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Anne-Marie Lézine, Kévin Lemonnier, Cheikh Abdoul Kader Fofana. Sahel environmental variability during the last millennium: Insight from a pollen, charcoal and algae record from the Niayes area, Senegal. Review of Palaeobotany and Palynology, 2019, 271, pp.104103. 10.1016/j.revpalbo.2019.104103 . hal-02370143

HAL Id: hal-02370143 https://hal.science/hal-02370143

Submitted on 20 Dec 2021

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1	Sahel environmental variability during the last millennium: insight from a pollen, charcoal
2	and algae record from the Niayes area, Senegal
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13	Abstract :
14	The pollen, algae and charcoal record from the "Baobab" depression near Mboro in the
15	Niayes area of Senegal shows that the last millennium was characterized by a gradual
16	aridification which started 1340 CE and culminated 1845 CE. The sub-Guinean gallery forests,
17	which survived the previous environmental crisis dated from the end of the Holocene Humid
18	Period, suddenly collapsed and the Niayes region suffered a dramatic loss of biodiversity. We
19	show that the tipping point between a forested and non-forested state occurred ca 400
20	centuries after the onset of aridity recorded by the lowering of the water level and the
21	progressive salinization of the aquatic environment.
22	
23	Key words : Pollen, Algae, Charcoal, last millennium, Sahel, Senegal

24

1. Introduction

26

Was the last major Sahelian drought in the 1970s unusual? By studying records of 27 environmental changes and associated forcing mechanisms in the recent past, it is possible 28 to situate this major climate crisis in the context of the natural variability of the west African 29 30 monsoon climate system. Natural archives such as lake sediments can fill gaps in understanding climate variability by providing access to a long-term perspective 31 32 encompassing periods out of any anthropogenic influence. In western Africa, such long-term records are rare (e.g., Nash et al., 2016 and references therein) and somewhat contradictory. 33 In particular, the well-known interval of the Little Ice Age (LIA - 1400-1850 CE) when 34 exceptionally cool temperatures were reached in the Northern hemisphere, was described 35 36 as a dry period in the Lake Bosumtwi area (Ghana) (Shanahan et al., 2009) and a humid period in the Saloum delta (Senegal) (Carré et al., 2018). 37

The littoral depressions of the Niayes area, along the northern coast of Senegal have 38 provided major information on the Holocene history of the West African environments 39 (Lézine et al., 2011; Hély and Lézine, 2014). Local environmental conditions linked to the 40 41 proximity of the coast have favored accumulation of organic matter and expansion of gallery forests in the bottom of the depressions during the Holocene (Lézine, 1989) which have 42 43 persisted up to recently thanks to the high level of the water table (Trochain, 1940; Pézeril et al., 1986). It has been shown that these gallery forests were highly sensitive to variations of 44 the level of the water table whatever the temporal resolution considered, from decades 45 46 (Aguiar, 2010) to centuries (Lézine and Chateauneuf, 1991). Here we present a study combining pollen and diatom analyses on a sedimentary series recovered NW of Mboro in 47 the center of the Niaye area. The aim of our study is to examine the hydrological variations 48

- during the last millennium and to discuss their impact on the local vegetation. We also aim
 at characterizing the signature of human activities on the natural environment.
- 51

52 **2. Regional setting**

The Niaye area consists of depressions formed between linear fixed and partially fixed 53 54 coastal dune ridges, located between 15° and 16° N, at roughly 17°W along the Atlantic coast of Senegal. These depressions belong to an extensive river system which took place during 55 the Holocene Humid Period ca 12 ka yr BP. (Lézine, 1989). Edification of coastal dunes 56 (Michel, 1973) gradually closed access to the sea of water courses and promoted the 57 development of thick, continuous peat accumulations. The presence of fresh, shallow 58 ground-water near the surface in the bottom of the interdune depressions (Lézine and 59 60 Chateauneuf, 1991) supports the preservation (though deeply degraded today by intensive agriculture activity) of gallery forests of Guinean phytogeographical affinity, which are the 61 last traces of the northern extension of tropical humid forests during the Holocene (Watrin 62 et al., 2009; Hély and Lézine, 2014). This is the "sub-Guinean domain" of Trochain (1940) 63 consisting of enclaves of aquatic vegetation in open water areas (Nymphaea, Potamogeton, 64 Pistia) and reed-swamp populations (Typha, Phragmites) associated with Syzygium 65 guineense in the flooded zone, surrounded by Guinean forest relicts at the foot of the dunes 66 67 (Elaeis guineensis, Ficus, Treculia, Phoenix) followed by Dodonaea, Alchornea, Zanthoxylum, Combretum toward the upper limit of the gallery forests, whereas typically Sahelian trees 68 grow at the top of the dunes (Acacia, Balanites) (Adam, 1953, 1958; Jaeger, 1949; Villiers, 69 70 1957; Michel et al., 1969; Raynal, 1963) (Figure 1A).

