

Vegetation and hydrology changes in Eastern Amazonia inferred from a pollen record

MAURO B. DE TOLEDO¹ and MARK B. BUSH²

¹Departamento de Geologia, Universidade Federal Fluminense (UFF),
Av. Gal. Milton Tavares de Souza s/n, 4° andar, Gragoatá, 24210-346 Niterói, RJ, Brasil
²Department of Biological Sciences, Florida Institute of Technology,
150 W. University Blvd. Melbourne, Florida 32901, USA

Manuscript received on September 19, 2006; accepted for publication on March 23, 2007; presented by ALCIDES N. SIAL

ABSTRACT

Pollen, charcoal, and C14 analyses were performed on a sediment core obtained from Lake Tapera (Amapá) to provide the palaeoenvironmental history of this part of Amazonia. A multivariate analysis technique, Detrended Correspondence Analysis, was applied to the pollen data to improve visualization of sample distribution and similarity. A sedimentary hiatus lasting 5,500 years was identified in the Lake Tapera. Because the timing of the hiatus overlapped with the highest Holocene sea-level, which would have increased the local water table preventing the lake from drying out, it is clear that sea-level was not important in maintaining the lake level. Lake Tapera probably depended on riverine floodwaters, and the sedimentary gap was caused by reduced Amazon River discharge, due to an extremely dry period in the Andes (8,000-5,000 years BP), when precipitation levels markedly decreased. The lack of Andean pollen (river transported) in the record after this event supports this interpretation. The pollen analysis shows that when sedimentation resumed in 1,620 cal. years BP, the vegetation around the lake was changed from forest into savanna. This record demonstrates the need to improve our understanding of climate changes and their associated impacts on vegetation dynamics.

Key words: vegetation change, hydrology change, Amazonia, pollen record, climate change.

INTRODUCTION

It is undisputed the large growing body of evidence for climate changes, such as oscillations on mean temperature and precipitation in the geological time. Records of climate and environmental changes obtained from proxies such as ice cores from polar and tropical regions (Grootes et al. 1994, Thompson et al. 1995, 1998, Petit et al. 1999), tree rings (Rodriguez et al. 2005, Sano et al. 2005, Therrell 2005), coral reefs (Dunbar et al. 1996, Gagan et al. 2000), diatoms (Steinitz-Kannan et al. 1993, Jenny et al. 2002, Fritz et al. 2004) and fossil pollen (Absy 1979, Absy et al. 1991, Behling 1995, Van der Knaap and Van Leeuwen 1997, De Oliveira et

Correspondence to: Mauro B. de Toledo

E-mail: mtoledo@igeo.uff.br

al. 1999, Haberle and Maslin 1999, Behling et al. 2004, 2005, Garcia et al. 2004), indicate that these changes took place within decadal to century time scales, and affected both relatively small geographic scales (e.g. the Little Ice Age in Europe) and entire hemispheres (e.g. the last glacial period).

Holocene climatic changes have been invoked to account for vegetation changes worldwide, and the variation in the range of Amazonian savannas is not an exception. A paleoecological study conducted on a mosaic landscape of forest-savanna ecotone in Bolivia revealed a forest expansion of c. 100 km in the last 3000 years (Mayle et al. 2000). Similarly, palaeoecological records from Colombian savannas also suggest a forest expansion in the last 4000 years (Behling and Hooghiemstra 1998, 1999, Berrio et al. 2002).

Even though climate change may be one of the most powerful drivers of modifications in the environment and vegetation, the anthropogenic impacts of occupation in the Amazonian region cannot be dismissed. Archaeological studies provided evidence that humans have lived in Amazonian lowlands for at least 11,000 years (Roosevelt et al. 1991, 1996). Further evidence that these Neotropical landscapes have been intensely used and modified by pre-Columbian human populations comes from savannas in the Bolivian Amazonia (Erickson 2000, 2001) and Upper Xingu (Heckenberger et al. 2003), in Southern Brazilian Amazonia. However, the full extent of the Pre-European manipulation of Amazonian habitats remains unclear (Meggers 2003).

The main goal of this study was to investigate how the vegetation of this portion of Amazonia responded to climate changes and human impacts during the Holocene. For this purpose, a sediment core was systematically analyzed for its pollen contents and micro-charcoal fragments in order to provide a record of vegetation dynamics and palaeo-fires (natural or anthropogenic in origin).

STUDY SITE

Lake Tapera (N 0°07′40.9" / W 51°04′47.8") is a small (1 km long and 700 m wide) shallow lake (1.5-2.5 m deep), located at north of Macapá, about 10 km from the Amazon River, and lying at less than 10 m above sea level (Fig. 1). This lake drains into a larger system, known as Curiaú, therefore being part of its watershed, but presumably a little higher and consequently more isolated. Lake Curiaú is an L-shaped basin 6.5 km long in the vertical axis (N-S), 3.5 km long in the horizontal axis (E-W), and 1.2 km wide; its watershed area is estimated to be at least 150 km². Even though Curiaú dries out seasonally, Lake Tapera is known to be a permanent water body.

The modern vegetation of the study site is a mosaic of dry and flooded savannas and patches of secondary and dry forests. Gallery forests and palm swamps are also present along the water drainages (locally called "igarapés"). Lake Tapera is surrounded by semi-deciduous and gallery forests, with *Mauritia* palms and *Montrichardia* growing on the margins as well. In a phytosociological study in this savanna region, Thomaz et al. (2004) identified 44 families, 83 genera and 119 species. The

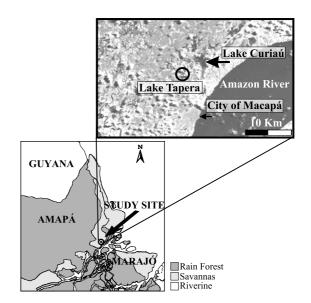


Fig. 1 – Map of the study area, showing the location of Lakes Curiaú and Tapera (the studied lake), the city of Macapá, and the Amazon River.

most important families were Cyperaceae (18 species), Poaceae (15 species), Fabaceae (7 species), Rubiaceae (3 species), Lentibulariaceae and Onagraceae (5 species each). It is noteworthy to mention that among the dominant species were *Sagittaria rhombifolia*, *Montrichardia arborescens* and *Mauritia flexuosa*, and the latter ones are found in isolated populations throughout the Curiaú watershed.

