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1 **Late MIS 3 interstadial vegetation in coversands at Saint-Vincent-de-Paul, southwest France.**

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11

12 **ABSTRACT**

13 An archaeological survey carried out at Saint-Vincent-de-Paul, about 11 km northeast of Dax (Landes,
 14 southwest France) made it possible to investigate a new sequence of aeolian sands overlying a Middle
 15 Pleistocene alluvial terrace of the Adour River. A peat level with wood fragments was found between
 16 2.2 and 2.4 m depth at the bottom of a gully incised in, and then filled with aeolian sands. The peat,
 17 ¹⁴C dated on wood fragments to $26,731 \pm 101$ BP (31,136-30,844 cal BP) and $28,066 \pm 106$ BP (32,821-
 18 31,655 cal BP), can be correlated with the interstadial GI-5.1 (or possibly GI-5.2) of the Greenland Ice
 19 Core Chronology, i.e., in the latter part of Marine Isotope Stage (MIS) 3. The scarcity of Pleistocene
 20 peats in the region makes this discovery highly valuable for documenting the palaeoenvironments
 21 contemporary with the Middle Gravettian. The peat was cross-studied for pollen, non-pollen
 22 palynomorphs (NPPs), charred particles and woody macroremains. At the beginning of peat formation,
 23 a reed bed associated with willows developed locally in a treeless landscape. The surrounding area was
 24 covered by steppe vegetation, dominated by sagebrush. Later, the gully was colonized by birch trees,
 25 accompanied by *Salix*, *Myrica*, *Calluna* and *Erica cf. tetralix*. Finally, a sphagnum bog colonized by
 26 *Myrica*, *Calluna* and *Erica cf. tetralix* developed. The phase of woodland growth was synchronous with
 27 an increase in charred particles. This study together with the previous analyses made on regional MIS
 28 3 peats strongly suggests that patches of boreal forest repeatedly developed in favorable locations in
 29 the Landes region during the interstadials of the Last Glacial and lends support to the assumption of
 30 discontinuous aeolian activity in the coversand area.

31 **Keywords:** Late MIS 3 peat, Landes, southwest France, pollen, xylology

32 **RESUME**

33 VEGETATION INTERSTADIAIRE DE LA FIN DU SIM 3 DANS LES SABLES EOLIENS LANDAIS A SAINT-
 34 VINCENT-DE-PAUL, SUD-OUEST DE LA FRANCE.

35 Des sondages archéologiques réalisés à Saint-Vincent-de-Paul, à environ 11 km au nord-est de Dax
 36 (Landes, sud-ouest de la France) ont permis d'étudier une nouvelle séquence de sables éoliens
 37 recouvrant une terrasse alluviale de l'Adour attribuée au Pléistocène moyen. Un niveau de tourbe
 38 avec des fragments de bois a été trouvé entre 2,2 et 2,4 m de profondeur au fond d'une ravine

39 incisée dans les sables éoliens, puis remplie par de nouveaux apports sableux. La tourbe, datée au ^{14}C
 40 sur des fragments de bois de $26\ 731 \pm 101$ BP ($31\ 136\text{-}30\ 844$ cal. BP), et $28\ 066 \pm 106$ BP ($32\ 821\text{-}31$
 41 655 cal. BP) peut être corrélée avec l'interstadiaire GI-5.1 (ou peut-être GI-5.2) de la chronologie
 42 glaciaire du Groenland, c'est-à-dire dans la dernière partie du stade isotopique marin (SIM) 3. La
 43 rareté des tourbes pléistocènes dans la région rend cette découverte d'un grand intérêt pour
 44 documenter les paléoenvironnements contemporains du Gravettien moyen. La tourbe a fait l'objet
 45 d'études croisées pour le pollen, les microfossiles non polliniques (NPPs), les particules végétales
 46 carbonisées et les macrorestes ligneux. Au début de la formation de la tourbe, une roselière associée
 47 à des saules s'est développée localement dans un paysage dépourvu d'arbres. La zone environnante
 48 était couverte d'une végétation steppique, guidée par l'armoise. Plus tard, le ravin a été colonisé par
 49 des bouleaux, accompagnés de *Salix*, *Myrica*, *Calluna* et *Erica cf. tetralix*. Enfin, une tourbière à
 50 sphaignes peuplée de *Myrica*, *Calluna* et *Erica cf. tetralix* s'est développée. La densification des
 51 boisements locaux et l'accroissement consécutif de la biomasse se sont accompagnés de
 52 l'augmentation des particules carbonisées. Cette étude, ainsi que les analyses précédentes
 53 effectuées sur des tourbes régionales du SIM 3, suggèrent fortement que des parcelles de forêt
 54 boréale se sont développées de manière répétée dans les endroits favorables des Landes pendant les
 55 interstades du Dernier Glaciaire et soutient l'hypothèse d'une activité éolienne discontinue dans la
 56 zone des sables de couverture.

57 **Mots-clés :** Tourbe de la fin du SIM 3, Landes, sud-ouest de la France, pollen, xylologie.

58

59 **1. Introduction**

60 While pollen sequences documenting the Last Glacial (e.g., Woillard, 1978; De Beaulieu and Reille,
 61 1984, 1992a, 1992b; Guiter *et al.*, 2003; Fletcher *et al.*, 2010; Sirocko *et al.*, 2016; Duprat-Oualid *et*
 62 *al.*, 2017 ; Kern *et al.*, 2022) remain scarce in western Europe, they are especially rare and
 63 discontinuous in Aquitaine, southwest France (Paquereau and Schoeller, 1959; Paquereau, 1961;
 64 Diot, 1999). As a result, vegetation contemporary with the Last Glacial remains very poorly
 65 documented there. The closest available data are mainly from the Massif Central (Reille and De
 66 Beaulieu, 1988, 1990; De Beaulieu and Reille, 1992b), the Pyrenees (Olfield, 1968; Andrieu-Ponel *et*
 67 *al.*, 1988; Jalut *et al.*, 1992; Jalut *et al.*, 1998; González-Sampériz *et al.*, 2006) and the Loire and
 68 Garonne basins recorded in the marine sediments of Bay of Biscay (Sanchez-Goñi *et al.*, 2008,
 69 Fletcher *et al.*, 2010). These records provide only an imperfect picture of lowland vegetation in
 70 Aquitaine, either because of the altitude (Massif Central and Pyrenees) or because the pollen inputs
 71 come from a wide geographical area (Bay of Biscay).

72 The Pleistocene peat levels discovered during preventive archaeological work offer the opportunity
 73 to document in detail, albeit during small time windows, the glacial palaeoenvironments of the
 74 region. Although discontinuous, these records provide information on the vegetation mosaic
 75 characterizing a region and may allow the identification of potential refuge areas for temperate
 76 trees. Due to poor drainage, the Landes plateau has yielded numerous occurrences of peat
 77 intercalated in the aeolian 'Sable des Landes' Formation. Some of these levels have been the subject
 78 of pollen analysis and numerical dating, which made it possible to relate them to interstadials of the
 79 first half of Marine Isotopic Stage (MIS) 3 (i.e., 55-40 ka; Cestas-Les Pins de Jarry, Bertran *et al.*, 2009,
 80 2020; Barp-Pot aux Pins, Sitzia *et al.*, 2015) (Fig. 1) or to the Middle Pleistocene (Hourtin, Sitzia *et al.*,

81 2015, north of the coversands not shown in Fig. 1). The discovery of a peat level about 15 cm thick at
 82 Saint-Vincent-de-Paul in the southern margin of the ‘Sable des Landes’ provides an opportunity to
 83 document another late MIS 3 interstadial and is the focus of this paper. The presence of wood
 84 fragments gives this peat an exceptional character.