Hydrological studies in the Niayes area have shown that the water level varies greatly in
 phase with local rainfall. Accentuated dryness associated with intensive water pumping in

recent decades has led to the lowering of the fresh water table and the penetration of salt
waters from the adjacent sea into the depressions closest to the coast (e.g., Faye, 1995;
Aguiar, 2008; Maugis et al., 2009).

76

77 3. Material and methods

A 4 m long sediment core was collected in the center of the so-called "Baobab" depression in 2016 (15.149132N; 16.909275W) under few tens cm of water using a "Russian" manual corer (Jowsey, 1966). Baobab lies SW of the main paleoriver channel of M'Boro at the NW extremity of a depression separated from the sea shore by a ridge of ca 2 km wide dunes of recent Holocene age (Michel, 1973) (Figure 1B). The sediment consists of homogeneous organic clay. Samples were taken at 5 cm intervals for pollen and algae analyses.

84 For pollen and non-pollen palynomorph (NPP) analyses samples were processed using the 85 standard HF method described by Faegri and Iversen (1977). After sieving in a 5 µm mesh, the residue was stained with safranin and mounted in glycerine. Pollen identification was 86 based on the reference slide and photo collections of the LOCEAN laboratory and the 87 Museum national d'Histoire naturelle in Paris as well as on pollen atlases of tropical African 88 flora from e.g., Maley (1970), APLF (1974), Ybert (1979) and Bonnefille and Riollet (1980). 89 Pollen nomenclature, which follows the standard defined by the African Pollen Database 90 91 (Vincens et al., 2007), was based on Lebrun and Stork (1991-2015). The pollen microflora 92 was composed of 86 taxa (Table 1): 44 trees, shrubs, palms and lianas, 17 herbs and 14 undifferentiated taxa mainly because of the low level of determination of the grains or when 93 94 the family or genus level corresponded to several life-forms. In addition, 4 aquatic pollen taxa and 4 ferns were encountered. Pollen taxa were grouped according to the 95 phytogeographical affinity of the parent species following Lézine et al. (2009) in four entities 96

according a decreasing rainfall gradient: Guineo-congolian, Sudanian, Sahelian and Saharan.
In addition, some taxa have been selected as indicators of water-table fluctuations according
to their distance from the open water area (Raynal, 1963; Michel et al., 1969; Lézine, 1987)
(Fig. 2).

Pollen percentages were calculated against a sum excluding aquatics and ferns, the percentages of which were calculated separately against a sum including all the pollen grains and fern spores counted.

Undifferentiated fungi and 4 green algae (*Botryococcus, Coelastrum, Pediatrum, Tetraedron*)
 represent the NPP group. NPP are presented as percentages (Fig. 3B) and micro-charcoal as
 influxes (nbr of particles/cm²/year) (Fig.4).

