



Relating pollen representation to an evolving Amazonian landscape between the last glacial maximum and late Holocene

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1 **Relating pollen representation to an evolving Amazonian landscape**
2 **between the last glacial maximum and Late Holocene**

3

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21 **ABSTRACT**

22 In contrast to temperate regions, relationships between basin characteristics (e.g. type/size)
23 and fossil pollen archives have received little attention in Amazonia. Here, we compare fossil
24 pollen records of a small palm swamp (Cuatro Vientos; CV) and a nearby large lake (Laguna
25 Chaplin; LCH) in Bolivian Amazonia, demonstrating that palm swamps can yield Quaternary
26 pollen archives recording the history of terrestrial vegetation beyond the basin margin, rather
27 than merely a history of localized swamp vegetation dynamics. The pollen assemblages from
28 these two contrasting basins display remarkable agreement throughout their late Quaternary
29 history, indicating past drier climates supported savanna landscape during the last glacial
30 maximum (LGM; 24,000-18,000 cal yr BP) and savanna/semi-deciduous forest mosaic during
31 the middle Holocene (7000-4750 cal yr BP) at both regional (inferred from LCH) and local
32 (inferred from CV) spatial scales. Additionally, the local-scale catchment of CV and the basin's
33 proximity to the riverine forests of the Río Paraguá enables exploration of the extent of
34 gallery/riverine forests during the LGM and middle Holocene. We show that, between 24,000-
35 4000 cal yr BP, riverine/gallery rainforests were substantially reduced compared with present,
36 challenging the hypothesis that gallery rainforests were important refugia for rainforest species
37 during the drier LGM and middle Holocene.

38

39 **KEYWORDS**

40 Paleoecology, Quaternary, Pollen, Bolivian Amazonia, Palm Swamp, last glacial maximum,
41 Holocene

42

43 **INTRODUCTION**

44 The role of paleoecology in determining how Amazonian ecosystems responded to long-term
45 past climate change is of paramount importance, given its relevance for understanding the
46 fate of Amazonia under future climate change. Particular focus should be given to ecotonal
47 regions of Amazonia, where humid evergreen forests form boundaries with, or grade into,
48 savannas and/or semi-deciduous tropical dry forests. Rainforest taxa at these ecotones exist
49 near to their climatic limits and should therefore be highly sensitive to climate change. Existing
50 paleoecological records have demonstrated this vulnerability, with evidence for climate-
51 induced expansion of savanna and/or dry forests during the last glacial maximum (LGM) and
52 the middle Holocene in ecotonal eastern Amazonia (e.g. Absy et al., 1991; Hermanowski et
53 al., 2012; Fontes et al., 2017; Reis et al., 2017) and southern Amazonia (Mayle et al., 2000;
54 Burbridge et al., 2004; Carson et al., 2014). However, the paucity of these paleo-records
55 means that considerable uncertainty exists as to the full nature and extent of these biome
56 shifts.

57
58 Unfortunately, finding suitable paleoecological sites is often a challenge in this region. The
59 dynamic hydrology of Amazonia means that long-lived, permanent lake basins are uncommon
60 (Colinvaux et al., 1985; Latrubesse, 2012). Small oxbow lakes are widespread, but they rarely
61 have sediment records that span the multi-millennial timescales needed to capture long-term
62 climate change (Toivonen et al., 2007; Latrubesse, 2012; Rodriguez-Zorro et al., 2015).

63
64 In the absence of suitable lake sediment records, bogs and palm swamps are often targeted
65 for paleoecological analysis. However, their value is often called into question due to
66 uncertainty over whether their pollen archives reliably capture the history of local terrestrial
67 vegetation beyond the bog/swamp, or instead merely reveal the history of swamp/bog
68 vegetation growing within the basin itself. In the latter case, they are of little use for
69 paleoecologists seeking to understand Holocene/Quaternary forest dynamics. Such concerns
70 are borne out by pollen records from sites such as the Pantano de Monica palm swamp in the
71 central Colombian Amazon (Behling et al., 1999) and the Vereda de Águas Emendadas palm
72 swamp in central Brazil (Barberi et al., 2000), both of which are dominated by swamp taxa
73 (e.g. palms and sedges) through much of the Holocene. Furthermore, because key pollen taxa
74 such as grass and sedge can only be identified to family level (Poaceae and Cyperaceae,
75 respectively), it is often unclear whether their presence signifies semi-aquatic species growing
76 within the swamp (e.g. floating sedge mat) or instead open, seasonally-flooded savanna
77 beyond the swamp.

78

79 Here, we present the results of a natural experiment, whereby analysis of fossil pollen records
80 from a small palm swamp and an adjacent large lake provides a rare opportunity to determine
81 the potential for palm swamps to reliably record glacial-Holocene terrestrial vegetation
82 histories, or merely a history of swamp vegetation. We present a 24,000-yr fossil pollen record
83 from Cuatro Vientos palm swamp located in Noel Kempff Mercado National Park (NKMNP),
84 north-eastern Bolivia (southern Amazonian forest-savanna ecotone) (Figs. 1 and 2). The
85 current state of knowledge of the late Quaternary paleoecology of this region is predominantly
86 based on the pollen records from two large lakes: Lagunas Bella Vista and Chaplin (Mayle et
87 al., 2000; Burbridge et al. 2004) (Figs. 1 and 2). These records demonstrate that most of the
88 regional catchments of these two lakes, which are today dominated by humid evergreen
89 rainforest, were previously characterized by a mosaic of savanna and semi-deciduous dry
90 forest communities during the LGM and early/middle Holocene under drier-than-present
91 climatic conditions. The regional climate became gradually wetter through the late Holocene,
92 causing the progressive replacement of savanna and dry forest by humid evergreen rainforest,
93 which expanded in the northern part of NKMNP ~3000 cal yr BP (around Laguna Bella Vista)
94 and attained current levels in the south of the park (around Laguna Chaplin) by ~750 cal yr
95 BP (Mayle et al. 2000; Burbridge et al. 2004).

96

97 Cuatro Vientos (CV) palm swamp is located only 6.5 km from Laguna Chaplin (Fig. 2), and
98 thus presents a unique opportunity to directly compare the paleoecological record of a palm
99 swamp with that of a neighboring large lake (~25 km² basin). Given the close proximity of
100 these two sites, located within the same vegetation type (humid evergreen rainforest), we
101 expect that they will have undergone the same climatic and regional vegetation changes in
102 the past. Therefore, our assumption is that any differences between the paleoecological
103 records can be attributed to the effects of basin type and/or basin size, thus enabling a robust
104 assessment of the potential value of palm swamps as repositories of paleoecological data in
105 southern Amazonia; e.g. whether they reflect a history of *terra firme*, climate-driven vegetation
106 change beyond the swamp, or merely a history of a palm swamp community controlled by
107 local hydrological conditions within the basin. The findings of our study may have implications
108 for the interpretation of other palm swamp records elsewhere in the neotropics, as well as
109 criteria for the selection of appropriate sites for paleoecological analyses and palaeo-data
110 syntheses.

111

112 If it is found that the Cuatro Vientos record does provide a long-term (multi-millennial) record
113 of vegetation changes beyond the swamp itself, the pairing of Cuatro Vientos and Laguna
114 Chaplin also provides an opportunity to explore the dynamics of local versus regional-scale
115 vegetation changes in the park. This strategy of pairing small and large neighboring

116 sedimentary basins has long been advocated as a sound approach for differentiating local
117 versus regional pollen catchments in mid to high latitude North America and Europe (Jacobson
118 and Bradshaw, 1981). In these temperate ecosystems, where most tree taxa are wind-
119 pollinated, modelling approaches based upon pollen productivity and dispersal data have led
120 to quantitative estimates of pollen catchment area, whereby large lakes ($>5 \text{ km}^2$) have
121 regional-scale pollen catchment areas ($> \sim 50 \times 50\text{--}100 \times 100 \text{ km}$) and are relatively insensitive
122 to localized or patch-size vegetation changes (Sugita, 1994; Sugita et al., 1999; Davis, 2000;
123 Sugita, 2007a), whereas small lakes ($< \sim 0.1\text{--}1 \text{ km}^2$) instead have local-scale pollen catchment
124 areas ($< \sim 10 \times 10 \text{ km}$) (Sugita, 2007b). These temperate ecosystem pollen catchment
125 estimates (e.g. 1 km^2 cut-off between local *versus* regional catchments) are unlikely to hold
126 true for humid tropical rainforests due to the different constituent taxa and far greater
127 complexity of pollination syndromes (wind, insects, bats, birds) associated with these more
128 biodiverse ecosystems. However, in our study area at least, modern pollen rain studies
129 (Gosling et al., 2005, 2009; Burn et al., 2010) show that wind-pollinated Moraceae pollen
130 dominates rainforest pollen assemblages in NKMNP. The general premise that large lakes
131 and small lakes capture regional- and local-scale pollen rain, respectively, therefore likely
132 holds true, corroborated by Carson et al. (2014). The area of the Cuatro Vientos swamp basin
133 is $\sim 5 \text{ km}^2$, compared with $\sim 25 \text{ km}^2$ for the neighboring Laguna Chaplin basin (Fig. 2), thus
134 enabling local-scale vegetation dynamics (Cuatro Vientos) to be differentiated from regional-
135 scale vegetation dynamics (Laguna Chaplin).

