologiques de deux sédiments du Nouakchottien des environs
 de Saint-Louis (Sénégal). Bulletin de l'Institut Fondamental d'Afrique noire 37, A (3),
 554 532-536.
- Médus, J., Monteillet, J., 1981. Données préliminaires sur la palynologie de l'Holocène du
 delta du fleuve Sénégal. *Geobios* 14 (6), 801-805. DOI: 10.1016/S0016-6995(81)801532
- Mercuri, A.-M., 2008. Human influence, plant landscape evolution and climate inferences
 from the archaeobotanical records of the Wadi Teshuinat area (Libyan Sahara). *Journal of Arid Environments* 72, 1950-1967. DOI : 10.1016/j.jaridenv.2008.04.008
- Michel, P., Assémien, P., 1969. Etudes sédimentologique et palynologique des sondages de
 Bogué (basse vallée du Sénégal) et leur interprétation morphoclimatique. *Revue de Géomorphologie dynamique* 3 (19), 97-113.
- 564 Michel, P., 1973. Les bassins des fleuves Sénégal et Gambie : étude géomorphologique.
 565 ORSTOM, Paris, 571 p.
- 566 MNHN databases, https://www.mnhn.fr; https://www.mnhn.fr/fr/bases-de-donnees 567 scientifiques-naturalistes; https://inpn.mnhn.fr/accueil/donnees-referentiels
- MolluscaBase (eds), 2022. MolluscaBase. https://www.molluscabase.org on 2022-09-09.
 doi:10.14284/448
- Monod, T., 1983. Sur la présence de graines de *Celtis* dans le Quaternaire de Mauritanie.
 Notes africaines 178, 3-4.
- Monteillet, J., Faure, H., Pirazzoli, P.A., Ravisé, A., 1981. L'invasion saline du Ferlo
 (Sénégal) à l'Holocène supérieur (1900 BP). *Palaeoecology of Africa* 13, 205–215.
- Ndeye, M., 2008. Marine reservoir ages in northern Senegal and Mauritania coastal waters.
 Radiocarbon 50 (2), 281–288. DOI: 10.1017/S0033822200033580
- 576 Parsons, K.M., Brett, C.E., 1991. Taphonomic Processes and Biases in Modern Marine
 577 Environments: An Actualistic Perspective on Fossil Assemblage Preservation. In:
 578 Donovan, S.K., (ed.) *The Processes of Fossilization*, Columbia University Press, New
 579 York, 22-65.
- 580 Past 4.03 software, https://www.nhm.uio.no
- 581 Pielou, E.C., 1966. Shannon's formula as a measure of specific diversity: Its uses and
 582 misuses. *American Naturalist*, 100 (914), 463–5.
- 583 Pointier, J.-P., Lamy, D., 1998. *Guide des coquillages des Antilles*. Découvre la Mer
 584 Collection, PLB Ed. Abymes, 225 p.
- Reimer, P., Austin, W.E.N., Bard, E., Bayliss, A., Blackwell, P.G., Bronk Ramsey, C., 585 586 Butzin, M., Cheng, H., Edwards, R.L., Friedrich, M., Grootes, P.M., Guilderson, T.P., 587 Hajdas, I., Heaton, T.J., Hogg, A.G, Hughen, K.A., Kromer, B., Manning, S.W., Muscheler, R., Palmer, J.G., Pearson, C., van der Plicht, J., Reimer, R.W., Richards, 588 D.A., Scott, E.M., Southon, J.R., Turney, C.S.M., Wacker, L., Adolphi., F, Büntgen, U., 589 590 Capano., M, Fahrni, S., Fogtmann-Schulz, A., Friedrich, R., Köhler, P., Kudsk, S., 591 Miyake, F., Olsen, J., Reinig, F., Sakamoto, M., Sookdeo, A., Talamo, S. 2020. The 592 IntCal20 Northern Hemisphere radiocarbon age calibration curve (0-55 cal kBP). 593 *Radiocarbon* 62. DOI : 10.1017/RDC.2020.41.
- Roger, J., Noël, B.J., Barusseau, J.-P., Serrano, O., Nehlig, P., Duvail, C., 2009. Notice *explicative de la carte géologique du Sénégal à 1/500 000, feuilles nord-ouest, nord-est et sud-ouest*. Ministère des mines, de l'industrie et des PME, Direction des Mines et de
 la Géologie, Dakar, 61 p.