The climate in this region is tropical humid, with mean annual temperatures of 25-27°C and 2500 mm of precipitation, falling mainly between December-August with a dry season from September to November (IBGE 2002).

MATERIALS AND METHODS

A sediment core (200 cm long) was obtained in 2002 using a Colinvaux-Vohnout corer (Colinvaux et al. 1999) from a platform supported by inflatable boats. The platform was anchored in the middle of the lake. The sealed core tubes were transported unopened to the laboratory and stored in a dark cold room until opened and the sediments described. A total of 79 samples (0.5 cm³), 39 for pollen and 40 for charcoal, were collected from the sediment core. Samples for C¹4 AMS dating were sent to the INSTAAR – AMS Radiocarbon Laboratory

at the University of Colorado at Boulder, and the resulting dates were calibrated to calendar years using the software CALIB 4.0 (Stuiver and Reimer 1993).

Standard pollen preparation procedures with HCl, KOH, HF, and acetolysis followed Faegri and Iversen (1989). Tablets of Lycopodium spores were added to the samples prior to processing, in order to calculate pollen concentrations (Stockmarr 1971). The pollen residues were mounted in glycerol and counts of at least 300 grains were conducted at $400\times$ and $1000\times$ magnification on a Zeiss Axioskop. Pollen grains were identified using the Florida Institute of Technology reference collection of modern pollen and published catalogs with photographs and morphological descriptions of pollen types (Hooghiemstra 1984, Roubik and Moreno 1991, Colinvaux et al. 1999).

The pollen sum, which included only the terrestrial taxa, pollen percentage and concentration were all calculated in TILIA (Grimm 1992). Afterwards, all the data were plotted in C^2 1.3 (Juggins 2003).

Samples for charcoal analysis were disaggregated in KOH (10%), and the resulting slurry washed through a 170 μ m sieve (particles > 170 μ m were retained). The charcoal analysis was performed under an Olympus dissection microscope (20× magnification) equipped with a video camera (Clark and Patterson 1997). Digital measurements were made via the video system and image recognition (NIH-IMAGE) software. This software provides the area of charcoal particles according to the number of pixels occupied by the fragment on the screen.

In order to investigate the complex relationships among the samples, Detrended Correspondence Analysis – DCA (Hill and Gauch 1980) was performed using PC-ORD 4.0 (McCune and Mefford 1999). The main goal of DCA is to reduce the number of dimensions, with a minimum loss of information, of a multidimensional space that describes the variables relationship (McCune and Grace 2002). Through the use of these more advanced numerical techniques, we are able to observe the most relevant of a possibly infinite number of patterns in the ecological data, identifying the similarity, association, and correlation among all the variables (Behling et al. 2005).

This technique was chosen basically because it is not subject to the infamous arch effect. Additionally, as very complex data sets with many rare taxa can hinder DCA, the data set was reduced to include only pollen taxa with percentage values equal to or higher than 1% and that were present in at least 3 samples throughout the sediment core. The percentage values were then standardized by square root transformation (McCune and Grace 2002). Because the eigenvalues provided by DCA cannot be used to demonstrate the proportion of variation explained by the axes, as in Principal Component Analysis (PCA), we used an after-the-fact coefficient of determination between relative Euclidian distance in the unreduced species space and Euclidian distance in the ordination space (McCune and Grace 2002) in order to evaluate the effectiveness of the ordination. The resulting sample scores from axis 1 were plotted against sample depths.

RESULTS

LITHOLOGY

The 200 cm-long sediment core from Lake Tapera (Fig. 2) is mostly composed of blue-grey clay rich in plant fragments in the bottom (200-65 cm) and black silt with no plant remains on the top 65 cm. In the bottom of the core (197-200 cm) there is a layer of dark-blue clay rich in black organic fragments, and between 69-76 cm depth the clay is very sandy (Table I).

Lake Tapera

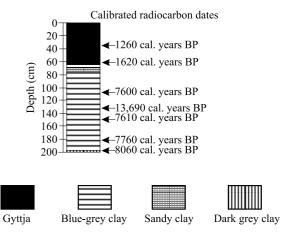


Fig. 2 – Lithology of sediment core from Lake Tapera (Amapá). Also showing location of dated samples on the sediment core (cal. years

TABLE I Sediment description of core from Lake Tapera (Amapá – Brazil).

| Depth (cm) | Sediment description | | |
|------------|--|--|--|
| 0–65 | Black organic silt, no plant remains present | | |
| 65–69 | Blue-gray clay rich in plant remains | | |
| 69–76 | Very sandy blue-gray clay | | |
| 76–197 | Blue-gray clay with some plant remains | | |
| 197–200 | Dark clay rich in charcoal fragments | | |

RADIOCARBON DATES

The sedimentary chronology for Lake Tapera was derived from 7 AMS radiocarbon dates (Table II). In order to account for variations in carbon isotopic fractionation throughout time, the resulting ages were calibrated to calendar years using the software CALIB 4.0 (Stuiver and Reimer 1993). For each calibrated age a maximum probability solution (type 2 solution) was adopted. All ages used henceforth will be interpolated (rounded up) calibrated years (cal. yrs) before present (BP), unless specifically noted otherwise. The basal age of 8,060 cal. yrs BP for Lake Tapera indicates that the sediments are Holocene in age. The age vs. depth relationship (Fig. 3) shows that sedimentation rates were not constant through time. As the age of 13,480-13,890 cal. yrs BP, obtained from sample OS-38390 at 132 cm depth (Table II), was probably originated from erosion of much older sediments from elsewhere, it was rejected because it does not represent the true time of deposition of the sedimentary layers.