85 **2. Geomorphological setting**

86 The study site is located in the Adour valley on the right bank, approximately 11 km northeast of Dax
 87 (southwest France) (Fig. 1). The 1:250,000 digital geologic map of Aquitaine (Bourbon, 2019)
 88 indicates that the bedrock is a high alluvial terrace of the Adour River (Fv, Lower to Middle
 89 Pleistocene). Aeolian sands several meters thick, which are an integral part of the ‘Sable des Landes’
 90 Formation (Sitzia *et al.*, 2017), overlie the alluvial deposits. The geotechnical drillings carried out as
 91 part of the development of the site did not reach the alluvium at a depth of 3 m. According to
 92 borehole BSS002EAWU located approximately 3.3 km to the southwest and stored in the Banque du
 93 Sous-Sol of the Bureau des Recherches Géologiques et Minières (<http://infoterre.brgm.fr/>), the sands
 94 would reach 5.5 m in thickness in the area. The aeolian cover has yielded two ages by Optically
 95 Stimulated Luminescence (OSL), one related to the Middle Pleistocene (~160 ka), the other to the
 96 Upper Pleistocene (~26 ka) in the Pontonx quarry 3 km to the southeast (Bertran *et al.*, 2020).

97 The topography visible from the 5 m DEM of the Institut Géographique National (RGEALTI® 5 m)
 98 shows that the area is covered by ridges 1 to 2 m high and oriented WNW-ESE, which correspond to
 99 the arms of elongated parabolic dunes (Fig. 2A). These degraded dunes have been dated to the
 100 Younger Dryas (~12 ka) at various points in the Landes, notably at Castets and Saint-Geours-de-
 101 Maremne (Bertran *et al.*, 2020), and mark the ultimate evolution of Pleistocene coversands. The
 102 relief was then dissected by a network of gullies connected to tributaries of the Adour River.

103 **3. Methods**

104 A peat block was collected in trench CM1-T10 (Fig. 3) for a multidisciplinary study including pollen
 105 and spores, non-pollen palynomorphs (NPPs), charred particles and woody macroremains. For the
 106 palynological study (pollen and NPPs), all samples were processed following the standard method
 107 developed by Faegri and Iversen (1975), which consists of chemical treatments with 10% NaOH, 37%
 108 HCl and 48% HF. Then a separation with a heavy liquid ($ZnCl_2$, density=2) has been carried out. No
 109 acetolysis was made in order to recognize any contamination by exogenous pollen stocks. Finally, the
 110 rest of the sediment was sieved using 10 μm and 200 μm mesh sieves.

111 The preparation involved a volume of 4 cm^3 of sediment. To estimate the richness in palynomorphs,
 112 the absolute pollen concentrations (number of pollen grains and spores per cm^3 of sediment) were
 113 calculated according to the method of Stockmarr (1971). Palynomorphs were mounted in a mobile
 114 medium (glycerine) and identified at 500x and 1000x magnification. Identification of spores and
 115 pollen material was made using determination keys (Faegri and Iversen, 1975; Moore *et al.*, 1991;
 116 Beug, 2004), photographic atlases (Reille, 1992, 1995, 1998) and the CReAAH pollen reference
 117 collection. The names of taxa followed by “type” refer to nomenclature of “The Northwest European
 118 pollen flora” for Asteraceae, Gentianaceae, Polygonaceae, Sparganiaceae families, and to
 119 nomenclature of Moore, Webb and Collinson for Plumbaginaceae family (Punt, 1975; Punt and
 120 Nienhuis, 1976; van Leeuwen *et al.*, 1988; Moore *et al.*, 1991; Punt and Hoen, 2009). Determination
 121 of non-pollen microfossils was carried out with the atlas published in 1998 based on the work of van
 122 Geel (van Hoeve and Hendrikse, 1998), supplemented by the synthesis of Miola (2012) and the

123 theses of Jouffroy-Bapicot (2010) and Cugny (2011). The diagram was built on a basic sum from
 124 which sedges (Cyperaceae), spores and indeterminate grains were excluded. NPPs are expressed as a
 125 percentage of the pollen sum. Microscopic charred particles (10-200 µm) observed in pollen sample
 126 residues were also counted on five lines at 500× magnification on each pollen slides. The
 127 concentration of charred particles in the sediment was obtained using an adaptation of the
 128 volumetric method (Cour, 1974). This method was chosen because the morphological features of
 129 charred particles differ from *Lycopodium* spores, which are spherical. The concentration of charred
 130 particles is expressed as the number of charred particles per cm³ of fresh sediment.

131 For the xylological study, macroremains recovery involved water sieving of the sediment on a column
 132 of 250 µm, 500 µm and 2 mm sieves. The volume of sieved sediment ranged from 100 to 400 cm³,
 133 depending on the available sampling volume (Table 1). Wood analysis was performed on the remains
 134 larger than 2 mm. A number of 40 to 50 individuals were analyzed in the majority of samples. The
 135 minimum number of fragments to be analyzed per sample was determined from the cumulative
 136 frequencies of taxa as a function of the number of individuals, the optimal number being reached
 137 when the frequencies of taxa are stabilized. Prior to taxonomic identification, macroscopic analysis of
 138 the wood fragments was carried out under a binocular microscope, i.e. length, width, thickness,
 139 presence of bark, pith, and stem starts associated or not with leaf scars, degree of ring curvature
 140 according to three classes (strong, intermediate and weak). The taxonomic identification was carried
 141 out with a transmitted light microscope, (x200 and x500) using xylological atlases (Schweingruber,
 142 1978, 1990). For the genus *Erica*, the arrangement of leaf scars and their number, observed under
 143 binocular microscope, was a criterion for the specific determination, using the flora of Rameau *et al.*
 144 (1989). The taxonomic identification of the root wood could not be realized as root wood does not
 145 have a uniform anatomical structure and cannot be identified except for rare cases (Schweingruber,
 146 1978).

147 **4. Results**

148 **4.1. Stratigraphy**

149 The trenches carried out during the archaeological survey allowed us to observe the sedimentary
 150 formations over a thickness of 3 m. The stratigraphy is relatively homogeneous on the land parcel. It
 151 is schematically composed of the following units, from the top to the bottom (Fig. 3):

152 (1) A Holocene podzol, about 1 m thick. The upper horizons have been mixed by agricultural work
 153 (black Ap horizon, 0.4 to 0.5 m thick). The underlying horizons (Bh, Bs) are poorly cemented by Fe
 154 oxides.

155 (2) Light brown sands becoming light grey at the base, with dominant subhorizontal bedding. Levels
 156 with deformed or slightly oblique bedding are present. This unit corresponds to aeolian sandsheet
 157 deposits (coversands).

158 Two trenches (CM1-T10 and CM2-T9) revealed a peaty level beneath the sandy cover between 2.2
 159 and 2.4 m deep (Fig. 2B). This level was described in detail and sampled in trench CM1-T10 (Fig. 3). It
 160 shows at the base (Fig. 4) alternating peaty and sandy beds (2 cm thick); then brown peat with
 161 abundant plant debris (4 cm) (Fig. 5), humified black peat with plant debris (3 cm), and finally, at the
 162 top, alternating sand and black peaty beds (5 cm). The base of the sequence consists of coarse
 163 subhorizontally bedded brown sands of alluvial origin.

164 The limited spatial extent of the peat suggests that it formed in a gully incised in aeolian deposits.
 165 Examination of the 5 m DEM (Fig. 2) indicates that the area is intersected by several small tributaries
 166 of the Adour River, notably the Pont-de-Paul creek that flows 400 m west of the parcel. These
 167 shallowly incised tributaries (2 to 3 m) form a roughly dendritic network, converging on a single,
 168 deeper talweg at the edge of the present Adour plain. The hypothesis of an ancient gully incised into
 169 the sands during an interstadial period, then largely filled by aeolian deposits during the Last Glacial
 170 Maximum can therefore be proposed to account for the localized presence of peat accumulation. In
 171 trench CM1-T10, two generations of aeolian sandy fill separated by a small incipient podzol can be
 172 distinguished. The more recent generation of sand is likely related to the emplacement of the
 173 Younger Dryas dunes.

174 **4.2. Chronology**

175 Two samples were dated by AMS-¹⁴C at the CIRAM laboratory (Martillac). A first large wood fragment
 176 (4 cm long, 1 cm wide) was taken from the median brown peat level. The age obtained (CIRAM-AV85)
 177 is $26,731 \pm 101$ BP, i.e. 31,136 to 30,844 cal BP (2, Intcal20). Another smaller willow fragment from
 178 the basal sand and peat beds was also dated. The age (CIRAM-4395) is $28,066 \pm 106$ BP, i.e. 32,821 to
 179 31,655 cal BP. The peat can therefore be correlated with one of the GI-5.1 or GI-5.2 interstadials of
 180 the Greenland ice core record (Rasmussen *et al.*, 2014), i.e., in the latter part of MIS 3. Insofar as the
 181 study of pollen assemblages (see *infra*) shows that the peat recorded only one phase of climate
 182 amelioration and no discontinuity in sedimentation is identifiable within the level, a correlation with
 183 GI-5.1 is favored here. According to this hypothesis, the small willow fragment taken from the base
 184 of the peat could correspond to slightly older material reworked at the bottom of the gully.