For diatom analyses samples were processed using a standard method (Battarbee, 1986). 107 108 The residu was diluted in 20 ml of water and 2 ml was taken, then dried and fixed with Naphrax. Diatom identification was based on e.g., Guermeur (1954), Servant-Vildary (1978), 109 Foged (1986), Gasse (1986), Compère (1991), Cocquyt et al. (1993), Bellinger and Sigee 110 (2015). In addition, Ouattara et al. (2000), Sow and Diène (2002), Sow (2001; 2004), Sow et 111 112 al. (2005) were used to precise the ecological requirement of taxa, particularly in Senegal. 113 Twenty-two diatoms were determined (Table 2). They correspond to three main hydrological freshwater/salt tolerant, 114 conditions: freshwater, brackish/marine waters and 115 undifferentiated. Diatoms are presented as percentages calculated against the sum of all the diatoms counted (Fig. 3A). 116

Both pollen and diatom data are presented under the form of diagrams using Tilia and CONISS (Grimm, 1991). Diversity indices were derived from PAST 3.23 software (Hammer et al., 2001).

120

121 **4. Age control**

Age control was established by 3 accelerator mass spectrometry (AMS) radiocarbon dating of total organic matter on the whole Baobab sequence, conducted at UMS-ARTEMIS AMS Facilities (France). An additional control point was given by the first occurrence of *Casuarina equisetifolia* pollen grains in our samples the parent trees of which were planted in 1948 in the Niaye area (Ndiaye et al., 1993) (Table 3). According to this, the last millennium is represented by the 121 uppermost centimeters of the 4m-long Baobab sedimentary section. Our analysis has a temporal resolution varying between 70 and 12 years.

129

130 **5. Results**

The pollen (Fig. 2), diatom (Fig. 3A) and NPP (Fig. 3B) records display several phases of environmental change over the last millennium with a remarkable coherency. They are described from the base to the top of the sequence.

134

135 **5.1 Pollen**

136 The pollen spectra are characterized by the dominance of Poaceae and the continuous 137 occurrence of Chenopodiaceae/Amaranthaceae and Asteraceae. This, together with the high number of other taxa, mainly herbaceous, from steppes, grasslands and wooded grasslands 138 139 point to an open and fairly dry regional landscape. Except Poaceae, all the other taxa are poorly represented. This is particularly true for tropical trees which are characterized by a 140 low pollen productivity and an enthomophilous way of pollen dispersal (e.g., Watrin et al., 141 142 2007). As a consequence, their percentages are rarely higher than 10% (e.g., Alchornea). 143 Zone 1, base-1340 CE: Zone 1 is typified by tropical tree taxa, e.g., Alchornea, Macaranga,

Allophylus, Detarium and Mitragyna inermis which reach their highest values (%). Aquatics

(Potamogeton, Nymphaea) are also present. All this depict an open water surface surrounded by a relatively developed gallery forest. By the end of this zone, in sub-zone 1b (1235-1340 CE) tropical trees start to decline whereas *Typha* starts to increase showing the onset of lake level lowering under drier conditions. Zone 1b marks the onset of *Elaeis guineensis* development.

Zone 2, 1340-1810 CE: Some tropical tree taxa disappear (*Allophylus, Mitragyna inermis*) or decline (*Alchornea, Macaranga*). In contrast, *Syzygium guineense* increases, as well as taxa from reed-swamp populations (*Tyhpa*, ferns) both recording low water stands and marsh encroachment.

Zone 3, 1810-1945 CE: This zone shows a dramatic environmental change with the disappearance of most of the plant taxa replaced almost exclusively by Poaceae (91%). As a consequence, biodiversity falls sharply and Simpson diversity index values, which fluctuated only slightly around 0.6 in the lowermost zones, fall to 0.177 at level dated 1845 CE. The peak of Fungi at this level (Fig. 3B) testify for the at least partial or seasonal desiccation of the Baobab depression.

160 Zone 4, 1945-2016 CE: This zone is characterized by the increase in markers of 161 anthropogenic activity with the appearance of pollen from the planted tree Casuarina equisetifolia. Charcoal remains, which steadily increased since 1370 CE, peaked at 1989 CE, 162 163 suggesting the dramatic increase in fire occurrence. Increase in diversity compared to zone 3 as well as significant percentages of tropical tree taxa suggests a certain recovery of the 164 gallery forest. However, these taxa never reach values as high as in zones 1 and 2. Typha and 165 166 Syzygium show that the level of the water table remained low during this zone. The increase of Suaeda in the uppermost levels (post 1980 CE: zone 1b) points to the setting of saline 167 conditions. 168

169

170 **5.2 Algae**

171 **5.2.1 Diatoms**

Aulacoseira granulata and Nitzschia (N. acicularis and N. scalaris) are the most dominant
 species and account for up to 95, 49 and 23% respectively (Fig. 3A).