PALAEOECOLOGICAL RECORD

The pollen and charcoal records from Lake Tapera (Fig. 4) allow the distinction of 2 main zones: zone T1, which lasts from c. 8,060 until before 1,620 cal. yrs BP; and zone T2 lasting from \sim 1620 cal. yrs BP until the present. The diagram zonation was based both on DCA scores and visual inspection.

T1A (200 cm-65 cm; 8,060-1,620 cal. yrs BP)

Sediments are composed of dark clay rich in fragments of charred wood on the bottom, and blue grey clay with plant fragments. Pollen concentration displays strong fluctuations, peaking at 190 cm (c. 32,500 grains cm⁻³ of

sediment), 165 cm and 150 cm (around 15,000 grains cm⁻³ of sediment), and decreasing to almost nothing upwards (Fig. 4). This zone lasts from 8,060 cal. yrs BP until c. 1,620 cal. yrs BP, and is characterized by low percentages of Asteraceae, Bignoniaceae, Cassia, Cecropia, Dilleniaceae, Euterpe, Mimosaceae, Macrolobium, Mauritia, Combretaceae type (Mel/Comb), Moraceae/Urticaceae 2-porate and 3-porate types (Mor/ Urt2, Mor/Urt3), Poaceae, Polygonum, Spondias, and Cyperaceae. These pollen types show higher values upwards. On the other hand, pollen of Coccoloba, Desmodium, Didymopanax, Lecythidaceae, Caesalpiniaceae, Papillionaceae, Machaerium, Malpighiaceae, Protium, Symphonia, Virola, and Rhizophora display higher percentage values in this zone (Fig. 4). Even though the fern spores display relatively small and constant percent values throughout the core, there is one distinct peak at 75 cm (128%). Pollen commonly found in mountain regions (e.g. in the Andes), such as Alnus, Hedyosmum, and Podocarpus are recorded with low percentage values in this zone. No charcoal particles were found in this zone.

T2 (65 cm-0 cm; c. 1,620 to present)

Sediments are composed of a layer of sandy blue-grey clay on the bottom, a transition from blue-grey clay rich in plant fragments to black organic silt with no plant fragments. Pollen concentration increases sharply and the values fluctuate around 10,000-20,000 grains cm⁻³ of sediment (Fig. 4). Pollen spectra are characterized by decreased percentage values of Asteraceae, Bignoniaceae, Cassia, Coccoloba, Desmodium, Didymopanax, Lecythidaceae, Caesalpiniaceae, Machaerium, Malpighiaceae, Protium, Rubiaceae, Symphonia, and Virola. Other taxa display increased percentages, such as Euterpe, Macrolobium, Mauritia, Mel/Comb, Polygonum, Spondias, Sagittaria, Typha, and Utricularia. The percentage of Poaceae pollen exhibits a 5-fold increase in this zone, and despite some fluctuation remains abundant. Cecropia pollen reaches its highest values at 55 cm (c. 17%), but does not increase above 10% anywhere else, displaying only small variation throughout this zone (Fig. 4). Pollen from Alnus is completely absent from this zone, and *Hedyosmum* and *Podocarpus* are recorded in only one sample at insignificant values.

| | | | - | |
|----------|------------|-----------------------|--|---------------------|
| Sample | Depth (cm) | ¹⁴ C yr BP | ¹³ C/ ¹² C ratio | Age (cal. years BP) |
| OS-38388 | 35 | 1320 ± 30 | -27.93 | 1230–1300 |
| OS-38389 | 65 | 1710 ± 35 | -27.37 | 1540–1700 |
| OS-40060 | 108 | 6720 ± 30 | -26.72 | 7570–7630 |
| OS-38390 | 132 | $11,750 \pm 55$ | -25.37 | *13,480–13,890 |
| OS-40061 | 150 | 6730 ± 35 | -27.86 | 7570–7630 |
| OS-38391 | 182 | 6930 ± 45 | -27.03 | 7670–7850 |
| OS-38392 | 197 | 7260 ± 35 | -23.39 | 8000–8120 |

TABLE II Radiocarbon dates from Lake Tapera.

^{*}Date rejected as an outlier.

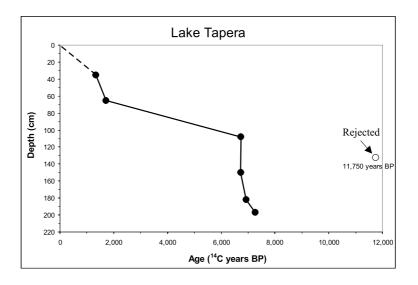


Fig. 3 – Radiocarbon ages from Lake Tapera plotted against depth (cm). The radiocarbon age that is represented by an open circle (11,750 years BP) was considered to be a contamination, and therefore rejected as an outlier. The dashed lines represent the extrapolated ages towards the present.

The accumulation of charcoal particles, measured here as $\rm mm^2~cm^{-3}$ of sediment, starts at 65 cm (1,620 cal. yrs BP), and displays a peak of c. 15 mm² at 60 cm. At the depth of 35 cm (1,258 cal. yrs BP) charcoal area is greatly decreased, and values remain small until the end of the zone.

MULTIVARIATE ANALYSIS

The resulting DCA scores from Lake Tapera were also plotted in two different ways. First, axis 1 *vs.* axis 2, then the sample scores from axis 1 were plotted with the corresponding depths (Fig. 5). The proportion of

variance explained by the first two axes, as given by the after-the-fact coefficient of determination, was 80%.

Axis 1 vs. Axis 2

The DCA polarized samples on axes 1 and 2 so that the bottom samples of the core are on the positive side of axis 1, representing zone T1, and the top samples of the core on the left side representing zone T2.