185 **4.3. Palynology**

186 4.3.1. Pollen and NPP

187 Pollen sedimentation conditions appear favorable, ensuring reliable and significant data from a
 188 palaeoecological point of view (Reille, 1990; Leroyer, 1997), as shown by high counts and absolute
 189 concentrations, good preservation of pollen stocks, and satisfactory taxonomic diversity (Table 2, fig.
 190 6). This has made it possible to draw up a floristic list of 44 taxa, to which can be added 26 NPPs (fig.
 191 6). The pollen assemblages (Table 3, fig. 6) allows us to identify three local pollen zones (LPZ), with
 192 the first two composed of only one sample each.

193 Characterized by the scarcity of tree pollen, the first local pollen zone (LPZ 1, sample at 15 cm from
 194 the top of the peat) covers the base of the organic level (peaty sand). The environment appears very
 195 open with the local presence of a reed bed-megaphorbia or a sedge meadow (Poaceae, Cyperaceae,
 196 Rosaceae, Ranunculaceae, *Thalictrum*), formations that thrives on waterlogged soils with little
 197 oxygen (Fare *et al.*, 2001). The absence of aquatics indicates that there was no perennial water body
 198 (Dutartre and Rebillard, 2015). In addition to pollen assemblages, the presence of this vegetation is
 199 corroborated by the predominance of *Clasterosporium carincinum* (HdV-126) and *Byssothecium*
 200 *circinans* (HdV-16), microfossils that are respectively associated with *Carex* (Cyperaceae) and
 201 Poaceae but are absent from very wet areas (van Geel, 1976; van Geel and Middeldorp, 1988; van
 202 Geel and Aptroot, 2006; Yeloff *et al.*, 2007). A few rare trees were present, mainly willows (*Salix*)
 203 whose occurrence *in situ* is confirmed by woody debris (Fig. 7) but also some rare hygrophilous
 204 shrubs (*Myrica*, Ericaceae, *Calluna*). In the surroundings, the drier areas hosted a low steppe
 205 vegetation, dominated by sagebrush (*Artemisia*) accompanied by other heliophilous herbaceous

206 species requiring well-drained soils. In light of the low pollen count (12%), the local presence of pine
 207 appears unlikely (Huntley and Birks, 1983). Its record is more likely to be the result of distant
 208 contributions in an open environment because of high pollen production and its capacity for long-
 209 distance dispersal (Heim, 1970). Similarly, the sporadic nature and low occurrence (< 0.5 %) of
 210 temperate hardwoods (*Fagus*, *Carpinus*, *Corylus*) also argue for distant contributions even though the
 211 maintenance of beech in southwestern France during the Late Pleistocene is plausible based on
 212 palaeogenetic data (de Lafontaine *et al.*, 2014). Although current pollen studies show that
 213 percentages of *Fagus* below 2% may indicate the local presence of isolated trees (Huntley and Birks
 214 1983), the persistence of beech during MIS 3 near St. Vincent de Paul cannot be assumed from our
 215 data, since it is only recorded in the basal peat sample, which also provides the greatest amount of
 216 pine pollen from distant sources.

217 The base of the brown peat (LPZ 2, sample at 10 cm) shows a profound local environmental change.
 218 The wetland was colonized by birch trees (*Betula*), accompanied by willows and rarer Ericaceae and
 219 marsh heather (*Calluna*). The development of trees created a filter to distant inputs of pine pollen,
 220 which supports the idea that pine was not established in the immediate vicinity. Similarly, the
 221 perception of steppe vegetation (*Artemisia*, Asteraceae, Chenopodiaceae) decreased, but it is not
 222 possible to assess whether it really declined. Although more constrained, a reed bed remained in the
 223 site. It evolved somewhat with the appearance of meadowsweet (*Filipendula*) and bur-reed
 224 (*Sparganium emersum* type) at the side of Cyperaceae and Poaceae. Bur-reed indicates, in the
 225 absence of true aquatic plants, soils that are often wet but without perennial water (Aniotsbehere,
 226 1999). This interpretation is corroborated by the microfossils HdV-128 and HdV-187d, the latter
 227 infatuated with *Phragmites* and *Carex*, both of which indicate shallow, eutrophic water (Pals *et al.*,
 228 1980; van Geel *et al.*, 1983). However, the appearance of copepod spermatophores could suggest the
 229 at least temporary occurrence of open water (van Geel, 1978). These small crustaceans can also be
 230 related to the establishment of sphagnum mosses (Mighall *et al.*, 2006), reported by their spores and
 231 the presence of *Tilletia sphagni* (HdV-27), which is dependent on them (van Geel, 1978).

232 The third pollen (LPZ 3, samples at 7.5, 5 and 2 cm) zone shows the development of a sphagnum bog,
 233 which is attested, in addition to the rise of *Sphagnum* spores, by *Tilletia sphagni* (HdV-27). Other
 234 microfossils typical of sphagnum bogs are present, e.g. *Assulina muscorum* (HdV-32a) and *Callidina*
 235 *angusticollis* (HdV-37). The former has broad ecological requirements, but has a high relative
 236 abundance in drier stages of bogs (Payne *et al.*, 2012). The evolution of vegetation appears to have
 237 been gradual with, initially (LPZ 3a), the retreat of birch trees to the benefit of willows, with also
 238 some Ericaceae and bog myrtle (*Myrica*). Then (LPZ 3b and c), the site was colonized by *Myrica*, a
 239 shrub specific to acidic and humid soils, especially peat bogs (Bournerias, 1979). Birch trees (LPZ 3b)
 240 and then willows (LPZ 3c) disappeared almost completely, but a few Ericaceae and *Calluna* remained.
 241 The development of a denser local vegetation led to the non-perception of pines. On the other hand,
 242 the persistence of a reed bed is testified by pollen of Poaceae and Cyperaceae but also by the
 243 presence of microfossils that are attached to them such as *Clasterosporium caricum* (HdV-126) and
 244 sometimes *Glomus cf. fasciculatum* (HdV-207) (van Geel *et al.*, 1989, 2003).

245 4.3.2. Palynofacies and charred particles

246 The palynofacies consists mainly of fresh non-woody plant debris, i.e. rootlets of Cyperaceae, tissues
 247 of aerial parts of Monocotyledons, indeterminate herbaceous tissues, and more rarely Bryophyte and
 248 *Sphagnum* tissues. Charred particles are rarer, especially in the two lower samples (15 and 10 cm),

249 where their concentrations are less than 1500 debris/cm³. However, the charred particles increase
 250 abruptly from 7.5 cm where they reach their maximum concentration (117,365/cm³) and then
 251 decrease to around 85,000/cm³. They include remains of woody and herbaceous nature, of which
 252 some correspond to tissues of aerial parts of Monocotyledons. Their size is small, mostly less than
 253 100 µm, but some herbaceous remains can reach up to 420 µm in length and 140 µm in width. The
 254 increase in the concentration of charred particles is synchronous with the densification of local
 255 woodlands, corroborating their association with a higher biomass characteristic of wetter
 256 interstadials (Daniau *et al.*, 2009, 2010).

257 **4.4. Macroremains**

258 **4.4.1. Representativeness of data**

259 While a minimum of 40-50 individuals was analyzed in most samples, the optimum number was not
 260 systematically reached (i.e. for sample 0-4 cm). Wood fragment concentrations increased from the
 261 base to the top (Table 1), reaching up to 243 fragments per 100 cm³ of sediment in the top sample.
 262 Only five taxa were identified, i.e. birch (*Betula pendula/pubescens*), willow (*Salix*), marsh heather
 263 (*Erica cf. tetralix*), heather (*Erica*) and Ericaceae (Table 3). The high rate of indeterminate debris results
 264 from the abundance of root fragments, which do not allow identification except occasionally for birch
 265 (Schweingruber, 1978, 1990).