136

137 The local-scale catchment of Cuatro Vientos is particularly pertinent given the location of this
138 site at the margin of the riverine forests of the Río Paraguá (Figs. 1 and 2), as this provides a
139 unique opportunity to investigate the Quaternary history of riverine/gallery rainforest. During
140 drier periods of the Pleistocene when humid evergreen rainforest cover was reduced, it has
141 previously been proposed that rainforest taxa may have survived within refugia provided by
142 riverine gallery rainforest, due to the more continuous water supply from the river (Meave et
143 al., 1991; Meave and Kellman, 1994; Pennington et al., 2000). These gallery rainforest refugia
144 may have provided important routes and source areas for the spread of plant and animal
145 species (Redford and da Fonseca, 1986; Meave et al., 1991; Costa, 2003), as well as
146 providing routes for human population expansion (Iriarte et al., 2017). Investigating the extent
147 of gallery rainforests in NKMNP through the late Quaternary may also help to explain the
148 mechanism of rainforest expansion in the late Holocene, e.g. whether the gallery rainforests
149 served to expedite the spread of rainforest taxa in response to climate change (e.g. Mayle et
150 al., 2007). However, the extent to which these gallery rainforests survived through the drier
151 climatic periods of the LGM and middle Holocene in NKMNP is uncertain, given that, until now,
152 only regional-scale vegetation records are available from pollen data from the two large lakes

153 in this area (i.e. Lagunas Chaplin and Bella Vista), which lack the spatial resolution to capture
154 changes in the extent of riverine vegetation.

155

156 This paper addresses the following questions:

157 1. How does the Quaternary paleoecological record from a small (~ 5 km²) Amazonian
158 palm swamp (Cuatro Vientos) in Noel Kempff Mercado National Park (NKMNP)
159 (ecotonal southern Amazonia) compare with that of a neighboring large lake (Laguna
160 Chaplin, ~25 km²), and what does this comparison reveal about the suitability of palm
161 swamps as fossil pollen archives for investigating Amazonia's Quaternary vegetation
162 history?

163 2. What does the palaeoecological record from Cuatro Vientos, located close to a river,
164 reveal about the extent of riverine gallery rainforest in NKMNP during the drier climatic
165 conditions of the LGM and middle Holocene when the interfluves were dominated by
166 savanna and/or semi-deciduous tropical dry forest?

167 3. What are the implications of this palm swamp study for assessing the role of gallery
168 forest as rainforest migration corridors or refugia under drier climatic conditions?

169

170 **STUDY AREA**

171 Noel Kempff Mercado National Park (NKMNP) is a 15,230 km² protected reserve located near
172 the southern margin of the Amazon basin in north-eastern Bolivia (Fig. 1) (Killeen and
173 Schulenberg, 1998). The park has been designated a UNESCO world heritage site due to its
174 exceptionally high *beta* (habitat) diversity and is largely undisturbed by modern anthropogenic
175 land use (Killeen et al., 2003; Heyer et al., 2018).

176

177 *Geomorphology and regional vegetation*

178 NKMNP is located on the western reach of the Precambrian Brazilian shield, the
179 geomorphology of which splits the park into two distinct landscapes (Fig. 1). To the east, the
180 park is dominated by the Huanchaca Plateau, a table-mountain ~600–900 m above sea level
181 (a.s.l.) comprized of Precambrian sandstone and quartzite. The plateau is predominantly
182 covered in upland *cerrado* savanna vegetation that has been present since at least the end of
183 the last glacial period (Maezumi et al., 2015). To the west lies a lowland peneplain, where the
184 Precambrian bedrock is blanketed by Tertiary and Quaternary alluvial sediments and is
185 covered predominantly in *terra firme* humid evergreen tropical forest (HETF). The clear-water
186 rivers of the Río Iténez and Río Paraguá form the north/eastern and western boundaries of
187 NKMNP, respectively. These rivers and other smaller streams in the park are lined by
188 evergreen riverine forests, usually on the natural levees that form from deposition events
189 during seasonal flooding. Patches of seasonally-inundated savanna occur near the rivers
190 where soil drainage is poor. The southern border of NKMNP defines the modern ecotone
191 between the HETF of southern Amazonia and the Chiquitano semi-deciduous tropical dry
192 forest (SDTF) of eastern lowland Bolivia. The term ‘semi-deciduous’ is used here to describe
193 the flexible phenologic response (deciduousness) of the constituent trees, depending on the
194 degree and duration of the dry season (Killeen et al., 1998; Killeen and Schulenberg, 1998).
195 In contrast to the HETF, the SDTF supports a denser understorey vegetation as more light
196 can penetrate the canopy.

197

198 *Climate*

199 The precipitation regime of the region is distinctly seasonal, predominantly controlled by the
200 South American Summer Monsoon (SASM; Zhou and Lau, 1998; Raia and Cavalcanti, 2008;
201 Silva and Kousky, 2012). The majority of the ~1400–1600 mm mean annual precipitation falls
202 during the wet season during austral summer, with a dry season lasting for ~4–6 months
203 during austral winter. Mean annual temperatures are ~25–26°C, with little monthly variation.
204 However, during austral winter, cold fronts (‘surs’ or ‘surazos’) originating in Patagonia can

205 reach the area and cause temperatures to drop below 10°C for several days (Killeen et al.,
206 2003).

207

208 *Site descriptions*

209 Cuatro Vientos (CV, 14°31'18.5"S, 61°7'11.3"W; elevation ~170 m a.s.l.) is a palm swamp, ~5
210 km² in area, located in western NKMNP, ~5 km from the Río Paraguá (Figs. 1 and 2). Although
211 it receives river flood waters during the rainy season, it is not an oxbow. As with the large
212 lakes in NKMNP (Lagunas Chaplin and Bella Vista), the oval-shaped CV likely formed either
213 as a solution hollow or subsidence along faults of the underlying siliceous rocks of the Pre-
214 Cambrian Shield. The surrounding vegetation (beyond the palm swamp) consists of *terra firme*
215 HETF to the east and riverine (riparian) forest of the Río Paraguá immediately to the west.
216 The riverine forests in NKMNP vary in their structure, from young pioneer communities, with
217 trees such as *Cecropia*, *Sapium* and *Acacia*, through to older communities with later
218 successional tree taxa, particularly from the Moraceae family (e.g. *Brosimum lactescens*,
219 *Pseudolmedia* spp., *Ficus* spp.). Most of these species are dioecious and wind-pollinated
220 (anemophilous) and are therefore over-represented in the pollen record due to their prolific
221 pollen production (Bush and Rivera, 2001; Burn et al., 2010). Although similar to communities
222 of *terra firme* evergreen forests, the riverine forests can be distinguished by their sparse
223 understoreys (due to seasonal flooding), smaller stature, flood-tolerant species and lower
224 overall species diversity (Killeen and Schulenberg, 1998; Burn et al., 2010). Growing within
225 the CV basin itself is a floating mat of sedge/grass swamp vegetation, interspersed with small
226 pools of open water and scattered clumps of *Mauritiella* palm trees.

227

228 Laguna Chaplin (LCH, 14°28'12"S, 61°2'60"W; elevation ~170 m a.s.l.) is a large (~12 km²),
229 shallow (2–2.5 m in the dry season), flat-bottomed lake (within a ~25 km² basin), located ~6.5
230 km north-east of CV (Mayle et al., 2000; Burbridge et al., 2004) (Fig. 2). The LCH basin is
231 surrounded by HETF, with a mix of seasonally-inundated riverine forest (around much of the
232 lake margin and along the small, ephemeral streams that flow in and out of LCH) and *terra*
233 *firme* (upland) HETF. Adjacent to the lake, in the southern half of the basin, lies a patch of
234 savanna wetland. Comparison of the modern pollen spectra of the surface sediments of the
235 lake (Burbridge et al., 2004) with pollen trap data from all the constituent plant communities in
236 NKMNP (Gosling et al., 2005; Burn et al., 2010; Jones et al., 2011) reveals that the modern
237 pollen assemblage of this lake originates from both the riverine and *terra firme* HETF
238 ecosystems in the lake catchment. Crucially, however, the regional-scale pollen source area
239 of this large lake means that differentiation of the relative extent of riverine versus *terra firme*
240 ecosystems is not possible. Both LCH and CV are located ~30 km from the modern
241 HETF/SDTF ecotone at the southern limit of NKMNP (Fig. 1).

242

243 **METHODS**

244 **Sediment core**

245 Cuatro Vientos (CV) was cored in August 1995 by FM with a modified square-rod Livingstone
246 piston corer (Wright, 1967). The core location was ~300 m from the eastern edge of the palm
247 swamp (Fig. 2), with the inherent difficulty in traversing swamp environments making it
248 impossible to penetrate further into the basin. The top 20 cm of the core site comprized a
249 floating mat of grasses and sedges. Below this was a ~1 m water column, the bottom of which
250 was well mixed with the soft uppermost sediment making it difficult to determine the depth of
251 the sediment-water interface. Therefore, core depths were recorded by reference to the top of
252 the floating mat vegetation (FMV). A 154 cm core was recovered, between 155 and 309 cm
253 below the surface of the FMV. Unfortunately, the sediment above 155 cm was too soft to be
254 recovered. Lithological descriptions are based on the color (using a Munsell soil color chart)
255 and texture of the sediment core. Loss-on-ignition (LOI) analysis was carried out at 4 cm
256 intervals through the CV core. After drying at 100°C for 24 hours, each 1 cm³ sample was
257 combusted at 550°C for 2 hours (LOI₅₅₀). The relative loss of weight before and after
258 combustion determines the percentage organic carbon content that was present in that
259 sample (Dean, 1974; Heiri et al., 2001).