The highest scores for axis 1 were yielded by *Pachi-ra*, Bignoniaceae, Malpighiaceae, *Symphonia*, and Euphorbiaceae. The species that scored lowest in this axis, and therefore were responsible for samples being placed

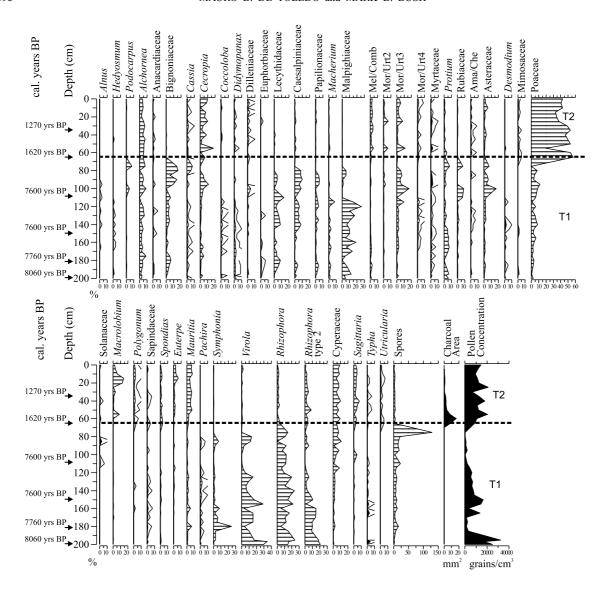


Fig. 4 – Pollen percentage diagram of Lake Tapera, Amapá (Brazil), showing the most representative taxa, including Andean taxa, area of charcoal particles (mm²/cm³ of sediment), pollen concentration (grains/cm³ of sediment), and radiocarbon dates (cal. years BP). The hollow curves are exaggerated 5 times.

on the more negative side of axis 1 on zone T2, were *Utricularia*, *Polygonum*, *Sagittaria*, Mimosaceae, and *Macrolobium*.

On axis 2 samples from zone T1 (bottom of the core) were subdivided into two groups. The species that scored highest in this axis were *Coccoloba*, *Typha*, *Desmodium*, *Didymopanax*, and *Virola*. The lowest species scores were yielded by Solanaceae, Bignoniaceae, Rubiaceae, Lecythidaceae, Papillionaceae, Asteraceae, Cyperaceae, Caesalpiniaceae, Myrtaceae, and *Symphonia*.

Axis 1 vs. Depth

Plotting axis 1 DCA sample scores against sample depth revealed a striking pattern, in which a transition between zones T1 and T2 occurred at 70 cm (between c. 7,600 cal. yrs BP and 1,618 cal. yrs BP). In zone T1, samples were placed on positive side of axis 1 and showed small oscillation upwards, with the exception of sample T-100 (100 cm) that scored much lower values. In zone T2, from 70 cm until 0 cm (before 1,620 cal. yrs BP until the present), samples present a smaller range of variation

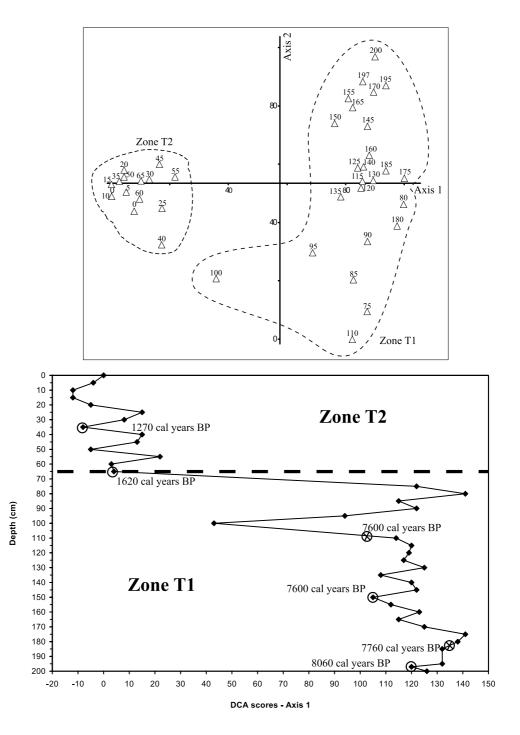


Fig. 5 – DCA scores from Lake Tapera. Top graph shows axis 1 vs. axis 2. Samples are encircled to display zones as interpreted from pollen diagrams. Bottom graph represents resulting DCA scores of axis 1 plotted vs. depth (cm) showing a strong polarization of samples into two main groups representing zones T1 and T2. Ages obtained from C^{14} AMS are also shown.

than in the bottom of the core, and were placed on the negative side of the axis.

DISCUSSION

SEDIMENTARY EVIDENCE

Stratigraphy alone indicates that the core from Lake Tapera provides a record of an environmental change, as it shows blue-grey clay in the bottom being replaced by black-organic sediments upwards (Fig. 2). The sharp shift of the sediments implied that this change was at least locally (basin scale) significant and may have been even abrupt. The blue-grey clay that constitutes the bottom of the sediment core was full of plant remains, wood and fragments of charred wood, indicating a shallow and unstable depositional environment. The coring location may have supported a gallery forest subjected to periodic river flooding, which may explain the presence of wood and plant fragments. The sandy layer at 69-76 cm in the core was another indication of environmental instability, suggesting drought and maybe associated erosion of nearby sediments.

Sedimentation rate was the fastest in the basal portion of the core, slowing down upwards (Fig. 3). Slowing of sedimentation in the middle of the core was so extreme that it almost certainly represents a long gap, or a series of depositional hiatuses that resulted in little net sedimentary accumulation between c. 7,610 and 1,620 cal. yrs BP.

The sharp change in sediment type, combined with a rapid change in community composition, and different sediment depositional rates below and above the boundary (65 cm) suggest the occurrence of a sedimentary hiatus in the lake. Such breaks in sedimentation are common in shallow tropical lakes (C. Listopad, unpublished data, Bush et al. 2007a) and are generally taken to indicate conditions in which lake water level fell.

Duration of the sedimentary hiatus at Tapera, given by the difference between interpolated ages estimated through sedimentation rates from the bottom upwards and from the top downwards is \sim 5, 500 years. From this observation, it is possible to infer that lake level fell during this time interval and was reestablished as a permanent water body only at c. 1,620 cal. yrs BP. With the flooding of Tapera, deposition of organic-rich mud initiated.