266 Preservation of woody material is variable across layers and taxa. For example, *Erica cf. tetralix* and
 267 Ericaceae fragments are generally well preserved in the four samples where they were identified (Table
 268 4). The condition of willow fragments is variable, but mostly quite poor, while birch fragments are
 269 poorly preserved in the three lower samples (7.5-10, 10-15 and 15-17 cm). The willow and birch
 270 fragments have a compressed cross section. The fragments are small with a length ranging from 0.06
 271 to 5.1 cm (mean 1.3 cm) and a width from 0.05 to 2 cm (mean 0.25 cm). In fact, the different species
 272 show slightly different sizes (Table 4). When the cross-section is complete, the diameter could be
 273 calculated. It never exceed 0.5 cm and is mostly smaller than 0.25 cm. The largest diameters
 274 correspond mainly to indeterminate remains, to which are added one fragment of birch and one of
 275 heather. The woody material is, therefore, of small size. When identifiable, the curvature of the rings
 276 is always strong, except for one individual with intermediate curvature, pointing to small branches.

277 **4.4.2. Evolution of the macroremain assemblages**

278 The general diagram (Fig. 8) shows the predominance of willow (*Salix*) ahead of birch (*Betula*) in the
 279 three lower samples (7.5 to 17 cm), followed by indeterminate remains (4 and 5 cm) and, finally,
 280 heather (*Erica*) at the top. *Salix* has its maximum representation at 10-15 cm then decreases somewhat
 281 (7.5-10 cm) and then disappears completely from 7.5 cm. Its evolution as woody fragments roughly
 282 follows that documented by the pollen assemblages, which shows the retreat of willow in the upper
 283 levels (Fig. 6). However, the relative abundance of *Betula* in the basal peaty sand beds raises questions,
 284 as it is not attested in pollen assemblages. Since this absence cannot be attributed to a deficit in pollen
 285 production and dispersal (Heim, 1970), it is likely that the birch fragments correspond to roots, which
 286 are identifiable for this species (Schweingruber, 1978, 1990). They would, therefore, derive from the
 287 birch stand that later established in the site.

288 *Erica* appears in the 7.5-10 cm sample. It is the only taxon recorded in the two overlying samples (0-5
 289 cm) that yielded mainly indeterminate woody debris (>90%). *Erica* becomes largely predominant at

290 the top, especially if we accept that the Ericaceae fragments refer to this genus. The woody debris thus
 291 documents an environment somewhat different from that depicted by palynology, as it includes no
 292 remains of *Myrica* but almost exclusively of *Erica cf. tetralix*. This difference is likely due to two factors,
 293 the pollination mode of Ericaceae, which are always under-represented in pollen assemblages (Heim,
 294 1970) even if in areas where *Erica sp* are locally abundant, pollen values >25% can be frequent (Huntley
 295 and Birks, 1983), and the very local information provided by the macroremains. It can be assumed that
 296 only marsh heather was growing at the sampled site, while bog myrtle was located a little further away,
 297 but probably in the immediate vicinity as *Myrica* usually has a low pollen representation, even where
 298 it is abundant and growing luxuriantly (Huntley and Birks, 1983).

299 **5. Discussion**

300 As mentioned above, continental records documenting the Last Glacial vegetation remain scarce in
 301 Western Europe. Vegetation is mainly documented by a few long continental sequences geographically
 302 distant from southwest France, supplemented by fragmentary records from northwest Europe coming
 303 from organic levels intercalated in glacial deposits (Fletcher *et al.*, 2010). The pollen analyses carried
 304 out on the peaty levels discovered during preventive archaeological work in the Landes region are
 305 closely related to the latter. Although discontinuous, these records have the advantage of being close
 306 to each other and of having been deposited in a similar geomorphological context, i.e. Pleistocene
 307 coversands. Also dated to MIS 3, they allow for comparison with the Saint-Vincent-de-Paul record.

308 Two sites near the town of Cestas (Fig. 1) have been analyzed and dated. In both sites, the peat levels
 309 are cryoturbated and have developed in interdune depressions in connection with the rise of the
 310 groundwater table. At Barp-Pot aux Pins, the peat was OSL-dated between 50 ± 3 ka and 55 ± 3 ka and
 311 correlated with GI-14 (Sitzia *et al.*, 2012, 2015). Pollen assemblages are dominated by *Pinus*, ahead of
 312 Ericaceae, Poaceae and *Myrica*. The internal evolution of the peat level attests to a rise in woody
 313 species (*Pinus*, *Myrica*, Ericaceae) at the expense of Poaceae. The peat sample analyzed at Cestas-Les
 314 Pins de Jarry was ^{14}C -dated on bulk sediment to $23,550 \pm 160$ BP (27,330-28,020 cal BP) and $22,390 \pm$
 315 120 BP (26,140-26,670 cal BP) (Bertran *et al.*, 2009). Nevertheless, an OSL date on aeolian sand made
 316 10 cm above the peat subsequently provided an age of 40.2 ± 2.7 ka (Bertran *et al.*, 2020), which
 317 strongly suggests that the ^{14}C ages were rejuvenated probably due to illuvial organic matter derived
 318 from the superficial podzol. Therefore, this level could correspond to a relatively old MIS 3 interstadial
 319 (GI-9 to GI-14). The single pollen assemblage obtained shows a pattern quite similar to the last phase
 320 seen at Saint-Vincent-de-Paul, and is dominated by Poaceae, *Myrica*, Ericaceae, *Pinus* and *Betula*. If
 321 the distribution of the different taxa is not completely similar (Ericaceae, *Pinus*, *Betula* and Poaceae
 322 are better perceived there than *Myrica*), it nevertheless refers to the same type of environment.

323 Overall, the MIS 3 peat levels described in the Landes show the development of localized woodlands
 324 during various interstadials, mainly composed of cold-tolerant species typical of the boreal forest
 325 (*Salix*, *Betula*) associated with hygrophilous plants such as *Myrica* and Ericaceae. *Pinus*, more or less
 326 abundantly represented in pollen assemblages, is probably present only at great distance from the
 327 sites (cf. Cheddadi *et al.*, 2006). It is only known in the region as macroremains (charcoal) during the
 328 Alleröd interstadial in palaeosols preserved beneath Younger Dryas dunes (Bertran *et al.*, 2009; Sitzia
 329 *et al.*, 2015). Better representation of *Pinus* during GI-14 than GI-5, which suggests greater regional
 330 abundance, is in line with the Bay of Biscay record (Sanchez-Goñi *et al.*, 2008; Fletcher *et al.*, 2010).
 331 The absence of thermophilous tree taxa such as *Quercus* and *Fagus* is also noteworthy. The single
 332 occurrence of a *Fagus* pollen grain in the basal peat sample cannot lend support to the debate about

333 the presence of beech stands in the Landes during MIS 3 proposed from genetic studies (de Lafontaine
 334 et al., 2014).

335 The interstadial vegetation dynamics recorded in southwest France do not differ significantly from
 336 those seen elsewhere in lowland western Europe, with the development of a few trees (*Salix*, *Betula*,
 337 *Juniperus*) in the context of an open vegetation dominated by Poaceae and Cyperaceae (Guter et al.
 338 2003; Fletcher et al., 2010; Kern et al., 2022). Comparison with the long French continental records (La
 339 Grande Pile, Les Echets, Velay maars, Sost, Barbazan, Lourdes) is not always easy because of the low
 340 resolution and poor chronological control that limit correlations to the various MIS 3 interstadials. The
 341 Velay sequences (Lac du Bouchet, Ribains), geographically closest but located at much higher
 342 elevation, reflect punctual extensions of *Pinus* and *Picea* into an open *Betula* and *Salix* boreal forest
 343 during the Upper Pleniglacial, which were attributed to long-distance transport or to older pollen
 344 stocks reworked by erosion (Reille and de Beaulieu 1988, 1990; de Beaulieu and Reille, 1992a). In the
 345 French Pyrenees, the Barbazan, Sost and Lourdes pollen sequences (Andrieu-Ponel et al., 1988; Jalut
 346 et al., 1992; Jalut et al., 1998) attest to the occurrence of *Picea* up to ca. 26 ka cal BP together with few
 347 mesothermophilous trees (mainly *Quercus* and *Corylus*). By contrast, the El Portelet sequence in the
 348 drier Spanish Pyrenees only records the presence of *Pinus* and *Juniperus*, while the few pollen grains
 349 from mesothermophilous trees were thought to reflect long-distance transport (Gonzalez-Sampeniz et
 350 al., 2006). At Bergsee (southwest Germany), sharp increases in *Betula* and *Pinus* are recorded,
 351 accompanied by *Juniperus* that evoke the development of open boreal forests (Duprat-Oualid et al.,
 352 2017). Pine played a notable role there, which has already been observed at La Grande Pile during the
 353 "Grand Bois Complex" correlated to the end of MIS 3 (de Beaulieu and Reille, 1992a). Recent reanalysis
 354 of the Füramoos sequence (southern Germany) allows better correlation with the late MIS 3
 355 interstadials, which were marked by rapid tree extensions including *Betula*, *Pinus* and sporadically
 356 *Picea*. However, vegetation remained dominated by Cyperaceae and Poaceae grasslands and
 357 sagebrush steppe (Kern et al., 2022).