260

261 **Chronology**

262 The chronological framework for CV is based on 9 Accelerator Mass Spectrometry (AMS)
263 radiocarbon (¹⁴C) dates (Table 1). Due to the absence of sufficient plant macrofossils, the
264 majority of the dates were obtained from non-calcareous bulk sediment. However, two of the
265 samples (Beta-467884 and Beta-467885) contained enough decayed plant remains during
266 pre-treatment to be dated. All samples selected for dating were treated to remove any
267 carbonates, and the plant remains were treated to remove mobile humic acids. Radiocarbon
268 ages were calibrated using the IntCal13 calibration curve (Reimer et al., 2013), and a
269 chronology was constructed using the Bayesian age modelling software Bacon v2.3.4 (Blaauw
270 and Christen, 2011). The IntCal13 calibration curve was chosen over SHCal13 because of the
271 hydrological connection of the study area to the northern hemisphere, via the SASM
272 (McCormac et al., 2004; Hogg et al., 2013).

273

274 **Pollen analysis**

275 The CV core was sub-sampled for pollen analysis at 4 cm intervals, apart from between 220–
276 252 cm where sub-samples were taken at 2 cm intervals. The last 29 cm of the core (280–
277 309 cm) was unsuitable for pollen analysis as the sediment had oxidized, preventing pollen
278 preservation. For each horizon, 1 cm³ of sediment was prepared for pollen analysis using
279 standard protocols (Faegri and Iversen, 1989), including hot treatments of 40% HF and 10%

280 NaOH. Samples particularly rich in clay were given pre-treatments of hot 5% sodium
281 pyrophosphate to help disperse the clays, but were not subjected to a fine-sieving stage to
282 ensure small grains (<5 µm) were retained. A known concentration of the exotic marker spore
283 *Lycopodium clavatum* was added to each sample so that absolute pollen concentrations could
284 be calculated (Stockmarr, 1971). Prepared samples were mounted in silicone oil and were
285 counted to the standard 300 Terrestrial Land Pollen (TLP) sum. Cyperaceae pollen was
286 included in the TLP sum (as per Laguna Chaplin, Burbridge et al., 2004) as this taxon is
287 important in the seasonally-flooded savannas of the study region. Pollen identifications were
288 made with reference to published tropical pollen atlases (Roubik and Moreno Patiño, 1991;
289 Colinvaux et al., 1999; Lorente et al., 2017), a freeware digital database of neotropical pollen
290 (Bush and Weng, 2007), and an extensive modern neotropical pollen reference collection of
291 >1500 specimens housed at the laboratory of the Tropical Palaeoecological Research Group,
292 University of Reading. Pollen of the Moraceae/Urticaceae families were grouped into a single
293 'Moraceae' category (with the exception of *Cecropia*). It is notoriously difficult to distinguish
294 between these families and their genera, and given the grains from CV were often obscured
295 or damaged, there was little confidence in genus-level identification, even with the help of
296 published morphological descriptions (Burn and Mayle, 2008). Zones for the pollen data were
297 drawn based on a stratigraphically constrained cluster analysis by incremental sum of squares
298 (CONISS; Grimm, 1987), with the number of statistically significant zones evaluated using the
299 broken-stick model (Bennett, 1996). All analyses and plotting of the pollen data were
300 performed in R (v.3.4.4), using the rioja (v.0.9-15.1) and vegan (v.2.4-6) packages (Juggins,
301 2017; Oksanen et al., 2018).

302

303 **Laguna Chaplin core**

304 Laguna Chaplin (LCH) was cored in 1998 by FM, with the methodology and results of the
305 paleoecological analyses presented in subsequent publications (Mayle et al., 2000; Burbridge
306 et al., 2004; Maezumi et al., 2018b). The pollen data from the analyses of LCH are presented
307 here and compared with those of CV to provide the necessary regional-scale, late Quaternary
308 vegetation and climate context for determination of the paleoecological significance of the CV
309 palm swamp fossil pollen record. We replot the LCH data with an updated age-depth model
310 because the original age-depth model was based on simple linear interpolation between
311 consecutive radiocarbon dates (Burbridge et al., 2004) – a method no longer favored in the
312 paleoenvironmental community (Blaauw et al., 2018). The chronological framework for LCH
313 presented here is based on 14 Accelerator Mass Spectrometry (AMS) radiocarbon (¹⁴C) dates
314 (Table 2) and, as with CV, uses the Bacon Bayesian age modelling software package (Blaauw
315 and Christen, 2011). Note that only the 0-24,000-year portion of the 40,000-year LCH pollen
316 record is plotted here, to allow direct comparison with the 24,000-year CV pollen record.

317 **RESULTS**

318 **Cuatro Vientos - core stratigraphy and chronology**

319 Figure 3 shows the age-depth model derived from Bacon. The model used 7 of the 9 ¹⁴C AMS
320 dates, with the dates at 240 cm and 276 cm rejected based on Bacon's outlier identification.
321 The date at 240 cm was based on a particularly small sample size of extracted decayed plant
322 remains, raising the possibility that the younger-than-expected age could be due to down-core
323 movement of the sample. The date at 276 cm is consistently rejected by multiple Bacon runs,
324 as well as through an exploratory run of OxCal's statistical outlier model (Bronk Ramsey, 1995,
325 2009), and may be anomalously old due to incorporation of older, reworked sediment. The
326 dates for the top (155 cm) and bottom (309 cm) of the core are based on extrapolation, and
327 so must be interpreted with care.

328

329 The sediments from CV can be split into three main stratigraphic sections (Fig. 3).

330 (1) 309–255 cm: comprized of inorganic greyish-brown silty clay, with a sedimentation rate of
331 ca. 0.05 mm/yr. The age range of this section is ca. 33,000–28,000 to 19,000–16,000 cal
332 years BP, corresponding to the late Pleistocene and including the last glacial maximum (LGM).
333 The upper boundary of 255 cm likely marks a hiatus in the core lasting from ca. 19,000–16,000
334 to 12,000–10,500 cal years BP.

335 (2) 255–230 cm: comprized of gray, silty clays, with some organic inclusions. This section
336 corresponds to the early Holocene, between ca. 12,000–10,500 and 8000–7500 cal years BP.
337 A particularly sandy layer of sediment is present within this section, between ca. 238–232 cm
338 where pollen preservation is very poor. Sedimentation rates increase to ca. 0.07–0.1 mm/yr.

339 (3) 230–155 cm: comprized of poorly humified black detrital peat with an increased
340 sedimentation rate of ca. 0.2–0.3 mm/yr. The age range of this section is ca. 8000–7500 to
341 4000–3000 cal years BP, spanning the middle Holocene and part of the late Holocene.

342

343 **Cuatro Vientos pollen data**

344 Figure 4 shows the fossil pollen data for the CV core between 155 and 280 cm (below FMV
345 surface). Three statistically significant zones were identified in the cluster analysis, but to aid
346 in interpretation, zone 1 was split into two sub-zones (before and after the hiatus) and an
347 additional zone was added to mark the period of poor pollen preservation between 239 and
348 230 cm (zone 2), thus giving a total of four pollen assemblage zones. The results of CV will
349 be discussed alongside the updated pollen diagram from LCH (Figs. -6).

350

351 *Zone 1a and Zone 1b: 280–240 cm, ca. 24,000–8750 cal yr BP (LGM to Early Holocene);*
352 *includes sediment hiatus ca. 18,000–11,000 cal yr BP*

353 This pollen assemblage has abundant grass (Poaceae; 40–60%) and sedge (Cyperaceae; 5–
354 10%) pollen, and the highest abundance of the herb taxa Asteraceae (5–10%), *Borreria* (~2%)
355 and Amaranthaceae (~1%). Levels of ‘cold-adapted’ taxa such as *Podocarpus*, *Alnus*, and
356 *Ilex* peak in this zone, though at low levels of up to 1%. This is the only zone to contain any
357 significant amounts of *Paullinia/Roupala*; levels of this pollen type are consistent at 3–5% for
358 most of the zone, rising to ~10% near the top of the zone. The savanna indicator *Curatella*
359 *americana* is present, but in low amounts (<1%). Other arboreal taxa are limited, with low
360 quantities (<3%) of Moraceae, *Celtis*, Arecaceae (palms), and *Alchornea* – although
361 *Alchornea* reaches its highest abundance in this zone. Few grains were recovered of the
362 aquatic/semi-aquatic taxa *Sagittaria* and *Isoetes*. In general, the pollen grains recovered in
363 these zones were often degraded. LOI₅₅₀ values are consistently low (~5–10%) throughout
364 these zones, reflective of the inorganic, silty clay sediment.