PALEOECOLOGICAL EVIDENCE

The pollen and charcoal records from Lake Tapera revealed two main contrasting zones. The most obvious biological distinction between them was the increase in Poaceae pollen, rising from < 10% in T1 to almost 60% (40-55%) in T2. Interestingly, these percentage values are comparable with proportions of Poaceae pollen (50-90%) found in the pollen rain of cerrados (Salgado-Labouriau 1973), suggesting the establishment of savanna vegetation.

A further indication of a significant vegetation change was the loss of *Rhizophora* pollen in zone 2. The declining proportion of *Rhizophora* pollen suggests a weakening of the fluvial-marine influence. Possibly, as the lake became more isolated from the Amazon, and fresher. Replacement of forest taxa, e.g. Bignoniaceae, Caesalpiniaceae, Cassia, Lecythidaceae, Machaerium, Malpighiaceae, Protium, Symphonia, and Virola (Marchant et al. 2002) in T2 with those from flooded savanna and swamp vegetation, such as Amaranthus/Chenopodiaceae, Asteraceae, Ludwigia, Macrolobium, Euterpe, Mauritia, Polygonum, Cyperaceae, Typha, and Utricularia, is consistent with a shift towards less dense forest. Additionally, the higher percentage values of Cecropia upwards suggest increased disturbance on the vegetation, as this tree is a well-known pioneer that occupies forest gaps (Marchant et al. 2002).

The sample scores derived from DCA when plotted against core depths provided an illustrative way to demonstrate the impact of environmental changes that took place in this system. Axis 1 was interpreted to show a hydrology gradient, changing from an environment subjected to seasonal river flooding, to a more permanently flooded one, accompanied by a landscape change from gallery forest to swamp forest, with strong influence of savanna. The timing of the environmental change was coincident with on set of fire near the lake. The replacement of *Symphonia* and *Pachira* in T1 by *Macrolobium*, *Polygonum*, *Mauritia* and Cyperaceae in T2 suggested changes within the community of wetland plants.

To sum up, the pollen and charcoal data showed a general environmental trend from gallery forest to savanna with swamp elements. Anthropogenic impacts on the environment were suggested by large quantities of charcoal particles after c. 1,600 cal. yrs BP in Lake

Tapera. Impacts on the vegetation were corroborated by the increasing abundance of *Cecropia* pollen in the upper part of the core.

INFLUENCE OF SEA-LEVEL VARIATION

In the early Holocene, relative sea level started to rise and reached its modern level at approximately 7,000 cal. yrs BP along the northern coast of Brazil (Suguio et al. 1985, Martin et al. 1986, Bezerra et al. 2003). The impacts of rising sea level on the water table changed the dynamics of rivers and also formed lakes in the Amazonian lowlands (Behling 1996, 2002 Behling and Da Costa 2000, 2001, Behling et al. 2001). The initial ponding of Tapera at \sim 8, 060 cal. years BP was probably induced by rapid sea level rise in the Pleistocene-Holocene boundary.

At c. 5,000 cal. yrs BP, even though relative sea level was at least 2 meters higher than modern levels (Suguio et al. 1985, Bezerra et al. 2003), Lake Tapera was dry at this time, which suggests that sea level was not the primary control of lake level. Therefore, it is probable that the lake relied on river floodwater to keep a high stand. It is important to note that Lake Tapera is slightly higher than Lake Curiaú, which makes it more isolated from river floodwaters. A further possibility is that the Tapera lake basin was not completely closed yet, so that the lake could be losing water through seepage.

ANDEAN DRY SPELL AND AMAZON RIVER FLOW

A drought or series of droughts influenced both Andean and Amazonian systems in the Mid-Holocene. Lake Titicaca (Peru-Bolivia) presented its lowest lake level in the Holocene at c. 5,500 cal. yrs BP (Baker et al. 2001). Though this drought probably began as early as 9,000 cal. yrs BP, and terminated as late as 3,400 yrs BP (Abbott et al. 1997), with 5,500 cal. yrs BP being the peak of the dry spell. Indeed, analyses of western Amazonian lakes suggest the last 1,000 years as being the wettest period of the Holocene (Bush et al. 2007b). A pollen record from Serra dos Carajás suggests that the interval 8,000-5,000 yrs BP was the driest period of the Holocene (Absy et al. 1991). High dust concentrations in the Sajama (Bolivia) ice core also reflect a period of Andean drought at this time (Thompson et al. 1998). As the climate of the southern tropical Andes, an important hydrological influence on Amazon River discharge was dry between 8,000 and 5,000 years BP, it is probable that the flow of the river was reduced. One of the impacts of this process would have been diminished river flooding, probably both spatially and temporally, during the wet season. That Tapera received floodwater is evident in the input of Andean pollen taxa, such as Podocarpus, Hedyosmum and Alnus. Therefore, as the lake probably relied on floodwaters from the Amazon River, it dried out, even though relative sea level was high at 5,000 yrs BP. This hypothesis is supported by the pollen record of Tapera that shows a relatively greater abundance of Andean pollen before the sedimentary hiatus than after, when sedimentation resumed. Even though some of these pollen types may be found in modern Amazonian forests (e.g. Hedyosmum and Podocarpus), their absence after the interpreted environmental change supports the assumption that they were river transported instead of airborne.

Although the climate change at the Andes was probably the main factor influencing the isolation of Lake Tapera, local sedimentary processes, such as the drowning of the Curiaú paleo-drainage system, probably played an important role. As relative sea level was high, and the Amazon flow was reduced, it is possible that a build up of sediments at the mouth of the Amazon and within the estuary induced a "coastal progradation" cutting the Curiaú hydrologic system, intensifying the effects of isolation from riverine floodwater, already experienced by the lake. A similar process was described for a mangrove in French Guiana, in which a coastal progradation at ca. 5,000 years BP was proposed to explain the decline in mangrove vegetation in the pollen record of a coastal lake during the peak of the Holocene sea-level highstand (Tissot et al. 1987).