358 Finally, late MIS 3 pollen assemblages from core MD04-2845, located offshore but not far from the
 359 Landes, indicate a development of deciduous oaks (*Quercus* deciduous type) together with boreal
 360 forest (*Pinus*, *Abies*, *Picea*, *Betula*) during the interstadials (Sanchez-Goñi et al., 2008; Fletcher et al.,
 361 2010), in sharp contrast to the Landes records. This obviously reflects a regional signal that includes
 362 refuge areas in the valleys of the Pyrenees where thermophilous trees were able to survive during the
 363 Glacial, and possibly in the Massif Central (although thermophilous trees are not recorded in the
 364 available pollen sequences). These also clearly demonstrate the specific character of the Landes
 365 vegetation, largely related to the acidic nature of the soils developed on aeolian sand.

366 The development of peat in the Landes region, although rarely preserved presumably due to deflation,
 367 strongly suggests that the aeolian dynamics was only intermittently active during the Last Glacial, or
 368 at least that sand transport was spatially restricted to a few favorable areas during the interstadials.
 369 Concurrently, overland flow and gully incision were active. The gullies, later plugged by new inputs of
 370 aeolian sand, provide favorable contexts for the preservation of peat. The discovery of peat levels in a
 371 similar context at Saint-Paul-les-Dax, dated to $28,230 \pm 140$ BP ($32,480 - 31,530$ cal BP), i.e. GI-5.2
 372 (Béague, 2014), and at Pontonx, dated to $32,410 \pm 450$ BP ($37,800 - 35,300$ cal BP), i.e. GI-8 (Bertran,
 373 unpublished report), allows us to anticipate the possibility that other interstadials may be documented
 374 in the future thanks to preventive archaeological work in the region.

375 **7. Conclusion**

376 The peat level discovered during preventive archaeological work at Saint-Vincent-de-Paul in the Landes
 377 region (southwestern France) documents the vegetation of interstadial GI-5.1, ca. 31 ka cal BP (or
 378 possibly GI-5.2). This level was preserved at the bottom of a gully incised in aeolian sands. During the
 379 early peat formation, a reed bed associated with willows developed in an open sagebrush steppe
 380 landscape. The gully was then colonized by birch trees, accompanied by willows, *Myrica*, *Calluna* and
 381 *Erica cf. tetralix*. *Sphagnum* peat finally formed. Comparison with other analyses of MIS 3 peat in the
 382 region shows that patches of boreal forest developed locally repeatedly during the interstadials. It
 383 shows as well that aeolian activity was discontinuous and probably restricted to spatially limited
 384 favorable areas during these periods. The sporadic discovery of other peat levels in similar contexts
 385 also dated to MIS 3 suggests the possibility of documenting other interstadials in the future if careful
 386 attention is paid to these finds.

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- 578

579 **Figure captions**

580 Figure 1: Map of aeolian deposits in southwest France, from Bertran et al. (2020) and location of the
 581 sites described in the text.

582 *Figure 1: Carte des dépôts éoliens du sud-ouest de la France, d'après Bertran et al. (2020) et localisation*
 583 *des sites décrits dans le texte.*

584 Figure 2: Topographic maps of the study area. Relief is from RGEALTI® 5 m (Institut Géographique
 585 National). A – General map; B – Location of trenches. The black dots correspond to the trenches that
 586 have yielded the studied peat level.

587 *Figure 2 : Cartes topographiques du secteur étudié. Le relief est tiré du RGEALTI® 5 m de l'IGN. A – Carte*
 588 *générale ; B – Localisation des tranchées. Les points noirs correspondent aux tranchées qui ont livré le*
 589 *niveau tourbeux analysé.*

590 Figure 3: Schematic stratigraphy. The horizon labels in CM1-T10 trench are from *Food and Agriculture*
 591 *Organization (FAO, 2006).*

592 *Figure 3 : Stratigraphie schématique. La nomenclature des horizons pour la tranchée CM1-T10 est celle*
 593 *de la Food and Agriculture Organization (FAO, 2006).*

594 Figure 4: View of the studied peat sample.

595 *Figure 4 : Photographie du bloc de tourbe analysé.*

- 596 Figure 5: Close up view of the plant debris.
- 597 *Figure 5 : Macrophotographies des débris végétaux.*
- 598 Figure 6: Pollen diagram (Cyperaceae, spores and indeterminates are excluded from the sum). A -
599 Pollen and spores; B - Major pollen types, non-pollen palynomorphs and charred particle
600 concentrations.
- 601 *Figure 6 : Diagramme pollinique (Cyperaceae, spores et indéterminés sont exclus de la somme). A -*
602 *pollen et spores ; B - principaux types polliniques, microfossiles non polliniques et concentrations en*
603 *particules carbonisées.*
- 604 Figure 7: Woody fragments. A – *Salix* (5-7.5 cm); B – *Erica cf. tetralix* (0-4 cm).
- 605 *Figure 7 : Fragments ligneux. A – Salix (5-7,5 cm) ; B – Erica cf. tetralix (0-4 cm).*
- 606 Figure 8: Xylological diagram.
- 607 *Figure 8 : Diagramme xylologique.*
- 608
- 609 **Table captions**
- 610 Table 1: Concentration in woody fragments and number of fragments analyzed.
- 611 *Tableau 1 : Concentrations en macrorestes ligneux et nombre de fragments analysés.*
- 612 Table 2: Sporo-pollen concentrations, number of morphotypes and preservation of pollen and spores.
- 613 *Tableau 2 : Concentrations sporo-polliniques, nombre de morphotypes et préservation du pollen et des*
614 *spores.*
- 615 Table 3: Description of pollen, spore and non-pollen palynomorph assemblages, charred particles
616 concentration and woody remains.
- 617 *Tableau 3 : Description des assemblages de pollen et spores, microfossiles non-polliniques,*
618 *concentration en particules carbonisées et macrorestes ligneux.*
- 619 Table 4: Preservation, size and curvature of woody remains.
- 620 *Tableau 4 : Conservation, taille et courbure des macrorestes ligneux.*
- 621
- 622