365

366 *Zone 2: 239–230cm, ca. 8750–7000 cal yr BP (early-middle Holocene)*

367 Pollen preservation in this zone was very poor, most likely a result of the coarse sandy
368 sediment damaging the grains.

369

370 *Zone 3: 230–206cm, ca. 7000–5500 cal yr BP (middle Holocene)*

371 During this mid-Holocene section of the core, pollen characteristic of SDTF become
372 established, including *Anadenanthera* (2–4%), *Astronium* (5–7%) and the understory taxon
373 *Clavija* (3–5%). At the same time, *Curatella americana* becomes more abundant (2–4%) and
374 Poaceae levels remain consistent at 50–60%. Other arboreal taxa remain at low levels,
375 although Moraceae does increase slightly from 4% to 10%. Palm taxa are uncommon in this
376 zone. Levels of weed and herb taxa (e.g. Asteraceae, *Borreria*) as well as the ‘cold-adapted’
377 taxa found in Zone 1, decrease to negligible amounts. The aquatic/semi-aquatic taxa
378 *Sagittaria* and *Isoetes* increase throughout this zone, with *Sagittaria* in particular becoming a
379 large proportion of the total pollen sum (~20%) between ca. 6000–5500 cal yr BP. A sudden
380 increase in LOI₅₅₀ occurs at ca. 6,000 cal yr BP (from <20 to ~80%), reflecting the switch from
381 clay to organic, peaty sediment (Fig. 3).

382

383 *Zone 4: 206–155cm, ca. 5500–3750 cal yr BP (middle to late Holocene)*

384 This zone is similar to zone 3, with only subtle differences in the abundances in some of the
385 taxa. Moraceae percentages stabilize at ~10%, with other HETF arboreal taxa remaining at
386 low/negligible levels. Palm (Arecaceae spp., *Mauritia/Mauritiella*) pollen grains have a more
387 consistent presence in the assemblage, being present in most samples within this zone, but
388 only at very low levels (<1%). *Curatella americana* decreases slightly towards the top of this
389 zone. *Sagittaria* becomes established at 20–30% of the total pollen sum, with *Isoetes*

390 decreasing slightly from Zone 3a to values of 0–5%. LOI₅₅₀ values remain consistent at ~90%
391 through this zone.

392 **INTERPRETATION and DISCUSSION**

393 **Comparison between pollen records of Cuatro Vientos palm swamp and Laguna** 394 **Chaplin**

395 *Last glacial maximum (ca. 24,000–18,000 cal yr BP)*

396 The paleoecological data indicate that the Cuatro Vientos (CV) basin was markedly different
397 during the LGM compared with today. The scarcity of palms (*Arecaceae* undiff. and
398 *Mauritia/Mauritiella*) in the pollen record is a clear indication that, unlike today, there were no
399 substantial stands of palm trees growing on or near the basin during this time. Additionally,
400 *Mauritiella* palm swamps, such as CV, are characterized by highly organic, peaty sediment.
401 However, this LGM section of the core is characterized by fine-grained inorganic clay sediment
402 (<10% LOI₅₅₀), suggestive of a low-productivity, low-energy lake rather than a peat swamp.
403 The absence of emergent macrophytes (e.g. *Sagittaria*, *Isoetes*) suggests that CV was
404 unsuitable for supporting aquatic/semi-aquatic vegetation, possibly indicating very low water
405 levels and the basin perhaps drying out seasonally. Low water levels would be consistent with
406 the regional paleoclimate reconstructions of a drier LGM in this region, not only from
407 neighboring LCH (Burbridge et al., 2004), but also Laguna La Gaiba ~500 km to the south
408 (Whitney et al., 2011; Metcalfe et al., 2014). An intermittently dry basin may also account for
409 the generally degraded nature of the pollen grains in this section of the core, whereby the
410 grains are exposed to short-term oxidation. We acknowledge the possibility that differential
411 pollen preservation of different taxa may conceivably introduce a degree of bias into these
412 pollen assemblages, although no significant taxonomic bias was apparent when counting the
413 pollen.

414

415 If our interpretation that CV was an open lake (rather than a palm swamp) at this time is
416 correct, the LGM pollen assemblage would reflect the local vegetation growing *outside* the
417 basin, rather than *within* the basin. This gives us confidence that CV is a useful repository of
418 paleoecological data during the LGM, reflecting the history of *terra firme* vegetation changes
419 beyond the basin. Therefore, any significant differences between the pollen records of LCH
420 and CV will most likely be a result of basin size (i.e. regional *versus* local pollen catchments,
421 respectively), rather than basin type (i.e. lake *versus* palm swamp). Although CV today
422 receives flood water from the neighboring clear-water Rio Paragua during the rainy season,
423 and likely did so throughout its late Quaternary history, we are confident that this riverine pollen
424 input is negligible compared with terrestrial pollen sources to the lake, based on modern
425 pollen-vegetation comparisons from several other flood-water lakes across lowland Bolivia.
426 The latter reveal modern pollen assemblages dominated by pollen inputs from local terrestrial

427 vegetation in the case of small lakes such as L. Granja (Carson et al., 2014) or regional
428 terrestrial vegetation with respect to large lakes such as L. Chaplin (Mayle et al., 2000), L. La
429 Gaiba (Whitney et al., 2011), and L. Oricore (Carson et al., 2014), despite receiving seasonal
430 flood waters from neighboring rivers.

431

432 The LGM pollen records from CV and LCH are remarkably similar (Figs. -6), both indicating
433 an open landscape covered with grasses, herbs and sparse tree cover typical of an open
434 savanna. There is abundant Poaceae alongside relatively high percentages of other terrestrial
435 herbs (*Borreria*, Asteraceae) and Cyperaceae. The modern pollen rain study of Jones et al.
436 (2011), based on pollen trap samples from 1 ha ecological plots within NKMNP, suggests that
437 this type of assemblage may be characteristic of an open seasonally-flooded savanna,
438 favoring herbaceous plants that cope better with contrasting seasonal water stresses.
439 Although they are negligible in the CV record, *Mauritia/Mauritiella* palms are slightly more
440 abundant in LCH, which Burbridge et al. (2004) and Jones et al. (2011) use as further evidence
441 of a seasonally-flooded savanna. Nevertheless, Jones et al. (2011) note that the differences
442 between seasonally-flooded and *terra-firme* cerrado savanna are subtle, and the LGM
443 landscape was most likely a mix of these two savanna types. The presence of the woody
444 savanna tree *Curatella americana* is noteworthy as this is a key indicator of savanna
445 environments. Within a seasonally-flooded environment, this species typically grows on top of
446 termite mounds to avoid waterlogging (Killeen and Schulenberg, 1998; Jones et al., 2011);
447 this limited growing area may explain the small quantities of *Curatella americana* in this section
448 of the core, though the fact this species is hermaphroditic and entomophilous (insect-
449 pollinated) will also be a key factor in it being under-represented in the pollen record. Overall,
450 the similarities in the pollen records of CV and LCH show that the local-scale vegetation (CV)
451 was similar to the regional-scale vegetation (LCH), corroborating the interpretations from
452 Burbridge et al. (2004) that much of southern NKMNP was covered in an open savanna during
453 the LGM in response to glacial aridity, lower atmospheric CO₂ levels (Monnin et al., 2001) and
454 cooler temperatures (Stute et al., 1995; Thompson et al., 1998; Whitney et al., 2011).

455

456 Burbridge et al. (2004) infer that the low levels of arboreal rainforest taxa at LCH most likely
457 indicate scarce communities of HETF regionally, most likely existing as gallery rainforests
458 bordering the Río Paraguá. However, the extent of these gallery rainforests is impossible to
459 determine with the regional-scale pollen catchment of LCH. Given that we have established
460 that the LGM pollen record from CV is representative of vegetation growing beyond its basin,
461 the smaller catchment size of CV and the basin's closer proximity to the Río Paraguá allows
462 us to gain more information about the gallery rainforests at this time. Modern pollen rain
463 studies suggest that a closed-canopy gallery rainforest would be expected to contain

464 Moraceae levels of at least 40%, alongside pioneer species such as *Cecropia* and potentially
465 small abundances of taxa such as *Pouteria*, *Sapium* and *Symmeria* (Gosling et al., 2005; Burn
466 et al., 2010). However, the LGM pollen assemblages of CV are not indicative of this kind of
467 gallery rainforest. In particular, Moraceae and *Cecropia* percentages never exceed ~5%,
468 which, given that these taxa are prolific pollen producers and are over-represented in pollen
469 assemblages (Gosling et al., 2005; Burn et al., 2010), is strong evidence against a substantial
470 gallery rainforest lining the nearby Río Paraguá.

471

472 Following Zone 1a, the sediment hiatus at CV from ca. 18,000–11,000 cal yr BP suggests a
473 period of very dry conditions, perhaps causing the basin to dry out completely. Similar hiatuses
474 or periods of low sedimentation have been identified in other basins across lowland Amazonia
475 during the last glacial period, for example: in NKMNP, Laguna Bella Vista (LBV; ca. 110 km
476 north of CV) records a hiatus between ca. 42,500–13,000 cal yr BP and LCH records very low
477 sedimentation rates during this period (Burbridge et al., 2004); in south-eastern Amazonia,
478 hiatuses are recorded on the Serra dos Carajás plateau between ca. 22,000–13,000 cal yr BP
479 (Sifeddine et al., 2001) and at Lago do Saci between ca. 18,200–9200 cal yr BP (Fontes et
480 al., 2017); several basins outlined in Ledru et al. (1998) from across Amazonia and southern
481 Brazil record low sedimentation or hiatuses spanning the LGM.