Zone 1, base-1340 CE: the diatom assemblage is almost exclusively represented by *A. granulata*, a planktonic species characteristic of well mixed fresh waters. This taxon is associated with *N. acicularis*, which confirms the presence of enriched turbid waters (zone 1a). Towards the top of Zone 1 at levels dated 1235 CE and above, Zone 1b shows an increase in diversity with *N. scalaris, Eunotia pectinalis* and *Thalassiosira decipiens* showing the appearance of more brackish/saline conditions.

Zone 2, 1340-1925 CE: This zone is characterized by the gradual decline of *A. granulata*,
 which reached its lowest values in year 1906 EC. Three main taxa successively increase: *N. scalaris* (Zone 2a), *T. decipiens* (Zone 2b), then *N. acicularis* (Zone 2c) suggesting the gradual
 increase of dry conditions initiated in Zone 1b.

Zone 2a (1340-1600 CE) is characterized by a noticeable increase in diversity and
 the appearance of periphytic (e.g., *Fragilaria construens*) or bentic (e.g., *F. pinnata, Pinnularia neomajor*) species. That together with the presence of species
 either cosmopolite or salt tolerant point to the lowering of the water level and
 the setting of dry conditions.

2one 2b (1600-1800 CE): *N. scalaris* decrease and is replaced by *T. decipiens* and
 C. stelligera showing that salt waters increased in the aquatic environment.

- Zone 2c (1800-1925 CE): *N. acicularis* dominate the diatom assemblage is
 association with *T. descipiens*. Meanwhile *A. granulata* reaches its lowest values.
 All this suggests that the water level reached its lowest position.

In Zone 3, 1925-2001 CE: The increase of *A. granulata* in the upper levels of the core suggests a certain recovery of the fresh water environment. This remained however highly variable and salty conditions are suggested in the upper level (Zone 3b) by the reappearance of *T. decipiens* and *S. Tenera* which were absent from Zone 3a.

198

199 **5.2.2 Green algae**

The green algae record (Fig. 3B), shows the opposite trend between *Tetraedron*, *Pediatrum*, *Coelastrum* which gradually decrease from the base to the top of the sequence, though fluctuating, and *Botryococcus*. This pattern illustrates the gradual depletion of nutrient and lowering of the water level. The absence of any other green algae except *Botryococcus* at levels post-1954 CE indicates the emergence of clearly oligotrophic conditions. This nutrient poor environment immediately followed the phase of lake level lowering indicated by the increase of fungi peaking at 1845 CE.

207

208 **6.** Discussion

209 Three main features emerge from our study:

210 **6.1** The dramatic collapse of the Guineo-Congolian gallery forest at 1640 CE.

The end of the HHP, from 4500 cal yrs BP onward (Lézine et al., 2011), was marked the decline of tropical plants that previously widely expanded throughout the Sahel and the Sahara (Watrin et al., 2009; Hély et al. 2014). The abruptness of the forest collapse in the Niayes area (Lézine, 1989) was interpreted as the result of crossing a hydrological threshold, 215 after a long period of increased seasonality in rainfall since the mid-Holocene. The tipping point between a forest environment and an open and degraded environment occurred ca. 216 2500 cal yrs BP (Lézine, 1989), i.e. approximately 2000 years after the end of the HHP in 217 218 Western Africa. The specific hydrological conditions due to the proximity of the ocean led to maintain a high level of the groundwater at the bottom of the depressions. They 219 220 compensated for a possible precipitation deficit by maintaining moisture conditions allowing 221 the forest to maintain at the foot of the dunes for an additional two millennia after the end of the HHP. 222