Although the volume of the Amazon River may have increased rapidly, it is improbable that floodwaters were reaching the lake in the same way as before the dry event, especially because of the lowering sea level. The replacement of blue-grey clay by black-organic sediments, indicative of autochthonous sedimentation, and the lack of Andean pollen from the lake record support this interpretation.

Another Mid-Holocene event that may have exerted an influence on this record is the oscillating intensity of the El Niño/Southern oscillation (ENSO). An increase in ENSO frequency, especially La Niña events, after 5,000 cal. yrs BP (Moy et al. 2002, Riedinger et al. 2002) might have caused drought in eastern Amazonia, and could have augmented the inferred pattern. It is probable that long-term shifts in ENSO intensity influenced both the Andes and Amazonia during the Holocene. However, as is not possible to parse out the ENSO signal from that of other factors in this study, we consider the reduction on river discharge and the resulting coastal progradation to be both more proximate and parsimonious explanations of the observed changes at Tapera.

CONCLUSIONS

The fossil pollen and charcoal records from a lake in eastern Amazonia revealed an environmental change that was initiated by a reduction in river flooding, consistent with a well-documented climate change in the central Andes. As the lake was dependent on flooding to maintain its hydrological balance, the reduced input resulted in a sedimentary hiatus lasting ~ 5 , 500 years at Lake Tapera.

The resume of sedimentation, indicative of local wet conditions, did not bring about a return to pre-drought conditions as a combination of factors, such as changed hydrology due to coastal progradation, human occupation and palaeo-fires may all have played a role in deflecting the system onto a new path.

Despite resumption of wet conditions, flooding of the system was never reestablished, consistent with the long-term change in hydrology induced by falling sea levels in the late Holocene. The rise of lake level within the last 2,000 years, even without significant white-water flooding, is consistent with the observation that the late Holocene has been an unusually wet period.

ACKNOWLEDGMENTS

Many thanks are due to several people from Brazil and the institutions they represent: Dr. Paulo Eduardo de Oliveira (Universidade Guarulhos), Dr. Odete F. da Silveira (UFPA), MS. Ronaldo L. Justo. The authors would also like to thank two anonymous reviewers who have contributed for the improvement of this manuscript. Financial support: Conselho Nacional de Desenvolvimento Científico e Tecnológico (CNPq) (200065/

99-8) and National Science Foundation, Division of Environmental Biology (NSF-DEB-9732951).

RESUMO

Análises de pólen, carvões e datações C14 foram conduzidas em um testemunho coletado no lago Tapera (Amapá) com o objetivo de interpretar a história paleoambiental desta parte da Amazônia. Uma das técnicas de análises multivariadas, Análise de Correspondência Destendenciada (DCA), foi utilizada a fim de melhor visualizar a distribuição e similaridade das amostras. Foi identificado um hiato sedimentar com duração de 5.500 anos no lago Tapera. Como o hiato ocorreu simultaneamente ao nível do mar mais alto do Holoceno, o que deveria ter aumentado o lençol freático, impedindo assim o lago de secar, é evidente que variações do nível do mar não foram importantes na manutenção do nível do lago. O lago Tapera provavelmente dependia de água das cheias fluviais, e o hiato sedimentar foi causado por uma redução da descarga do Amazonas, devido a um período extremamente seco nos Andes (8.000-5.000 anos cal. AP), quando os níveis de precipitação diminuíram drasticamente. A ausência de pólen transportado dos Andes pelo rio Amazonas no registro sedimentar, após este evento, apóia esta interpretação. A análise palinológica demonstra que quando a sedimentação reiniciou, em 1.620 anos cal. AP, a vegetação em torno do lago havia mudado de floresta para savana. Este registro demonstra a necessidade de melhorar nosso entendimento sobre a extensão e os impactos das mudanças climáticas sobre a evolução da vegetação.

Palavras-chave: mudança na vegetação, mudança na hidrologia, Amazônia, registro palinológico, mudança climática.

REFERENCES

ABBOTT MB, BINFORD MW, BRENNER M AND KELTS K. 1997. A 3500 14C yr high-resolution record of water-level changes in Lake Titicaca, Bolivia/Peru. Quat Res 47: 169–180.

ABSY ML. 1979. A palynological study of Holocene sediments in the Amazon basin. University of Amsterdam, Amsterdam, 86 p.

ABSY ML, CLIEF A, FOURNIER M, MARTIN L, SERVANT M, SIFEDDINE A, FERREIRA DA SILVA M, SOUBIÈS F, SUGUIO KT AND VAN DER HAMMEN T. 1991. Mise en évidence de quatre phases d'ouverture de la forêt dense dans le sud-est de L'Amazonie au cours des 60,000 dernières années. Première comparaison avec d'autres ré-

- gions tropicales. Compt Rend Acad des Sci Paris, Series II 312: 673–678.
- BAKER PA, SELTZER GO, FRITZ SC, DUNBAR RB, GROVE MJ, TAPIA PM, CROSS SL, ROWE HD AND BRODA JP. 2001. The history of South American tropical precipitation for the past 25,000 years. Science 291: 640–643.
- BEHLING H. 1995. A high resolution Holocene pollen record from Lago do Pires, SE Brazil: vegetation, climate and fire history. J Paleolim 14: 253–268.
- BEHLING H. 1996. First report on new evidence for the occurrence of Podocarpus and possible human presence at the mouth of the Amazon during the late-glacial. Veg Hist Archaeobot 5: 241–246.
- Behling H. 2002. Impact of the Holocene sea-level changes in coastal, eastern, and central Amazonia. Amazoniana 17: 41–52.
- Behling H and da Costa ML. 2000. Holocene environmental changes from the Rio Curuá record in the Caxiuanã region, eastern Amazon Basin. Quat Res 53: 369–377.
- BEHLINGH AND DA COSTA ML. 2001. Holocene vegetational and coastal environmental changes from the Lago Crispim record in northeastern Pará State, eastern Amazonia. Rev Palaeo Pal 114: 145–155.
- BEHLING H AND HOOGHIEMSTRA H. 1998. Late Quaternary paleoecology and paleoclimatology from pollen records of the savannas of the Llanos Orientales in Colombia. Palaeo Palaeo Palaeo 139: 251–267.
- BEHLING H AND HOOGHIEMSTRA H. 1999. Environmental history of the Colombian savannas of the Llanos Orientales since the Last Glacial Maximum from lake records El Pinal and Carimagua. J Paleolim 21: 461–476.
- BEHLING H, KEIM G, IRION G, JUNK W AND NUNES DE MELLO J. 2001. Holocene environmental changes in the central Amazon Basin inferred from Lago Calado (Brazil). Palaeo Palaeo Palaeo 173: 87–101.
- BEHLING H, PILLAR VD, ORLÓCI L AND BAUERMANN SG. 2004. Late Quaternary Araucaria forest, grassland (Campos), fire and climate dynamics, studied by high-resolution pollen, charcoal and multivariate analysis of the Cambará do Sul core in southern Brazil. Palaeo Palaeo Palaeo 203: 277–297.
- BEHLING H, PILLAR VD AND BAUERMANN SG. 2005. Late Quaternary grassland (Campos), gallery forest, fire and climate dynamics, studied by pollen, charcoal and multivariate analysis of the São Francisco de Assis core in western Rio Grande do Sul (southern Brazil). Rev Palaeo Pal 133: 235–248.