482

483 *Early to middle Holocene (ca. 11,000–7000 cal yr BP)*

484 The end of the sediment hiatus at CV occurred at ca. 11,000 cal yr BP and is concurrent with
485 the slight change in the lithology of the sediment, with grayer clays and some organic
486 inclusions. Nevertheless, there is no evidence that the basin changed significantly from the
487 shallow open lake of the LGM; the LOI₅₅₀ values remain low and there is no change in the
488 levels of sedge, aquatic or palm taxa. The hiatus termination may indicate slightly wetter
489 conditions in the region, allowing water levels in the basin to rise and for local runoff to
490 increase, inputting more sediment to the basin. The latter is consistent with the paleoclimatic
491 interpretation of increased precipitation levels from ca. 12,200 cal yr BP at Laguna La Gaiba
492 (Whitney et al., 2011) and with the hiatus termination at LBV in the north of NKMNP (Burbridge
493 et al., 2004). Holocene temperatures were ca. 5°C higher than the LGM, with deglacial
494 warming of tropical South America occurring from ca. 19,500 cal yr BP (Seltzer et al., 2002;
495 Whitney et al., 2011). Atmospheric CO₂ levels in the Holocene were ca. 76 ppm higher than
496 the LGM (Monnin et al., 2001). Both temperature and CO₂ levels remained relatively stable
497 throughout the Holocene, prior to the industrial period (Indermühle et al., 1999; van Breukelen
498 et al., 2008; Whitney et al., 2011).

499

500 The CV pollen assemblage in Zone 1b (ca. 11,000–8750 cal yr BP) does not differ significantly
501 from that of the LGM Zone 1a, which suggests that the open savanna persisted in the area
502 into the early Holocene. This pollen assemblage is consistent with that of LCH, which is
503 relatively stable from the LGM through most of the Holocene (including through the hiatus
504 phase at CV). Therefore, even with the increase in precipitation at this time (restarting
505 sedimentation at CV), it clearly wasn't enough to support a humid arboreal landscape. The
506 period of poor pollen preservation in the CV core from ca. 8750 to 7000 cal yr BP (Zone 2) is
507 associated with a layer of sandy sediment. This may reflect a period where fluvial dynamics
508 caused a change in river course so that it flowed near, or even into, CV, therefore creating a
509 higher-energy deposition environment. A higher-energy environment would inhibit deposition
510 of pollen-size particles, instead favoring the deposition of the larger, heavier sand particles,
511 thus potentially explaining the lack of pollen in this section of the core. Additionally, agitation
512 of the pollen grains against the large sandy grains could have caused mechanical damage
513 and poor pollen preservation (Twiddle and Bunting, 2010). In the absence of bracketing C-14
514 dates, the rate of accumulation of this sandy layer is uncertain, although it is conceivable that
515 it was deposited very rapidly, perhaps as a single flood pulse from the neighboring river.

516

517 *Middle Holocene (ca. 7000–5500 cal yr BP)*

518 It has been well established that the middle Holocene was associated with a significantly drier-
519 than-present climate across much of southern hemispheric tropical South America, with peak
520 dryness occurring at ca. 6000 cal yr BP (Baker et al., 2001; Wang et al., 2007; Whitney and
521 Mayle, 2012; Cheng et al., 2013; Kanner et al., 2013; Bernal et al., 2016). The drier climate
522 has been attributed to lower southern-hemispheric summer insolation levels at this time,
523 driven by the precessional cycle of Earth's orbit (Berger and Loutre, 1991), which would have
524 acted to restrict the southerly migration of the Inter-tropical Convergence Zone (Haug et al.,
525 2001) and decrease the strength of the South American summer monsoon (Cruz et al., 2009;
526 Baker and Fritz, 2015).

527

528 The mid-Holocene section of the CV core is marked by significant changes to the CV basin,
529 in particular, a switch to highly organic (LOI₅₅₀ values > ~80%), peaty sediment ca. 6000 cal
530 yr BP (Fig. 3). Concurrent with this dramatic lithological change is increased abundance of the
531 aquatic/semi-aquatic macrophytes *Sagittaria* and *Isoetes*. These changes suggest a change
532 from a clear, shallow, open lake to a more eutrophic environment with high levels of deposition
533 of organics. We infer that this is the start of the transition of CV from a lake to a palm swamp.
534 The timing of this switch is interesting, given that it occurs at the peak of the mid-Holocene
535 drought at ca. 6000 cal yr BP and previous dry conditions during the LGM were associated
536 with low sedimentation rates and a sediment hiatus at CV. A drier mid-Holocene climate would

537 most likely cause a decrease in water levels at CV, but unlike at the LGM, the closer river
538 channel (as argued for in the early-Holocene Zone 2) would cause intermittent flooding at CV
539 inundating the basin with organic matter and maintaining an anaerobic environment.
540 Nevertheless, it is important to note that there is a great diversity of successional pathways in
541 swamp environments that can be caused by a variety of different factors (Behling and
542 Hooghiemstra, 1999; Kelley et al., 2013; Roucoux et al., 2013) which may or may not be
543 related to changes in precipitation. It is possible that this change from inorganic lacustrine
544 sediments to a peat swamp environment reflects hydrarch succession of the basin, whereby
545 a critical ecological threshold or 'tipping point' has been exceeded.

546

547 Despite the change of CV to a swamp basin at this time, we can remain reasonably confident
548 that vegetation growing in the swamp is not masking the influx of pollen from vegetation
549 beyond the swamp. Other than the increase in emergent aquatics (*Sagittaria*, *Isoetes*), there
550 are no increases in other taxa that would be expected to grow in a swamp environment and
551 dominate the pollen rain (e.g. Cyperaceae, *Mauritia/Mauritiella*). Increases in pollen
552 percentages of arboreal taxa that do not grow in a swamp environment (e.g. *Astronium*,
553 *Anadenanthera*) are especially significant, as pollen of these taxa must have come from the
554 surrounding *terra firme* area beyond the perimeter of the basin. Therefore, as with the LGM,
555 this gives us confidence that CV is a useful repository of mid-Holocene paleoecological data,
556 reflecting the history of *terra firme* vegetation changes beyond the basin.

557

558 Considering the clear contrast in the type (palm swamp *versus* lake) and size (5 *versus* 20
559 km²) of the CV basin compared with the neighboring LCH basin, it is perhaps surprising that
560 the pollen records of these two sites are so similar. Both records are indicative of a savanna-
561 SDTF mosaic landscape during this mid-Holocene pollen assemblage. The increased levels
562 of the savanna tree *Curatella americana*, decreased levels of herbs (e.g. Asteraceae) and
563 negligible amount of palm taxa at both sites may suggest that the savanna component was
564 more indicative of a woody cerrado (non-flooded) savanna, rather than the more open
565 seasonally-inundated savanna of the LGM (Jones et al., 2011). This interpretation is plausible
566 given that a weaker mid-Holocene summer monsoon would likely mean less flooding (from
567 the neighboring Río Paraguá) in the rainy season and longer dry seasons. The establishment
568 of *Anadenanthera* and *Astronium* is good evidence of SDTF being present around the basins
569 at this time, both locally (CV) and regionally (LCH), as these are key components of modern
570 SDTF. The *Anadenanthera* pollen type is most likely *Anadenanthera colubrina*, a key drought-
571 tolerant species that is dominant in the modern Chiquitano SDTF region (Killeen and
572 Schulenberg, 1998; Gosling et al., 2009) and a key dry forest indicator, given its absence from
573 both rainforest and savanna ecosystems (Gosling et al., 2009). Both *Anadenanthera* and

574 *Astronium* are often under-represented in pollen assemblages (<1% in modern pollen traps;
575 Gosling et al., 2009); therefore, the relatively high percentages of these taxa (5–7%) suggests
576 they would have been abundant in the area. The similarity between the CV and LCH pollen
577 records provides good evidence that the drier climate of the middle Holocene caused a
578 widespread savanna-SDTF mosaic landscape in the area, at both local (evidenced from CV)
579 and regional (evidenced from LCH) spatial scales.

580

581 The shifting river course into and out of CV that we infer from Zones 2 and 3 suggests that the
582 Río Paraguá ran just as close, if not closer, to CV during the middle Holocene compared with
583 present. Therefore, we may expect that CV would capture a strong gallery rainforest signal in
584 the pollen record. However, as with the previous pollen zones, there are only low levels of
585 arboreal rainforest pollen taxa at CV (10% Moraceae), certainly not at the levels (> 40%
586 Moraceae) expected from a significant gallery rainforest (Burn et al., 2010). We therefore infer
587 that there was insufficient gallery rainforest in NKMNP to provide significant refugia for
588 rainforest species during the drier climate of the middle Holocene. This is somewhat
589 corroborated by other records in southern Amazonia; for example, Laguna Granja, a small
590 oxbow lake ca. 300 km northwest of NKMNP, also shows reduced extent of gallery rainforests
591 during the middle Holocene (Carson et al., 2014).