Over the past millennium, tropical plant types stabilized at 30%, reflecting that sub-Guinean 223 224 gallery forests remained only moderately developed in the Niayes area as compared to the HHP during which these values were at least 80% (Lézine, 1989). These % dramatically fall at 225 226 1640 CE to a minimum at 1845 CE pointing to the setting of a severe environmental crisis: 227 the biodiversity dramatically declined and the Baobab depression dried up, at least partially. The disappearance of green algae and their replacement by fungi in the lake sediments as 228 well as the high values of salt-tolerant diatoms clearly show that hydrological conditions 229 230 were more evaporated including pronounced seasonal drying of the depression (Fig. 4). This drought event in the Niayes area of Senegal is contemporaneous with the repeated dry years 231 in the inner delta of the Niger River as recorded by documentary sources since 1639 232 (Coquery-Vidrovitch, 1997) and continuing into the 19th century, particularly between years 233 1800 and 1850 CE (Nicholson et al., 2012 in Nash et al., 2016). Similarly, the level of Lake 234 Chad underwent multiple lowering phases (Maley, 1976) and drought conditions developed 235 at Lake Bal, Nigeria (Holmes et al., 1999) during the 17th century. 236

237 6.2 The drying trend from 1375 CE to 1900 CE

The forest collapse 1640 CE occurred after several centuries of increasing drought starting from 1340 CE, as revealed by the development of marshy plant populations and of saline conditions in the Baobab depression.

The most prominent feature of the hydrological evolution of the Baobab depression during the last millennium is the development of *Syzygium* and *Typha* populations both resulting from the lowering of the water table responsible for a marsh development. As the water level dropped, *Syzygium* trees were able to expand towards the center of the depression. This situation also occurred in the Niayes area during the dry phases of the Holocene as shown by the well preserved woods of *Syzygium* trees north of Baobab, around Diogo (Figure 5).

The Syzygium development at Baobab occurred during the time interval 1340-1810 CE with a 248 249 peak at 1490 CE. As the level dropped further, these trees progressively declined and were 250 replaced by reed-swamp (Typha) populations that reached their maximum at 1688 CE. The Typha:Aquatic Pollen ratio used to illustrate the hydrological evolution of the Baobab 251 depression (Figure 4B) confirms the gradual lowering of the water level. This evolution is also 252 confirmed by the emergence of benthic and periphytic diatom taxa reflecting the lowering of 253 the water level during the time-interval 1600-1925 CE (Diatom Zone 2b and 2c), i.e., during 254 the drought event revealed by the pollen and fungi record. In the meantime, diatom taxa of 255 256 cosmopolite or saline ecological tolerance reached their maximum testifying for increased evaporation. The development of the freshwater taxon *Typha* at the lake margin during this 257 period is not incompatible with the increase in water salinity revealed by the diatomaceous 258 259 flora since Typha can tolerate brackish water for long periods of time and even develop on dry soils affected by seawater infiltration (Adam, 1964). 260

Our Baobab record does not show the humid event described in the Sahel by Maley (1977) at lake Chad around 1600 CE and Carré et al (2018) in the Saloum area of Senegal, 1500-1800 CE as well as in Eastern Africa by e.g., Verschuren et al., 2000 and Stager et al., 2005. However, it closely matches the Lake Bosumtwi record (Shanahan et al., 2009) which identifies a drought event between 1400 and 1750 CE, *"similar in timing to the Little Ice Age* (*LIA, ~1400 to 1850 CE*)". From our record, the LIA was a period of increased dryness in the Niaye area of Senegal with a particularly severe drought event peaking at 1845 CE.

268 **6.3 The anthropogenic impact**

Micro-charcoal fragments are insignificant in number at levels prior to 1600 CE. Then, during the LIA, they increase slightly, reflecting repeated fires during marked and probably longer dry seasons. In contrast, their unprecedented increase over the past two decades suggests a radical change in regional economic activities, probably in relation to the intensification of charcoal production around the cities of Thiès and St-Louis (Gueye, 2000).

274

275 **7. Conclusion**

Our pollen, charcoal and algae record of the Baobab sedimentary sequence near Mboro 276 shows that, during the last millennium, the Niayes area of Senegal underwent a gradual 277 aridification which culminated 1845 CE. This drying trend occurred during the Little Ice Age 278 279 during which the Northern Hemisphere experienced cool climate conditions. In the Niayes area, the sub-Guinean gallery forests, which survived the previous environmental crisis 280 dated from the end of the Holocene Humid Period 2500 years ago, suddenly collapsed and 281 282 the region suffered a dramatic loss of biodiversity. We show that the tipping point between a forested and a non-forested state occurred ca 400 centuries after the onset of aridity 283 recorded by the lowering of the water level and the progressive salinization of the aquatic 284

environment. This response time of tropical ecosystems to climate change can have critical
 implications for biodiversity and forest management strategies.