■ Lateglacial-Holocene dunes
■ Pleistocene coversands

■ Bordeaux

Arcachon

Atlantic
Ocean

Cestas-Les Pins de Jarry

Cestas-Pot aux Pins

Garonne

Leyre

Roquefort

Castets

St-Vincent-de-Paul

Mont-de-Marsan

Adour

Camiade

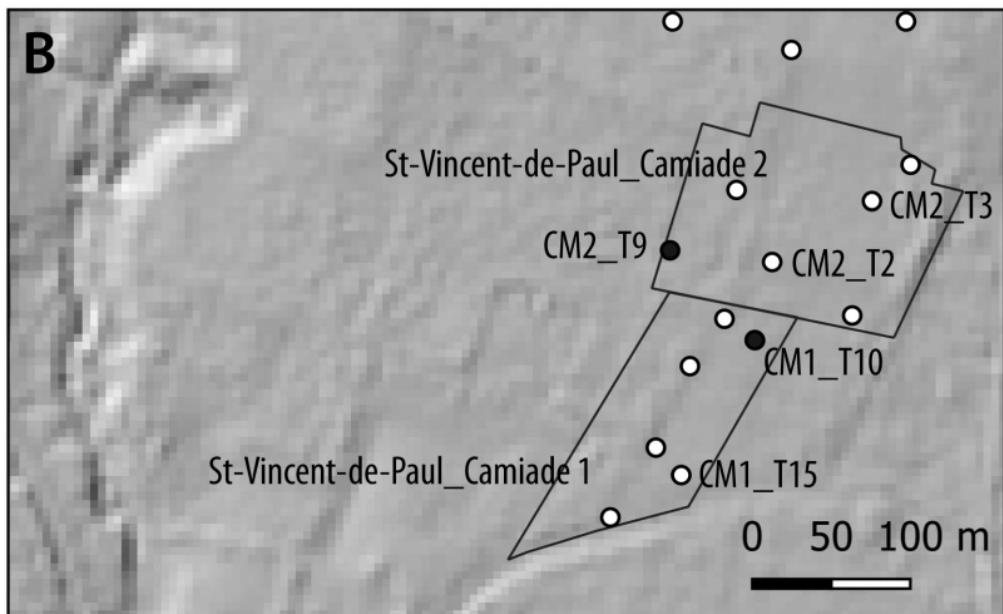
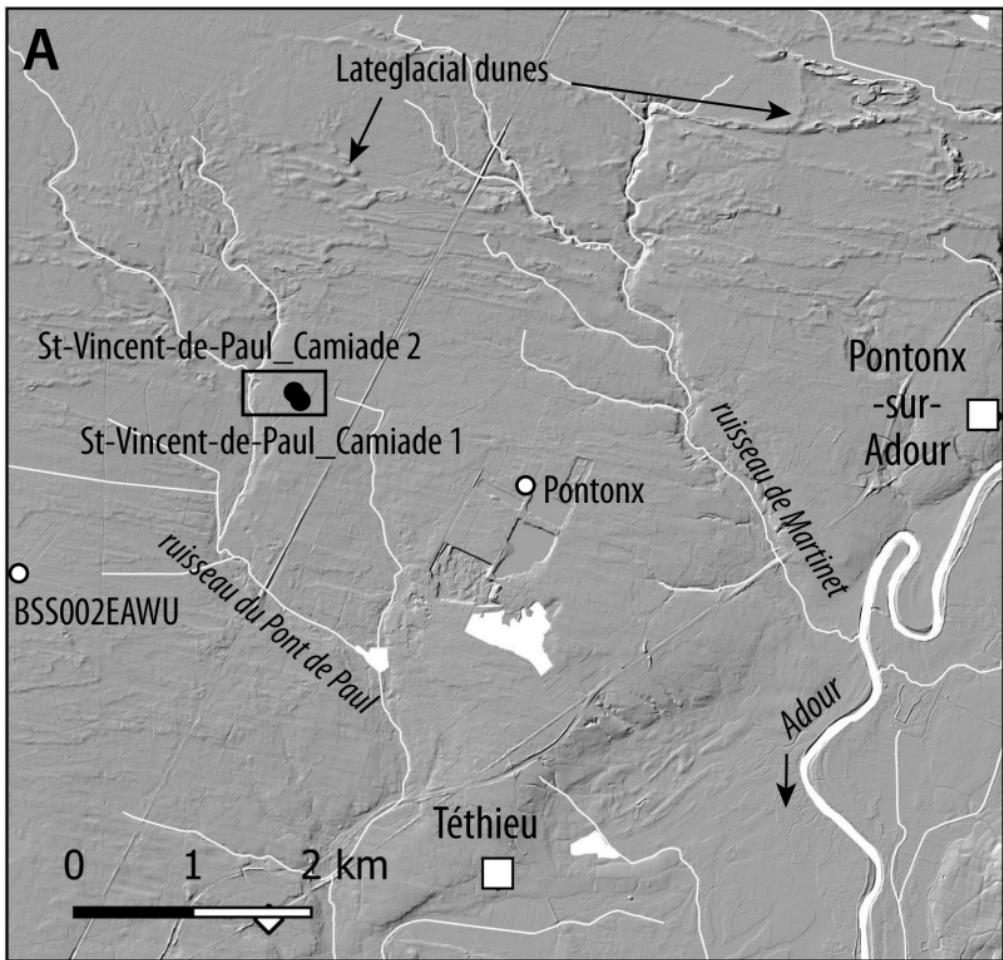
St-Paul-les-Dax