592

593 *Middle to late Holocene (ca. 5500–3750 cal yr BP)*

594 Following the peak of the mid-Holocene dry period at ca. 6000 cal yr BP, the climate in the
595 region gradually became wetter through the middle to late Holocene (especially after ca. 4,000
596 cal yr BP) (Baker et al., 2001; Wang et al., 2007; Whitney and Mayle, 2012; Cheng et al.,
597 2013; Kanner et al., 2013; Bernal et al., 2016) in response to progressive strengthening of the
598 SASM driven by gradually increasing insolation levels (Berger and Loutre, 1991; Cruz et al.,
599 2005; Baker and Fritz, 2015). In CV Zone 4, the organic, peaty sediment with consistently high
600 LOI₅₅₀ values is now well established, indicating that the basin has remained a swamp
601 throughout this zone. Aquatic vegetation is well represented, with consistently high levels of
602 *Sagittaria*, possibly outcompeting *Isoetes* for space and indicating a continued eutrophic
603 status. However, given that the levels of Cyperaceae and palm taxa remain mostly
604 unchanged, it is unlikely that the basin has yet become a palm swamp analogous to that of
605 today (with the floating mats of grass/sedges and clumps of palms growing throughout the
606 basin). The small increase in percentages of *Mauritia/Mauritiella* pollen is noted, although if
607 these palms were growing abundantly across the CV basin we would expect much higher
608 levels than the 1–3% seen here. Our interpretations are hampered by the absence of surface-
609 sediment samples from CV, which prevents us from determining the pollen signature of the
610 present-day palm swamp. Nevertheless, *Mauritia/Mauritiella* pollen percentages of between

611 10–40% are common for other palm swamps across Amazonia (Behling et al., 1999; Meneses
612 et al., 2015; Rodriguez-Zorro, 2017; Maezumi et al., 2018a) and are therefore likely
613 representative of the modern CV palm swamp as well.

614

615 At both CV and LCH, only small changes occur in the pollen assemblages between Zone 3
616 and 4, with no changes to the overall interpretation of a savanna-SDTF mosaic vegetation
617 cover both locally (around CV) and regionally (inferred from LCH). There are some subtle
618 differences, however, that may indicate some minor changes to the vegetation cover. The
619 small increase in Moraceae in Zone 4 may signify a greater proportion of SDTF relative to
620 savanna in the region, given that: (a) an increase to ~40% Moraceae would be expected from
621 significant expansion of HETF or gallery rainforest (Burn et al., 2010), and (b) the presence of
622 Moraceae in the modern pollen rain of savanna ecosystems is predominantly due to long-
623 distance wind-blown transport from the nearby HETF that was absent in the middle Holocene
624 (Gosling et al., 2009; Jones et al., 2011). The slight increase in *Mauritia/Mauritiella* at CV is
625 concurrent with larger increases seen at LCH, which may suggest the resumption of seasonal
626 flooding at some low-lying areas around the basins. Unfortunately, the CV record terminates
627 at ca. 3750 cal yr BP due to the difficulty in acquisition of uppermost sediments beneath a
628 floating mat of sedge/grass. This means that we cannot corroborate the timing of the increase
629 in HETF at LCH from ca. 2500 to 750 cal yr BP (Burbridge et al., 2004) or determine when the
630 current hydrology developed or when the expansion of palms across the swamp occurred.

631

632 **Implications of the paleoecological history of Cuatro Vientos**

633 Although great strides have been taken in recent years, the number of paleoecological sites
634 that provide information about the Quaternary vegetation history of tropical South America is
635 well below that of the temperate regions of North America and Europe. A recent mid- to late
636 Holocene multi-proxy vegetation reconstruction synthesis by Smith and Mayle (2018) reports
637 110 sites across southern hemispheric tropical South America, although many of these sites
638 were non-pollen based and were clustered in south-east Brazil, with significant gaps across
639 eastern and central Amazonia. Far fewer sites extend to glacial times, with Marchant et al.
640 (2009) reporting only 34 sites for the whole of Latin America in a pollen-based biome
641 reconstruction of the LGM. In contrast, there is good spatial coverage of several hundred mid-
642 Holocene sites across North America (e.g. Prentice et al., 1993; Sawada et al., 2004; Viau et
643 al., 2006) and Europe (e.g. Davis et al., 2003; Wu et al., 2007; Roberts et al., 2018), with
644 growing numbers in east Asia (e.g. Ni et al., 2010; Tian et al., 2017). As a result, tropical South
645 America remains poorly represented in global syntheses (e.g. Gajewski, 2008; Bartlein et al.,
646 2011) and paleodata–model inter-comparison projects (e.g. Kohfeld and Harrison, 2000;
647 Harrison and Prentice, 2003; Braconnot et al., 2012), despite the important role the Amazon

648 rainforest plays in global biogeochemical cycling (Phillips et al., 2009; Pan et al., 2011; Aragão
649 et al., 2014, 2018).

650

651 However, increasing the number of sites in tropical regions such as Amazonia is a complicated
652 task and selecting new target sites for paleoecological analysis is limited by site availability.
653 The challenging logistics of field work in Amazonia means that field seasons are often months
654 long and may only yield data from one or two sites. Therefore, researchers may be reluctant
655 to spend limited time and resources to investigate palm swamps for paleoecological study,
656 given the aforementioned concerns over their suitability for recording terrestrial vegetation
657 history from beyond their basin. Consequently, palm swamps are often viewed as 'sub-
658 optimal' compared with lakes as targets for paleoecological study. However, key lessons to
659 be drawn from the CV record are that: a) Amazonian swamp pollen records can provide useful
660 millennial-scale archives of climate-driven, terrestrial vegetation change beyond the swamp
661 margin, and b) one cannot assume that a palm swamp has always been a palm swamp – a
662 static wetland plant community reflecting purely local-scale basin hydrology, unchanging
663 through time, and unrelated to climate-driven vegetation dynamics elsewhere. Our CV study
664 reveal the importance of considering the potentially dynamic limnological histories of such
665 basins and shows that their present-day characteristics may not be representative of the entire
666 Quaternary sedimentological or catchment history; i.e. Pleistocene lakes with clay sediments
667 have evolved into palm swamps accumulating peat. Within the context of the regional-scale
668 Quaternary vegetation history from neighboring LCH, we have shown that the CV palm swamp
669 was once a lake and contains a fossil pollen archive of local-scale, terrestrial, climate-driven
670 vegetation dynamics extending to the LGM, rather than a localized Quaternary history of a
671 palm swamp plant community. Given the scarcity of Amazonian sedimentary records that
672 extend to the LGM, palm swamps may therefore hold considerably greater value for
673 reconstructing Amazonian Quaternary vegetation change than commonly assumed.

674

675 With regards to the history of riverine/gallery rainforests in NKMNP, the CV pollen record
676 shows that gallery (riverine) rainforest was either absent, or highly limited in extent, along the
677 neighboring Paragua river during the LGM and middle Holocene. Instead, our pollen data
678 reveal that during the LGM and middle Holocene, both the interfluves and riverine areas
679 (presently covered by humid rainforest) were instead covered by a mosaic of savanna and dry
680 forest. Therefore, our findings do not support the hypothesis, at least in our ecotonal area of
681 Amazonia, that during the LGM and middle Holocene wide ribbons of gallery rainforest lined
682 the rivers, providing important refugia for rainforest species. Narrower or non-existent gallery
683 forests during the middle Holocene may have implications for the migration routes of pre-
684 Colombian humans. For example, forest dwelling cultures such as the Tupi-Guarani likely

685 used gallery forests as routes for expansion through non-forested landscapes (Iriarte et al.,
686 2017). If the reduced extent of riverine gallery forests in NKMNP is representative of rivers
687 across ecotonal southern Amazonia, as well as the Cerrado savanna biome to the southeast,
688 it would support the hypothesis that the late Holocene expansion of gallery rainforest (e.g.
689 Silva et al., 2008) linking the Amazonian and Atlantic forest biomes facilitated the trans-
690 continental migration of the forest-dependent Tupi-Guarani culture from southern Amazonia
691 to southern Brazil ca. 2000–3000 cal yr BP (Iriarte et al., 2017).

692

693 The vulnerability of the gallery rainforests in NKMNP to drier mid-Holocene climatic conditions,
694 revealed from the CV record, raises concern over the fate of ecotonal areas of Amazonia
695 under drier climate scenarios predicted for the mid-to-late 21st century (Christensen et al.,
696 2017; Joetzjer et al., 2013; Boisier et al., 2015). Modern field-based ecological impact
697 analyses have shown that tree mortality increases significantly in Amazonian forests in
698 response to severe drought events, although regrowth occurs in subsequent wet years
699 (Phillips et al., 2009; Doughty et al., 2015; Feldpausch et al., 2016). However, these drought
700 events are likely to become more frequent under a future drier climate and if gallery rainforests
701 are not likely to provide refugia for rainforest species, then the resilience of ecotonal, southern
702 Amazonian rainforest would likely be reduced.