287

288 Acknowledgements

This work was initiated in the frame of a CNRS INSU LEFE research project headed by M. 289 290 Carré and an IRD scientific network coordinated by A. Sifeddine (CLIMACTE). It is an 291 "ACCEDE" Belmont Forum contribution (18 BELM 0001 05). Thanks are due to the Institut de Recherche pour le Développement (IRD) in Senegal and the UCAD Geological Department for 292 logistic support in the field and authorizations and to the ECLAIRS international laboratory, 293 Dakar and the LOCEAN laboratory for fundings. AML and KL are funded by CNRS, CAKF by 294 UCAD. Raw data will be stored at NEOTOMA and are also available on request from the first 295 296 author.

297

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422 **Table and Figure captions:**

Table 1: List of pollen taxa recovered in the Baobab last millennium sequence. Nomenclature
follows Vincens et al., 2007.

425 Table 2: List of Diatom taxa recovered in the Baobab last millennium sequence.

Table 3: Radiocarbon measurements (UMS-ARTEMIS AMS Facilities, France) and additional control points: 1948 CE: first appearance of *Casuarina equisetifolia* pollen grains in the sediments; 2016 CE: date of the core recovery).

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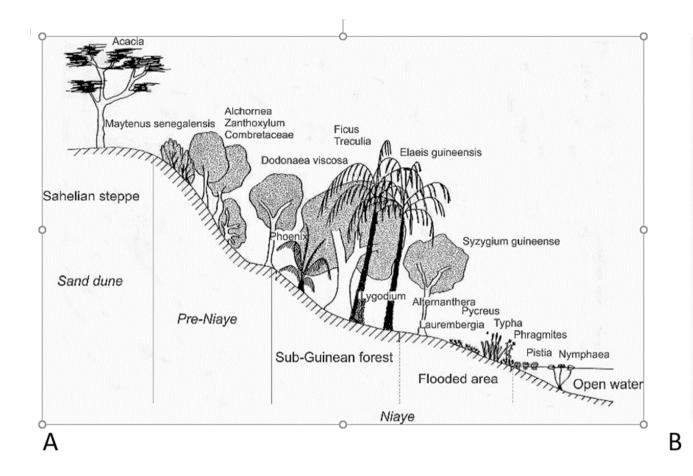
- Figure 1: The Niaye gallery forest. A: Schematic distribution of plants on dune slopes from the Sahelian steppes at the top of the dunes to the open-water surface at the bottom of the depressions (Raynal, 1963; Michel et al., 1969); B: The Mboro complex of paleodrainages and interdunal depressions filled by organic matter and the location of the study site (black dot) (ORSTOM, 1962).
- Figure 2: Synthetic pollen diagram of the Baobab (last millennium) showing from left to right
 versus time (in chronology CE) selected plant taxa. On the right, the Simpson D-1 index
 of diversity.

Figure 3: Synthetic diagram of the Baobab (last millennium) diatom and NPP record showing
from left to right versus time (in chronology CE) A: selected diatom taxa, main groups
of diatoms according to their ecological significance, the Simpson D-1 index of diatom
diversity and B: the NPP (Fungi and Green Algae) record.