- BERRIO JC, HOOGHIEMSTRA H, BEHLING H, BOTERO P AND VAN DER BORG K. 2002. Late Quaternary savanna history of the Colombian Llanos Orientales from Lagunas Chenevo and Mozambique: a transect synthesis. The Holocene 12: 35–48.
- BEZERRA FHR, BARRETO AMF AND SUGUIO K. 2003. Holocene sea-level history on the Rio Grande do Norte State coast, Brazil. Mar Geol 96: 73–89.
- BUSH MB, SILMAN MR AND LISTOPAD CMCS. 2007a. A regional study of Holocene climate change and human occupation in Peruvian Amazonia. J Biogeogr 34: 1342–1356.
- BUSH MB, SILMAN MR, DE TOLEDO MB, LISTOPAD C, GOSLING WD, WILLIAMS C, DE OLIVEIRA PE AND KRISEL C. 2007b. Holocene fire and occupation in amazonia: Records from two lake districts. Philos T Roy Soc B 362: 209–218.
- CLARK JS AND PATTERSON WAI. 1997. Background and local charcoal in sediments: Scales of fire evidence in the Paleorecord. Sediment Records of Biomass Burning and Global Change. NATO ASI Series. Series 1: Global Environmental Change vol. 51. (CLARK JS, CACHIER H, GOLDAMMER JG AND STOCKS B (Eds)), Springer, Berlin, p. 23–48.
- COLINVAUX PA, DE OLIVEIRA PE AND MORENO JE. 1999. Amazon Pollen Manual and Atlas. Harwood Academic Press, New York, 332 p.
- DE OLIVEIRA PE, BARRETO AMF AND SUGUIO K. 1999. Late Pleistocene/Holocene climatic and vegetational history of the Brazilian caatinga: the fossil dunes of the middle Sao Francisco River. Palaeo Palaeo Palaeo 152: 319–337
- DUNBAR RB, LINSLEY BK AND WELLINGTON GJ. 1996. Eastern Pacific corals monitor El Niño/Southern Oscillation, precipitation, and sea-surface temperature variability over the past three centuries. Climatic variations and forcing mechanisms over the last 2000 years (JONES PD, BRADLEY RS AND JOUZEL J (Eds)), Springer, Berlin, p. 375–407.
- ERICKSON CL. 2000. An artificial landscape-scale fishery in the Bolivian Amazon. Nature 408: 190–193.
- ERICKSON CL. 2001. Pre-Columbian roads of the Amazon. Expedition 43: 21–30.
- FAEGRI K AND IVERSEN J. 1989. Textbook of Pollen Analysis, 4th ed., FAEGRI K, KALAND PE, KRZYWINSKI K (Eds), J Wiley & Sons, New York, 328 p.
- FRITZ SC, BAKER PA, LOWENSTEIN TK, SELTZER GO,

- RIGSBY CA, DWYER GS, TAPIA PM, ARNOLD KK, KU TL AND LUO S. 2004. Hydrologic variation during the last 170,000 years in the southern hemisphere tropics of South America. Quat Res 61: 95–104.
- GAGAN MK, AYLIFFE LK, BECK JW, COLE JE, DRUFFEL ERM, DUNBAR RB AND SCHRAG DP. 2000. New views of tropical paleoclimates from corals. Qua Sci Rev 19: 45–64.
- GARCIA MJ, DE OLIVEIRA PE, SIQUEIRA E AND FERNANDES RS. 2004. A Holocene vegetational and climatic record from the Atlantic rainforest belt of coastal State of São Paulo, SE, Brazil. Rev Palaeo Pal 131: 181–199.
- GRIMM E. 1992. TILIA Software, Version 1.12., Illinois State University, Springfield.
- GROOTES PM, STEIG EJ, STUIVER M, WADDINGTON ED AND MORSE DL. 1994. A new ice core record from Taylor Dome, Antarctica. EOS Transactions, p. 75–225.
- HABERLE SG AND MASLIN MA. 1999. Late Quaternary vegetation and climate change in the Amazon basin based on a 50,000 year pollen record from the Amazon fan, PDP site 932. Quat Res 51: 27–38.
- HECKENBERGER MJ, KUIKURU A, KUIKURU UT, RUSSELL JC, SCHMIDT M, FAUSTO C AND FRANCHETTO B. 2003. Amazonia 1492: pristine forest or cultural parkland? Science 301: 1710–1714.
- HILL MO AND GAUCH HG. 1980. Detrended correspondence analysis: an improved ordination technique. Vegetatio 42: 47–58.
- HOOGHIEMSTRA H. 1984. Vegetational and climatic history of the high plain of Bogota, Colombia: A continuous record of the last 3.5 million Years, Gantner Verlag, Vaduz, 368 p.
- IBGE. 2002. Mapa de Clima do Brasil. Instituto Brasileiro de Geografia e Estatística, Rio de Janeiro, RJ, Brasil.
- JENNY B, VALERO-GARCÉS BL, VILLA-MARTÍNEZ R, URRUTIA R, GEYH M AND VEIT H. 2002. Early to Mid-Holocene Aridity in Central Chile and the Southern Westerlies: The Laguna Aculeo Record (34°S). Quat Res 58: 160–170.
- JUGGINS S. 2003. C2 User guide. Software for ecological and palaeoecological data analysis and visualisation. University of Newcastle, Newcastle upon Tyne.
- MARCHANT R ET AL. 2002. Distribution and ecology of parent taxa of pollen lodged within the Latin American Pollen Database. Rev Palaeo Pal 121: 1–75.
- MARTIN L, MÖRNER NA, FLEXOR JM AND SUGUIO K.