703

704 **CONCLUSIONS**

705 The fossil pollen data from the Cuatro Vientos (CV) palm swamp provide a local-scale, late
706 Quaternary vegetation history for southern Noel Kempff Mercado National Park (NKMNP),
707 Amazonian Bolivia, spanning the last glacial maximum (LGM) to the middle Holocene. This
708 local-scale vegetation history complements the previously published, regional-scale
709 vegetation history obtained from the adjacent large lake, Laguna Chaplin (LCH; Mayle et al.,
710 2000; Burbridge et al., 2004). Our results from CV demonstrate that palm swamps in southern
711 Amazonia have the potential to yield Pleistocene-age paleoecological records that provide
712 information about vegetation on *terra firme* landscapes beyond the basin itself, rather than
713 simply recording a history of wetland vegetation within the swamp. Comparison between the
714 CV and LCH pollen records reveals both local- and regional-scale evidence for savannas
715 during the LGM, and a savanna/SDTF mosaic during the middle Holocene. These results
716 demonstrate that the paleoecological value of tropical palm swamps, such as CV, is
717 considerably greater than often assumed – with the potential to yield local-scale, glacial-
718 interglacial histories of climate-driven, terrestrial vegetation dynamics. Although a palm
719 swamp today, the CV site was previously an open-water lake during the LGM, demonstrating
720 that the pollen taphonomy and catchment of this basin has changed markedly through time.

721

722 Due to its local-scale pollen catchment, and close proximity to the Río Paraguá, the CV pollen
723 record also reveals the history of riverine vegetation in ecotonal, southern Amazonia. We find
724 that drier climatic conditions of the LGM and middle Holocene supported expansion of open
725 savanna, not only in the interfluves, but in riverine areas too, challenging the common
726 assumption that rainforest persisted as refugia in ribbons of gallery rainforest lining the rivers.
727 The absence of significant gallery rainforest during past drier climatic conditions raises
728 concerns that gallery rainforest may not be resilient to projected future increased drought and
729 may therefore not be relied upon to serve as rainforest migration corridors, as has previously
730 been proposed (e.g. Mayle et al., 2007).

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1208 **LIST OF TABLES**

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1222 riverine rainforest and the inter-fluvial *terra firme* (non-flooded) rainforest. The red dots
1223 show the coring locations of the two sites. The photo shows *Mauritiella* palm and the
1224 floating sedge/grass mat in the CV palm swamp.

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1227 Vientos

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1229 **Figure 4** – Pollen percentage diagram of taxa from Cuatro Vientos, plotted against
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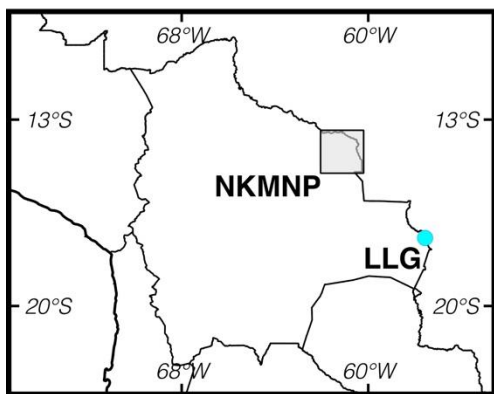
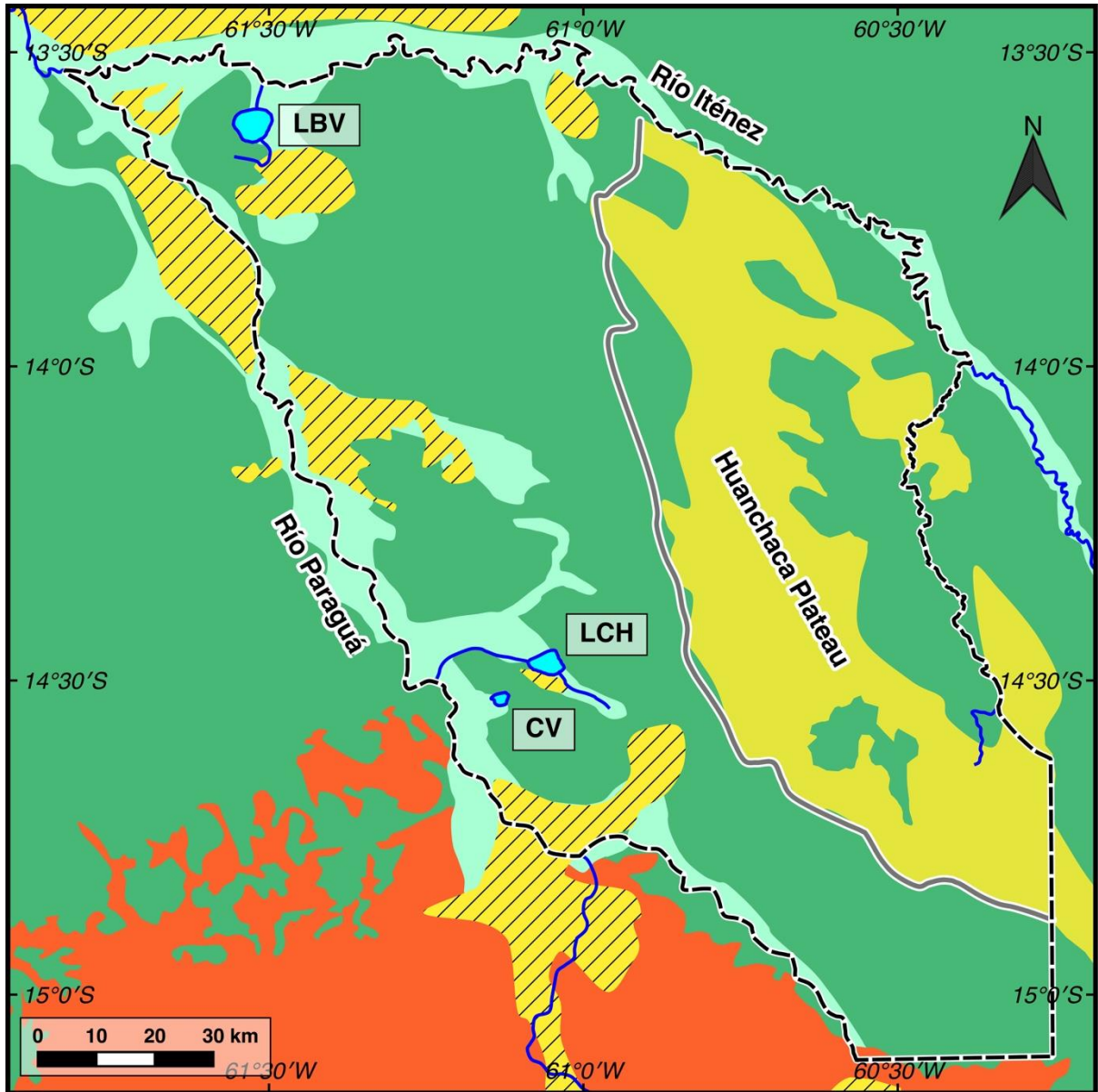
1232 **Figure 5** – Pollen percentage diagram of taxa from Laguna Chaplin, plotted against
1233 calibrated years BP. Dots signify <1% abundance. Zonations are based on the pollen zones
1234 of Cuatro Vientos to aid in comparison.









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1236 **Figure 6** – Summary percentage diagrams for (a) Cuatro Vientos and (b) Laguna Chaplin,
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1238 Groupings as in Figs. 4 and 5: Humid Evergreen Tropical Forest (HETF), Semi-deciduous
1239 Tropical Forest (SDTF), Savanna (SAV), Palm trees (PALM), Cold Adapted Taxa (CAT),
1240 Herbs, weeds and shrubs (HERB), Aquatic/Semi-Aquatic (AQ).

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-  NKMNP boundary
-  Lakes referred to in text
-  Huanchaca Plateau edge
-  Humid evergreen tropical forest
-  Seasonally inundated (riverine) evergreen tropical forest
-  Semi-deciduous tropical forest
-  Seasonally inundated savannah
-  Upland Cerrado savannah