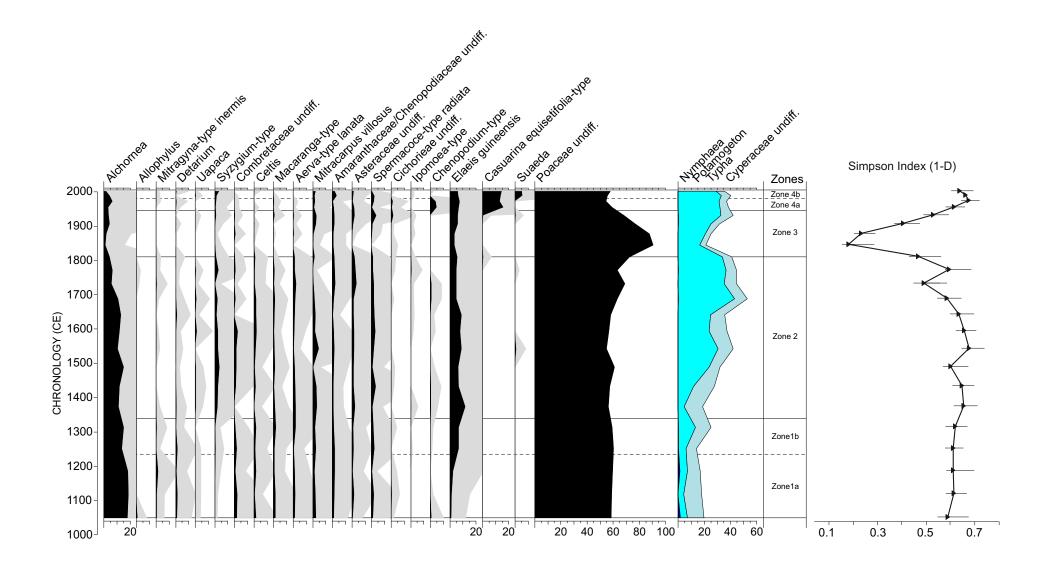
Figure 4: Main paleoenvironmental features at Baobab, (Mboro sector of the Niaye area,
Senegal) during the last millennium. A: Pollen and Fungi% and Charcoal influxes
illustrating the drying trend which led to the main environmental crisis peaking at 1845
CE in the Niayes area of Senegal and the increase of the anthropogenic activity during

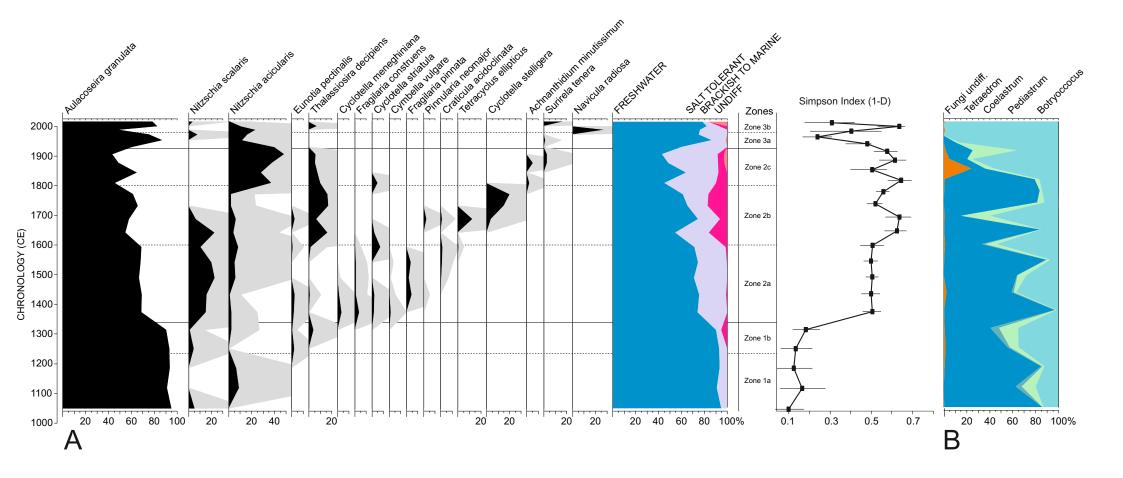
the last centuries; B: pollen markers of lake level lowering; C: diatom markers ofincreased salinity.