- Serrand, N., 2002. Exploitation des invertébrés marins et terrestres par les populations
 Saladoïdes et post-Saladoïdes du Nord des Petites Antilles (env. 500 B.C. 1200 A.D.).
 Etude de cas et comparaisons. PhD, Paris I Panthéon Sorbonne, Paris, 1317 p.
- Spalding, M., Blasco, F., Field, C., 1997. *The mangrove atlas*. The International Society for
 Mangrove Ecosystems, Okinawa, Japan, 178 p.
- Stuiver, M., Reimer, P.J., 1993. Extended 14C Data Base and Revised CALIB 3.0 14C Age
 Calibration Program CALIB rev. 8. *Radiocarbon* 35, 215-230. DOI:
 10.1017/S0033822200013904
- Taylor, J., Glover, E., 2021. *Biology, evolution and generic review of the chemosymbiotic bivalve family Lucinidae*. The Ray Society 182, London, 320 p.
- Tomadini, N., 2018. Hommes et animaux dans les colonies françaises des petites Antilles du *XVIIe siècle à la fin du XIXe siècle : changements, résiliences et adaptations mutuelles.*PhD, MNHN, Paris, 492 p.
- Trochain, J., 1940. Contribution à l'étude de la végétation du Sénégal. Mémoires de l'Institut
 Français d'Afrique Noire. Larose, Paris, 433 p.
- Vache-Grandet, C., 1959. Problème agraire en Mauritanie : l'aménagement du lac Rkiz. Les
 Cahiers d'Outre-Mer 12 (47), 251-264.
- Vincens, A., Lézine, A.-M., Buchet, G., Lewden, D., Le Thomas, A., 2007. African Pollen
 Database inventory of tree and shrub pollen types. *Review of Palaeobotany and Palynology* 145 (1-2), 135-141. DOI: 10.1016/j.revpalbo.2006.09.004
- Walter, L.M., Burton, E.A., 1990. Dissolution of Recent Platform Carbonate Sediments in
 Marine Pore Fluids, American Journal of Science 290, 601-643. DOI:
 10.2475/ajs.290.6.601
- Watrin, J., Lézine, A.-M., Hély, C., 2009. Plant migration and ecosystems at the time of the
 "green Sahara". *Comptes Rendus Geosciences* 341, 656-670. DOI:
 10.1016/j.crte.2009.06.007
- White, F. 1986. La végétation de l'Afrique : mémoire accompagnant la carte de végétation de
 l'Afrique, UNESCO/AETFAT/UNSO, ORSTOM Paris, 385 p.
- WoRMS Editorial Board, 2022. World Register of Marine Species. DOI:10.14284/170
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629 List of Figures

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- 631 Figure 1: Lake Rkiz. A: The general map is from en-us.topographic-map.com. The dashed
- 632 rectangle indicates the Lake Rkiz area detailed in Figure 1B. B: The detailed map of Lake
- Rkiz is redrawn after Vache-Grandet (1959). The shaded area within the lake indicates the
- area of the lowest level identified by the author in 1954. The Lake was entirely dry at the timeof coring. The star indicates the location of core Rkiz1 (this study).
- 636
- Figure 2: Core Rkiz1: Lithostratigraphy and ¹⁴C ages.
- 639 Figure 3: Core Rkiz 1 Minimum Number of Individuals (MNI) versus age. From left to right,
- 640 this figure shows the MNI of all the faunal remains encountered. On the right, the
- 641 preservation index and the Shannon diversity index are shown.642
- Figure 4: Core Rkiz 1 synthetic diagram of MNI of the mollusk taxa *versus* age. On the right,
 the preservation and diversity indexes are shown. Pollen zones are shown to the right of the
 figure for easy comparison.

646

- 647 Figure 5: Core Rkiz 1 synthetic diagram showing, versus age from left to right, the main
- 648 pollen and NPP taxa. Percentages are shown in black, gray shading shows the same results,

multiplied by 10 for highlighting. Mollusk zones are shown to the right of the figure for easycomparison.

651

Figure 6: Location map of sites attesting to the marine influence in the area of the Senegal
River and its delta during the Holocene. Black circles indicate pollen sites, gray circles
indicate marine shells, and white squares represent peat samples. At Lake Rkiz, the black
square show our coring site, while the diamonds indicate shell samples from Faure et
al.(1980). Satellite image from Google Earth.

658 Figure 7: Dated evidence of marine intrusion along the Senegal River and Lakes Guiers and 659 Rkiz (S-Table 3). The colored circles correspond to dates extracted from marine shells collected in 3 major locations: from the West and East: the delta, Lake Guiers and the Ferlo 660 river, and Lake Rkiz. Red circles correspond to the dates of this study at Lake Rkiz, pink 661 662 circles to those of Faure et al. (1980). Squares correspond to the dates of the long cores from 663 the bed of the Senegal River (white squares are dates on bulk sediment (peat); blue squares 664 are dates from shells. A: the dates are presented as a function of time and longitude in order to evaluate the distance of the marine intrusion from the coast, B: as a function of (non 665 666 corrected) altitude/depth when indicated by the authors. The dark gray arrows indicate the 667 average position of the two Holocene marine transgressions. The light gray double arrow highlights the difference in depth between the sea level records in the Senegal River bed 668 669 (Eudoxie, Leo, Keur Marsal) and in Lake Rkiz.

670

Figure 8: Synthetic diagrams showing the main features of the local (on the right) and 671 672 regional (on the left) environment of Lake Rkiz from 10 ka to the present. On the right, 673 mangrove pollen percentages (dark purple) are calculated against the sum of all pollen and 674 fern spores. Light shading shows the same results, multiplied by 10 for highlighting. 675 Percentages of mangrove (purple triangles) and marine (black dots) fauna are calculated 676 against the sum of all the determined mollusk shells (in black: power trend line). Only 677 samples with more than 20 determined remains are represented. On the left, pollen data are 678 presented under the form of phytogeographical group of plants from the most humid 679 (Guinean/Sudanian tropical taxa) to the most arid (Saharan desert) (aquatics and ferns 680 excluded from the pollen sum).

681

682 Figure 9: Modern (red dots) and Holocene (green dots) distribution of *Celtis toka* in tropical

683 West Africa. Pollen data are from the APD (https://africanpollendatabase.ipsl.fr), modern data

are from the GBIF (https://www.gbif.org) and MNHN (https://www.mnhn.fr) databases, and
the literature (Trochain, 1940; Hutchinson and Dalziel, 1954-1972). Simplified vegetation
map from White (1986).



