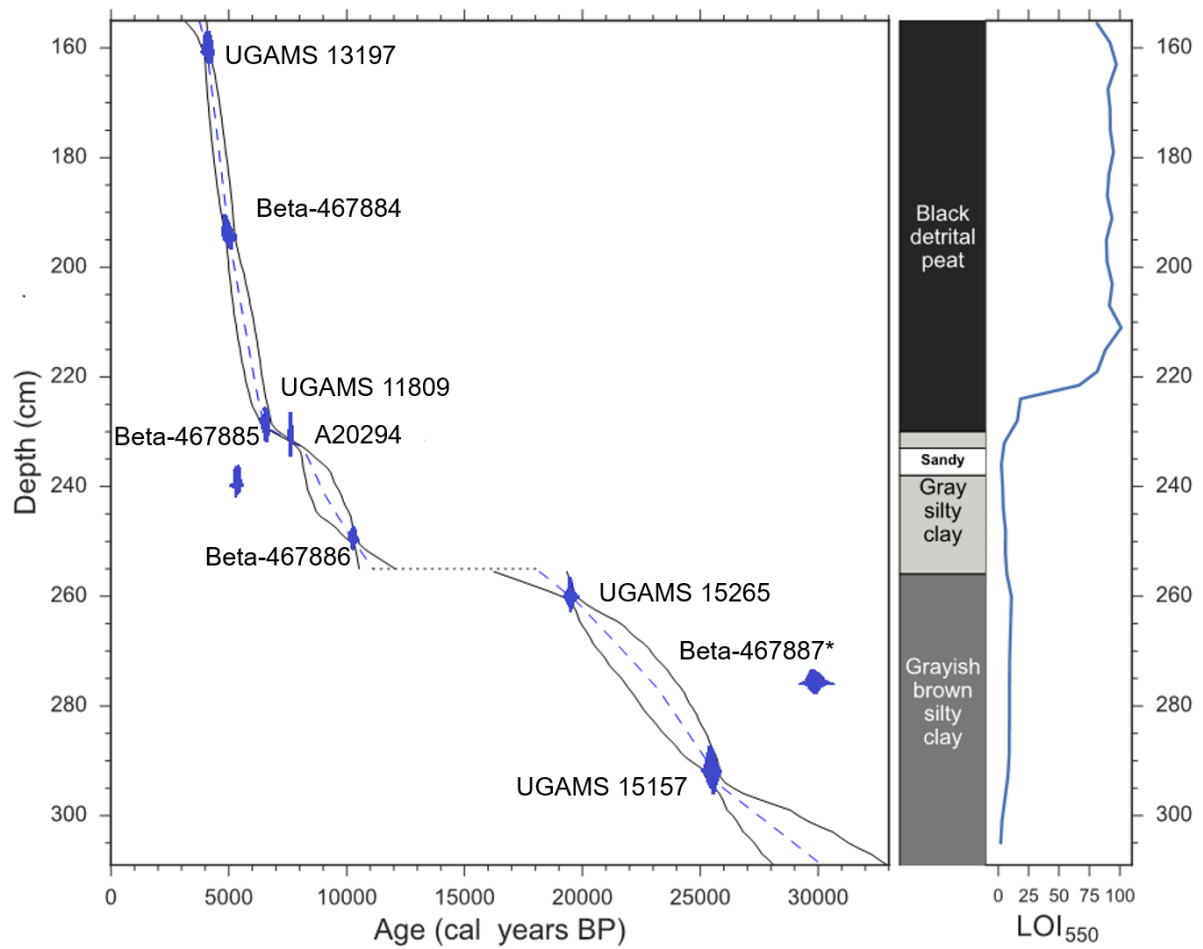
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Figure 1



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Figure 2



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1253 **Figure 3**

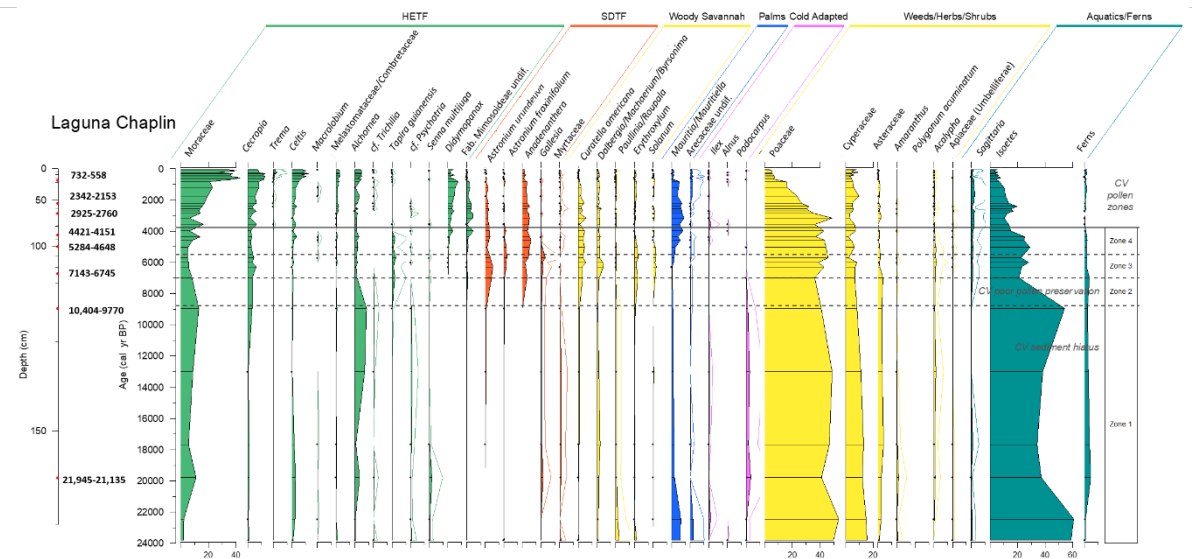
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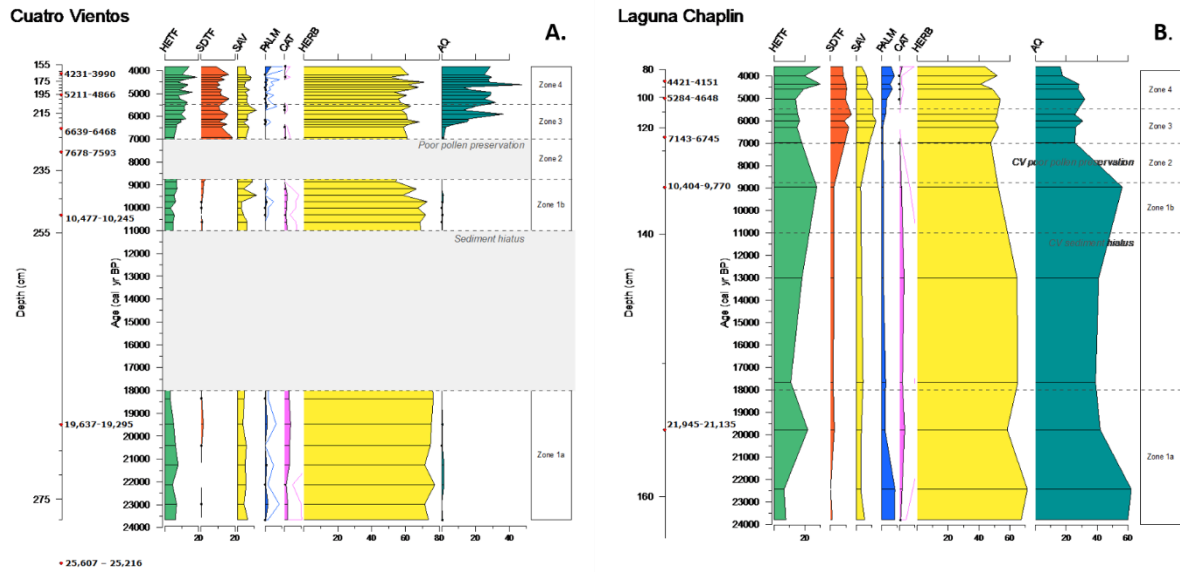
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1259 **Figure 5**

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Figure 6

Table 1 – List of the accelerator mass spectrometry radiocarbon dates from the Cuatro Vientos sediment core

Laboratory code	Sample depth (cm below FMV)	Dated material	AMS ¹⁴ C age (yr BP ± 1σ)	Calibrated age range (cal yr BP) ± 2 σ
UGAMS 13197	161.5	Bulk sediment	3760 ± 25	4231 – 3990
Beta-467884	195	Plant remains	4400 ± 30	5211 – 4866
UGAMS 11809	229	Bulk sediment	5750 ± 30	6639 – 6468
A20294	231.5	Bulk sediment	6800 ± 26	7678 – 7593
Beta-467885*	240	Plant remains	4640 ± 30	5465 – 5307
Beta-467886	250	Bulk sediment	9180 ± 30	10,477 – 10,245
UGAMS 15265	260	Bulk sediment	16,140 ± 40	19,637 – 19,295
Beta-467887*	276	Bulk sediment	25,700 ± 90	30,276 – 29,502
UGAMS 15157	292	Bulk sediment	21,070 ± 50	25,607 – 25,216

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*dates not included in age-depth model

Table 2 – List of the accelerator mass spectrometry radiocarbon dates from the Laguna Chaplin sediment core, taken from Burbridge et al. (2004)

Laboratory code	Sample depth (cm)	Dated material	AMS ¹⁴ C age (yr BP ± 1σ)	Calibrated age range (cal. yr BP) ± 2 σ
Beta-137570	36.5	Bulk sediment	710 ± 50	732 – 558
AA39700	51.5	Bulk sediment	2240 ± 40	2342 – 2153
AA39701	69.5	Bulk sediment	2740 ± 40	2925 – 2760
AA39702	85	Bulk sediment	3870 ± 50	4421 – 4151
AA39703	100	Bulk sediment	4330 ± 80	5284 – 4648
AA39704	125	Bulk sediment	6040 ± 50	7143 – 6745
AA39705	135	Bulk sediment	9000 ± 100	10404 – 9770
AA39706	155	Bulk sediment	17820 ± 140	21945 – 21135
AA39707	175	Bulk sediment	31060 ± 440	35941 – 34190
AA39708	195	Bulk sediment	34820 ± 700	41160 – 37945
AA39709	213	Bulk sediment	37750 ± 970	43801 – 40455
AA39710*	250	Bulk sediment	43400 ± 1900	-
AA39711*	285	Bulk sediment	41200 ± 1400	-
AA39712*	296	Bulk sediment	38100 ± 1000	-

1272 *dates not included in age-depth model

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