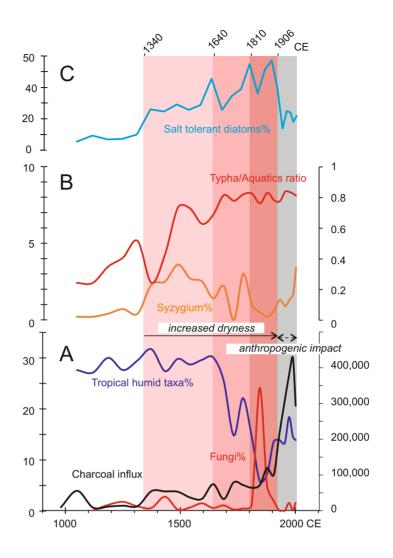
Figure 5: Fossil woods of *Syzygium guineense* of late Holocene age preserved in the organic levels of the Niaye sediments (Mandeng-Yogo, work in progress). A: the trunks emerge thanks to the digging of a well; B: the red arrows show the traces of the base of the *Syzygium* trees that formed a dense forest at the bottom of the interdunal depressions.















TREES and SHRUBS	UNDIFFERENCIATED
Acacia	Acalypha
Adansonia digitata	Aeschynomene
Alchornea cordifolia	, Amaranthaceae/Chenopodiaceae undiff.
Allophylus	Asteraceae undiff.
Anacardiaceae undiff.	Cichorieae undiff.
Anogeissus-type leiocarpus	Hibiscus
Anthostema-type	lpomoea-type
Bridelia-type	Justicia-type
Capparis fascicularis-type	Polygala-type
Casuarina equisetifolia-type	Rubiaceae undiff.
Celastraceae undiff.	Solanum-type
Celtis	Suaeda
Chrysobalanus icaco/Parinari	Triumfetta-type
Cissampelos-type	Undetermined
Cocculus pendulus-type	HERBS
Combretaceae undiff.	Achyranthes-type aspera
Combretum-type aculeatum	Aerva-type lanata
Combretum-type molle	Alternanthera
Commiphora africana-type	Centaurea-type perrottetii
Detarium senegalense	Cerastium-type
Dictyandra	Chenopodium-type
Diospyros	Cleome-type gynandra
Elaeis guineensis	Commelina-type benghalensis
Ficus	Commelina-type forskalaei
Hymenocardia	Gomphrena
Khaya-type senegalensis	Justicia-type flava
Lannea-type	Mitracarpus villosus
Macaranga-type	Poaceae undiff.
Mallotus-type	Polycarpaea-type
Mitragyna-type inermis	Polycarpon-type
Morelia senegalensis	Spermacoce-type radiata
Myrianthus-type	Tribulus
Paullinia pinnata	AQUATICS
Phoenix reclinata-type	Cyperaceae undiff.
Prosopis-type africana	Laurembergia tetrandra
Salacia	Ludwigia-type stolonifera
Sapotaceae undiff.	Nymphaea lotus-type
Syzygium-type	Polygonum senegalense-type
Tabernaemontana	Potamogeton
Tamarix	Typha
Tapinanthus-type	FERNS
Tetracera	Lygodium microphyllum
Tinospora bakis	Monolètes à épines et non lisses autres
Иараса	Monoletes undiff.
Zanthoxylum-type zanthoxyloides	Triletes undiff.

FRESHWATER

Aulacoseira granulata Craticula acidoclinata Cymbella vulgare Fragilaria construens Fragilaria pinnata Navicula densa Navicula radiosa Pinnularia divergens Pinnularia neomajor Tetracyclus ellipticus SALT TOLERANT

Achnanthidium minutissimum

Cyclotella meneghiniana Cyclotella stelligera Cyclotella striatula Eunotia pectinalis Nitzschia acicularis Nitzschia scalaris Rhopalodia gibba BRACKISH TO MARINE Navicula groveoides Surirela fastuosa

Thalassiosira decipiens

UNDIFF

Surirela tenera

Lab number SacA	Sample (cm)	Nature	Median Age CE	Range 1 σ
49827	BAOBAB 2016 (99- 100)	мо	1346	1317-1407
49828	BAOBAB 2016 (299- 300)	мо	-1547	1607-1503
49829	BAOBAB 2016 (399- 400)	мо	-2349	2454-2289

Additional control points (Age CE)						
	BAOBAB 2016 (0.5)	surface	2016			
		First record of				
	BAOBAB 2016 (30.5)	Casuarina	1948			
		pollen grains				