Dax

Saint-Geours-
de-Maremne

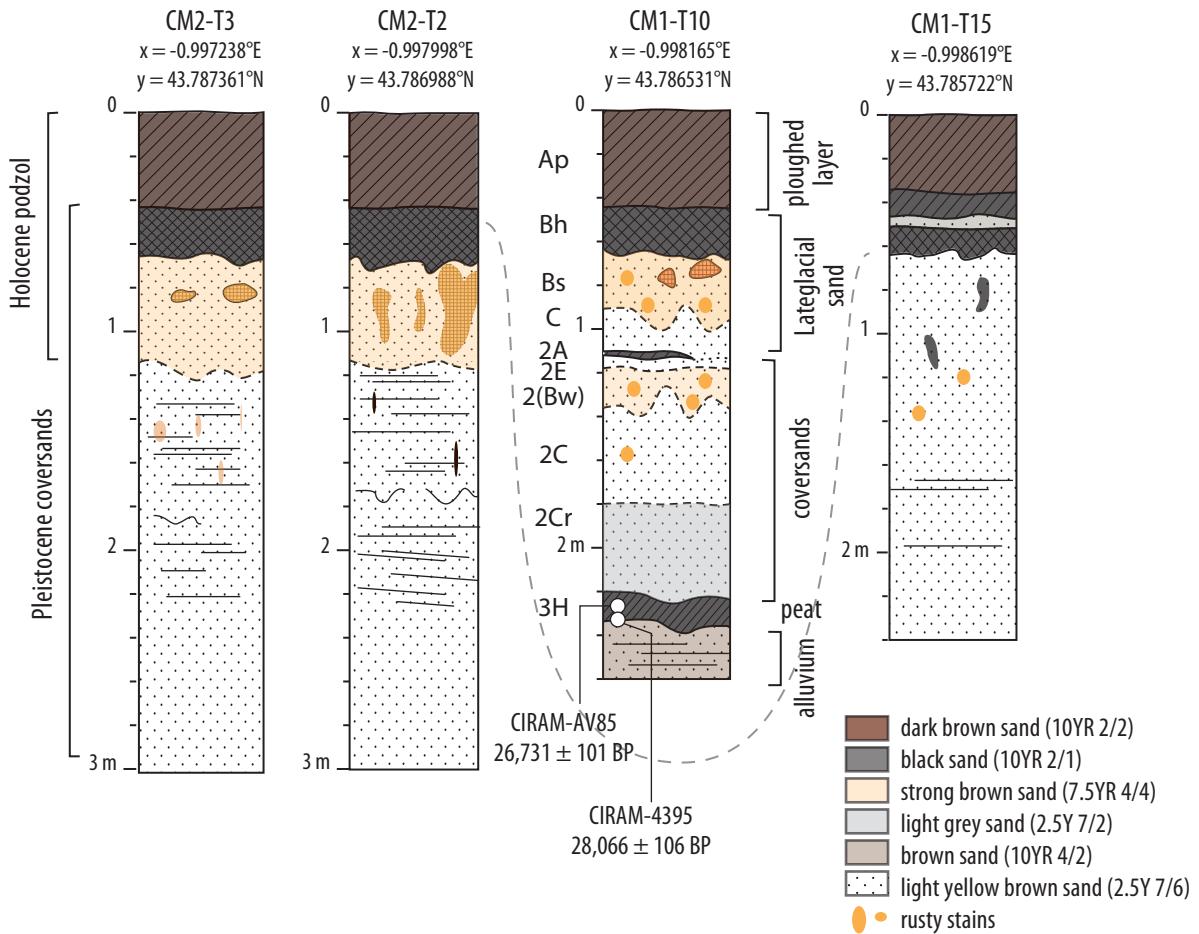


0 10 20 km



N

S





Alternating sand and
black peat beds

Black humified peat
with wood fragments

Brown peat with
wood fragments

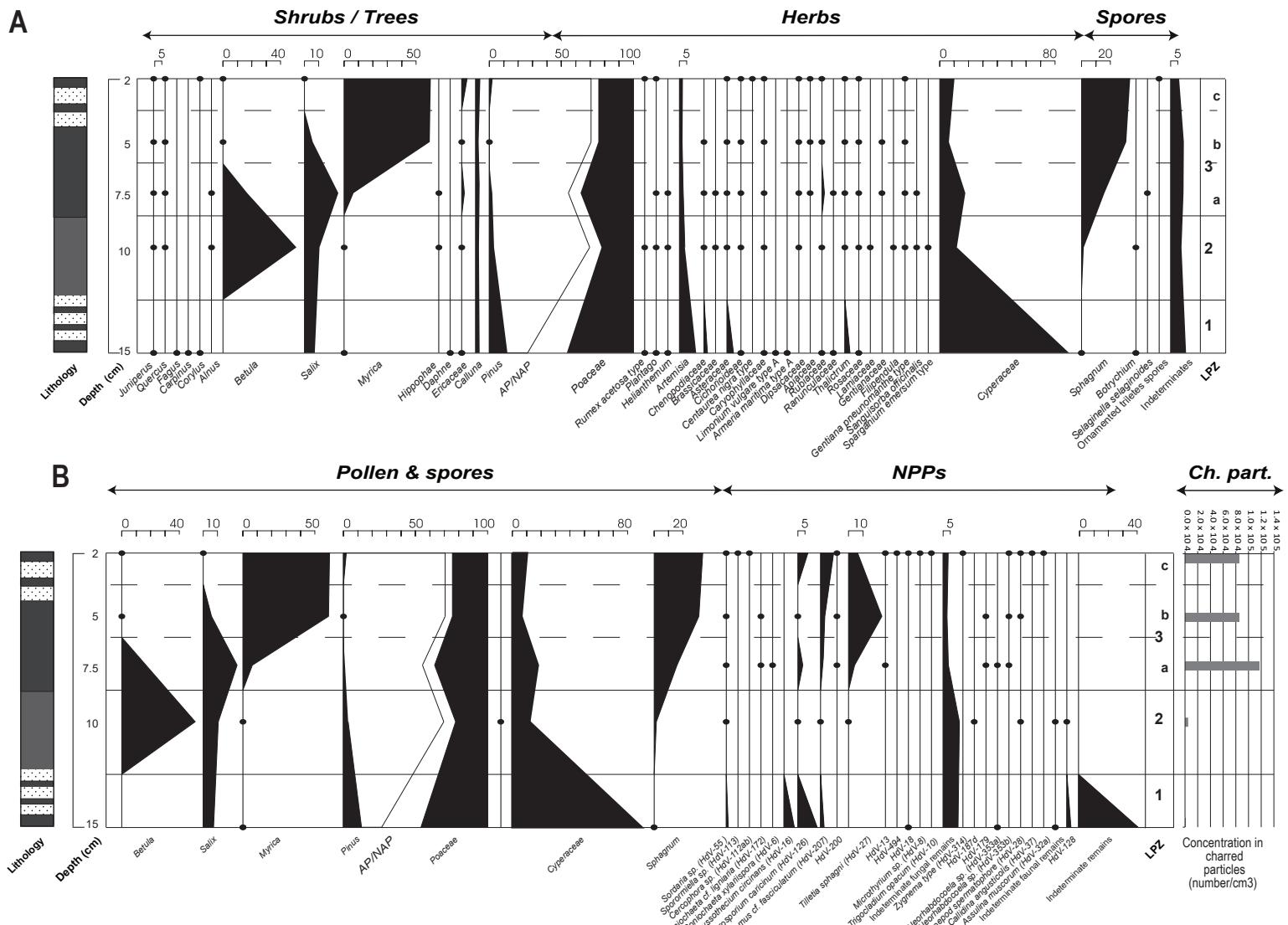
Alternating sand and
black peat beds

A



B

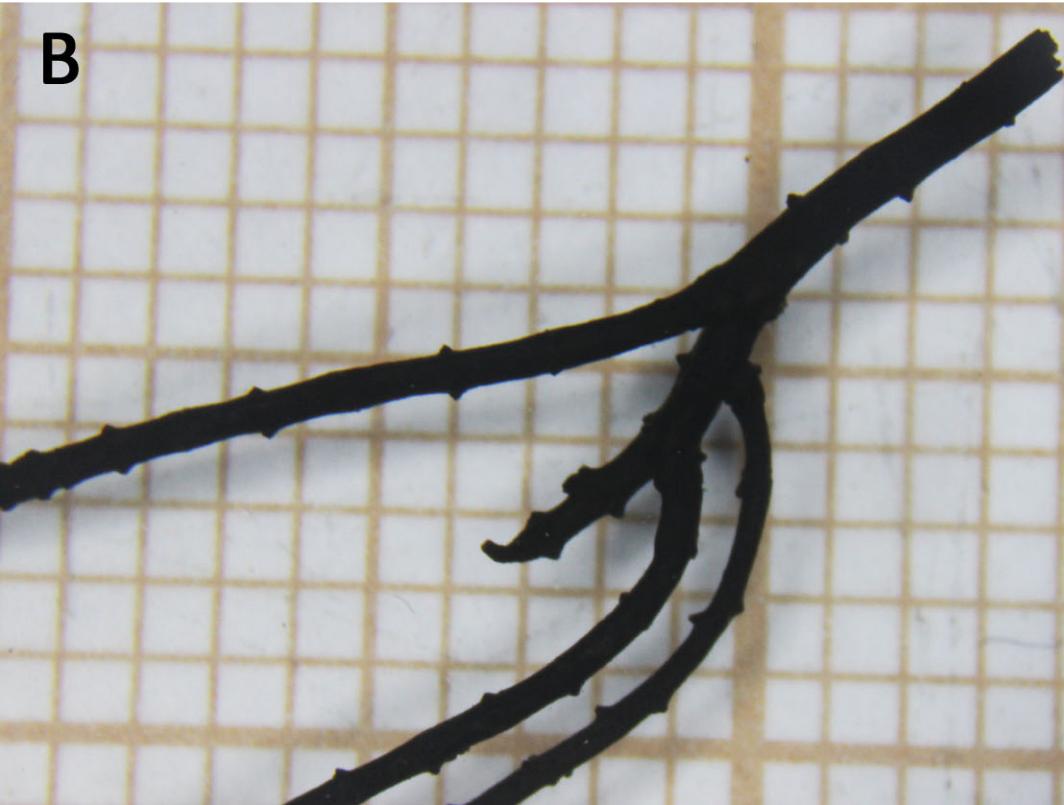


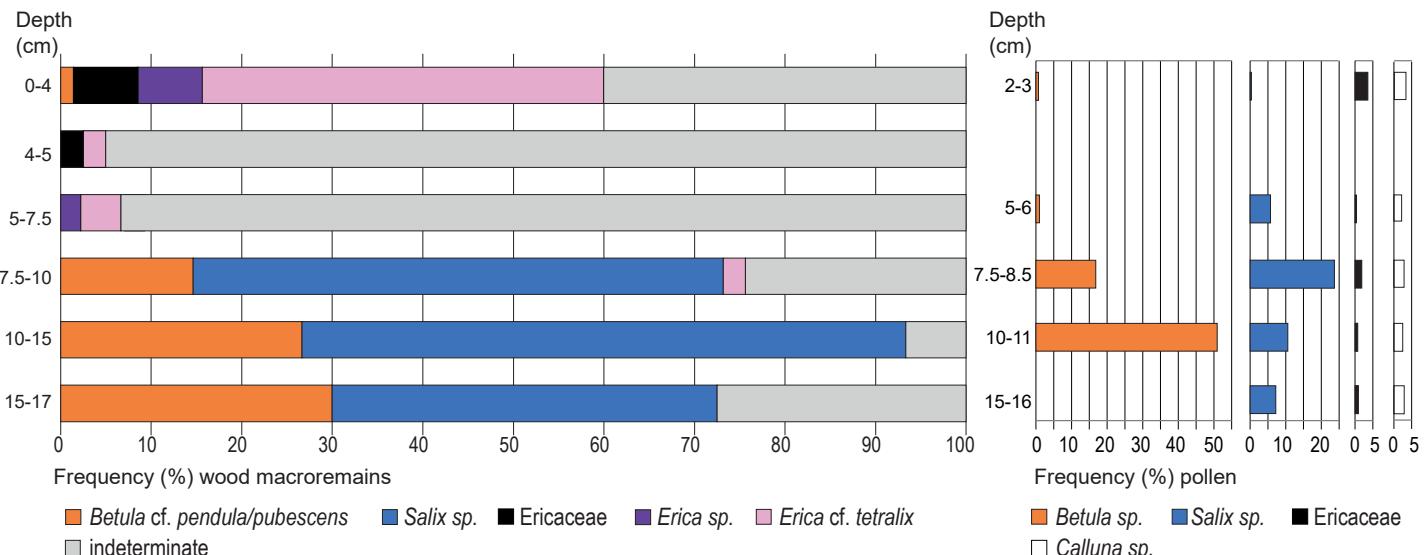


A



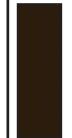
B





Sample number	Depth (cm)	Volume of sieved sediment (cm ³)	Total number of fragments	Concentration (fragment number/100 cm ³)	Number of fragments analysed
1	0 - 4 cm	250	607	243	70
2	4 - 5 cm	100	56	56	40
3	5 - 7.5 cm	400	130	33	45
4	7.5 - 10 cm	400	131	33	41
5	10 - 15 cm	300	82	27	45
6	15 - 17 cm	400	88	22	40