- 1986. Fundamentos e reconstruções de antigos níveis marinhos do Quaternário. Bol Geo USP 4: 1–161.
- MAYLE FE, BURBRIDGE R AND KILLEEN TJ. 2000. Millennial-scale dynamics of southern Amazonian rain forests. Science 290: 2291–2294.
- MCCUNE B AND GRACE JB. 2002. Analysis of Ecological Communities, MJM, Gleneden Beach, Oregon, 300 p.
- MCCUNE B AND MEFFORD MJ. 1999. PC-ORD. Multivariate Analysis of Ecological Data, MjM Software, Gleneden Beach, Oregon.
- MEGGERS BJ. 2003. Revisiting Amazonia circa 1492. Science 301: 2067.
- MOY CM, SELTZER GO, RODBELL DT AND ANDERSON DM. 2002. Variability of El Niño/Southern Oscillation activity at millenial timescales during the Holocene epoch. Nature 420: 162–165.
- PETIT JR, JOUZEL J, RAYNAUD D, BARKOV NI, BARNOLA J-M, BASILE I, BENDER M AND CHAPPELLAZ J. 1999. Climate and atmospheric history of the past 420,000 years from the Vostok ice core, Antarctica. Nature 399: 429–436.
- RIEDINGER MA, STEINITZ-KANNAN M, LAST WM AND BRENNER M. 2002. A 6100 14C yr record of El Niño activity from the Galapagos Islands. J Paleolim 27: 1–7.
- RODRIGUEZ R, MABRES A, LUCKMAN B, EVANS M, MASIOKAS M AND EKTVEDT TM. 2005. "El Niño" events recorded in dry-forest species of the lowlands of northwest Peru. Dendrochronologia 22: 181–186.
- ROOSEVELT AC, HOUSLEY RA, SILVEIRA MI, MARANCA S AND JOHNSON R. 1991. Eighth millennium pottery from a prehistoric shell midden in the Brazilian Amazon. Science 254: 1621–1624.
- ROOSEVELT AC ET AL. 1996. Paleoindian cave dwellers in the Amazon: the peopling of the Americas. Science 272: 373–384.
- ROUBIK DW AND MORENO E. 1991. Pollen and Spores of Barro Colorado Island. Monographs in Systematic Botany, v. 36, Missouri Botanical Garden.
- SALGADO-LABOURIAU ML. 1973. Contribuição à Palinologia dos Cerrados. Acad Bras Cienc (Eds), Rio de Janeiro, RJ, Brasil, 291 p.
- SANO M, FURUTA F, KOBAYASHI O AND SWEDA T. 2005. Temperature variations since the mid-18th century for western Nepal, as reconstructed from tree-ring width and density of Abies spectabilis. Dendrochronologia 23: 83–92.

- STEINITZ-KANNAN M, NIENABER M, RIEDINGER M AND KANNAN R. 1993. The fossil diatoms of lake Yambo, Ecuador. A Possible Record of El Niño Events. Bull Inst Fr Etudes Andines 22: 227–241.
- STOCKMARR J. 1971. Tablets with spores used in absolute pollen analysis. Pol Spor 13: 615–621.
- STUIVER M AND REIMER PJ. 1993. Extended 14C database and revised CALIB radiocarbon calibration program. Radiocarbon 35: 215–230.
- SUGUIO K, MARTIN L, BITTENCOURT ACSP, DOMINGUEZ JML, FLEXOR J-M AND AZEVEDO AEG. 1985. Flutuações do nível relativo do mar durante o Quaternário Superior ao longo do litoral Brasileiro e suas implicações na sedimentação costeira. Rev Bras Geo 15: 273–286.
- THERRELL MD. 2005. Tree rings and 'El Ano del Hambre' in Mexico. Dendrochronologia 22: 203–207.
- THOMAZ DO, NETO SVC AND TOSTES LCL. 2004. Inventário florístico das ressacas das bacias do Igarapé da Fortaleza e do rio Curiaú. Diagnóstico de ressacas do Estado do Amapá: Bacias do Igarapé da Fortaleza e do rio Curiaú. (TAKIYAMA LR AND SILVA AQ (Eds)), JM Editora Gráfica, Macapá, AP, p. 13–32.

- THOMPSON LG, MOSLEY-THOMPSON E, DAVIS ME, LIN P-N, HENDERSON KA, COLE-DAI J, BOLSAN JF AND LIU KG. 1995. Late glacial stage and Holocene tropical ice core records from Huascaran, Peru. Science 269: 46–50.
- THOMPSON LG ET AL. 1998. A 25,000-year tropical climate history from Bolivian ice cores. Science 282: 1858–1864.
- TISSOT C, DJUWANSAH MR AND MARIUS C. 1987. Evolution de la mangrove en Guyane au cours de l'Holocène étude palynologique. 10th Symposium APLF, Vol. 25 Institut Français Pondicherry, Bordeaux, p. 125–137.
- VAN DER KNAAP WO AND VAN LEEUWEN JFN. 1997. Late Glacial and early Holocene vegetation succession, altitudinal vegetation zonation, and climatic change in the Serra da Estrela, Portugal. Rev Palaeo Pal 97: 239–285.