Depth (cm)	Number of pollen grains and spores counted	Number of pollen grains and spores identified	Concentration (number of pollen grains and spores/cm ³ of fresh sediment)	Preservation state	Diversity (Pollen - Spores + NPPs)
2-3	931	896	238013	good, some corroded pollen grains	27 + 18
5-6	818	766	259602	good	23 + 10
7,5-8,5	636	594	97557	good, some corroded pollen grains	31 + 12
10-11	651	608	82075	good	32 + 9
15-16	2084	2059	55434	good	28 + 9

Lithology	Dep. (cm)	Pollen & spores	NPPs	Charred particles (10 µm-200 µm)	LPZ	Wood macroremains
	2-3	Myrica & Sphagnum stable // decrease of Salix.	Decrease of <i>Tilletia sphagni</i> (HdV-27) // rise of <i>Glomus cf. fasciculatum</i> (HdV-207) & <i>Clasterosporium carincinum</i> (HdV-126). Increase of <i>Assulina muscorum</i> (HdV-32a) & <i>Callidina angusticollis</i> (HdV-37).	Charred particles stable.	c	0-4 cm Increase of <i>Erica cf. tetralix</i> and <i>Erica</i> > und. woods > Ericaceae > <i>Betula</i>
	5-6	Extension of Myrica // decline of Salix, Betula, Pinus & Ericaceae. Decrease of Poaceae, Cyperaceae & Rubiaceae. Rise of Sphagnum.	Increase of <i>Tilletia sphagni</i> (HdV-27) // decrease of <i>Clasterosporium carincinum</i> (HdV-126).	Charred particles stable.	3 b	4-5 cm Undetermined woods > <i>Erica cf. tetralix</i> & Ericaceae.
	7.5-8.5	Increase of Myrica & Sphagnum Recovery of Poaceae & Cyperaceae. Slight development of Rubiaceae and appearance of Apiaceae.	Rise of <i>Tilletia sphagni</i> (HdV-27), <i>Clasterosporium carincinum</i> (HdV-126) & <i>Glomus cf. fasciculatum</i> (HdV-207). Appearance of <i>Coniochaeta cf. lignaria</i> (HdV-172), <i>Coniochaeta xylariispora</i> (HdV-6), <i>Nearhabdocoela</i> sp. (HdV 353b), HdV-13, HdV-179 & HdV-200.	Expansion of charred particles	a	5-7.5 cm Rise of undetermined woods > <i>Erica</i> sp. and <i>Erica cf. tetralix</i>
	10-11	Rise of <i>Betula</i> // slight decrease in <i>Pinus</i> but <i>Salix</i> and <i>Calluna</i> stable. Herbaceous decline: mainly Cyperaceae but also Asteraceae, Chenopodiaceae & <i>Thalictrum</i> . Diversification towards other herbs. Slight development of Sphagnum.	Disappearance of <i>Byssothecium circinans</i> (HdV-16), <i>Clasterosporium carincinum</i> (HdV-126), <i>Glomus cf. fasciculatum</i> (HdV-207) & undetermined remains. Appearance of <i>Tilletia sphagni</i> (HdV-27) & copepod spermatophores (HdV-28) and HdV-187d.	Slight increase of charred particles.	2	7.5-10 cm Decrease of <i>Betula</i> < <i>Salix</i> . Appearance of <i>Erica cf. tetralix</i>
	15-16	NAP > AP (scarce <i>Pinus</i> , <i>Salix</i> & <i>Calluna</i>) Cyperaceae > Poaceae > <i>Artemisia</i> > Asteraceae - Chenopodiaceae - <i>Thalictrum</i>	Predominance of undetermined remains (including faunal remains) <i>Clasterosporium carincinum</i> (HdV-126) > <i>Byssothecium circinans</i> (HdV-16) > <i>Glomus cf. fasciculatum</i> (HdV-207) > HdV-128 - <i>Sordaria</i> sp. (HdV-55)	Charred particles very scarce	1	10-15 cm Rise of <i>Salix</i> > <i>Betula</i> 15-17 cm <i>Salix</i> > <i>Betula</i> > undetermined woods

Depth	Taxa	Number	Percent	Preservation state			Mean length/min./max. (mm)	Mean width/min./max. (mm)	Diameter (mm)	Curvature of wood rings		
				good	medium	low				0-2.5	2.5-5.0	high
0-4 cm	<i>Betula</i> sp.	1	1.4		1		11.0	2.0			1	
	<i>Erica cf. tetralix</i>	31	44.3	31			14.4/4.0/27.0	1.3/0.5/2.5	28	1	31	
	<i>Erica</i> sp.	5	7.1	4	1		14.5/0.8/26.0	0.8/0.5/3.0	4		5	
	Ericaceae	5	7.1	5			21.6/16.0/35.0	1.7/1.0/2.0	3		5	
4-5 cm	indetermined	28 (including 21 root fragments)	40.0	3	20	5	15.5/5.0/40.0	3.1/1.0/6.0	4	7	23	5
	<i>Erica cf. tetralix</i>	1	2.5	1			18.0	2.0	1		1	
	Ericaceae	1	2.5	1			6.0	1.5	1		1	
	indetermined	38 (including 28 root fragments)	95	3	28	7	10.6/5.0/24.0	2.4/1.0/5.0	10	2	28	10
5-7.5 cm	<i>Erica cf. tetralix</i>	2	4.4	2			10.5/9.0/12.0	0.7/0.5/1.0	2		2	
	<i>Erica</i> sp.	1	2.2	1			11.0	1.0	1		1	
	indetermined	42 (including 23 root fragments)	93.3	11	13	18	13.9/5.0/50.0	3.8/1.0/20.0	7	3	27	15
	<i>Betula cf. pendula/pubescentis</i>	6	14.6	1	5		10.8/6.0/20.0	4.2/3.0/5.0	1		5	
7.5 - 10 cm	<i>Salix</i> sp.	24	58.5	4	18	2	16.6/6.0/51.0	3.4/2.0/5.0	5	3	16	
	<i>Erica cf. tetralix</i>	1	2.4	1			19.0	1.0	1	1		
	indetermined	10 (including 2 root fragments)	24.4	1	6	3	10.7/6.0/18.0	3.4/2.0/5.0	3		7	
	<i>Betula cf. pendula/pubescentis</i>	12	26.7	8	4		13.0/6.0/37.0	3.5/2.0/5.0	1	1	3	9
10-15 cm	<i>Salix</i> sp.	30	66.7	18	12		14.6/4.0/38.0	4.5/2.0/17.0	11		19	
	indetermined	3	6.7				8.7/6.0/10.0	3.0/2.0/4.0	2	1	3	
	<i>Betula cf. pendula/pubescentis</i>	12	30	1	2	9	11.0/6.0/20.0	3.0/2.0/5.0	1		5	7
	<i>Salix</i> sp.	17	42.5	8	2	7	12.3/5.0/30.0	2.1/1.5/3.0	13		4	
15-17 cm	indetermined	11	27.5	5	2	4	11.1/5.0/18.0	2.3/2.0/3.0	1	6	5	