

**Past climate
variability on fire and
vegetation, cerrádo
savanna ecosystem**

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The effects of past climate variability on fire and vegetation in the cerrádo savanna ecosystem of the Huanchaca Mesetta, Noel Kempff Mercado National Park, NE Bolivia

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Received: 14 November 2014 – Accepted: 1 December 2014 – Published: 30 January 2015

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Published by Copernicus Publications on behalf of the European Geosciences Union.

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1 Introduction

The cerrádo savanna of central South America is the largest, richest, and likely most threatened savanna in the world (Da Silva Meneses and Bates, 2002). The cerrádo is the second largest biome in South America covering 1.86 million km² and is home to over 10 000 plant species (Myers et al., 2000). The tropical forest–savanna ecotones within the cerrádo biome are of considerable interest to biologists because of their high habitat heterogeneity (*beta* diversity), importance in rainforest speciation (Smith et al. 1997) and sensitivity to climate change (IPCC, 2014). According to current estimates however, only 20 % of the cerrádo remains undisturbed and only 1.2 % of the area is preserved in protected areas (Mittermeier et al., 2000). Additionally, cerrádo savannas have a significant role in the modern global carbon cycle because of high CO₂ loss associated with frequent natural fire activity (Malhi et al., 2002). Currently savanna fires are considered the largest source of natural pyrogenic emissions, with the most fire activity of all major global land cover types (Pereira, 2003). In the last few decades, deforestation for agriculture and increased drought have resulted in increased burning in savannas, contributing to approximately 12 % of the annual increase in atmospheric carbon (van der Werf et al., 2010).

The cerrádo biome comprises forest, savanna, and campestre (open field) formations (Abreu et al., 2012; Mistry, 1998). Cerrádo sensu stricto is characterized as a woody savanna formation composed of dense, thin, and rocky outcrops with cerrádo physiognomies that are distinguishable based on their densities, heights, and scattered tree-shrub covers with roughly 50 % trees and 50 % grass (Abreu et al., 2012). The principal determinants of the growth and development of the cerrádo vegetation types are largely related to edaphic factors (Colgan et al., 2012). For example the distribution of major cerrádo vegetation types are closely related to the geomorphology of the Precambrian Brazilian shield in South America (Killeen, 1998a). The development of the variety of cerrádo vegetation communities is largely the result of heterogeneous nature of the edaphic features (Killeen, 1998a) including the depth of the water table,

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such as tundra and deserts, fire is rare, either because of unsuitable climate conditions or lack of biomass to burn. Fire-sensitive ecosystems such as tropical rainforests, are damaged by fire, which disrupts ecological processes that have not evolved with fire (Hardesty et al., 2005). Fire-dependent systems such as the well-drained grasslands of the cerrádo biome, have evolved in the presence of periodic or episodic fires and depend on fire to maintain their ecological processes (Hardesty et al., 2005). Fire-dependent vegetation is fire-adapted, flammable and fire-maintained (Miranda et al., 2009; Pivello, 2011).

The study of fire and vegetation change in the cerrádo is increasingly important as population, agricultural activity, and global warming create pressing management challenges to preserve these biodiverse ecosystems (Mistry, 1998). The long-term role of humans on vegetation and fire regimes of the cerrádo remains unclear. During the Late Holocene (3000 cal yr BP) there is increasing evidence for the increase in *Mauritia flexuosa* and fire activity in Bolivia, Colombia, Venezuela and Brazil that has been attributed to both natural and anthropogenic drivers (Behling and Hooghiemstra, 1999; Berrio et al., 2007; Kahn and de Castro, 1985; Kahn, 1987, 1988; Montoya and Rull, 2011; Rull, 2009; Da Silva Meneses et al., 2013).

To investigate the drivers of vegetation and fire in the cerrádo a long-term perspective is needed. The past few decades have experienced increased global temperatures, increased atmospheric CO₂, and unprecedented levels of deforestation (Malhi et al., 2002). These recent changes heavily influence modern ecological studies, thus limiting the understanding of the role of natural variability in these systems. Long-term paleoecological studies can provide baseline information on processes shaping forest-savanna fire-vegetation dynamics from centennial-to-millennial timescales (Mayle and Whitney, 2012). These long-term studies can inform whether recent shifts in ecotones are the result of a minor short-term oscillation around a relatively stable ecotone or a longer-term (e.g. millennial scale) unidirectional ecotonal shift forced by climate change (Mayle et al., 2000; Mayle and Whitney, 2012). Additionally, long-term paleoecological records help form realistic conservation goals and identify fire

management strategies for the maintenance or restoration of a desired biological state (Willis et al., 2007).

In this study, the long-term paleoecological perspective provides a context for understanding the role of centennial to millennial climate variability in the evolution of fire and vegetation in cerrádo savanna ecosystems. The purpose of this research is to explore long-term environmental change of cerrádo savanna palm swamps in Bolivia from the Late Glacial (ca. 15 000 cal yr BP) to present. Paleoecological proxies including lithology, magnetic susceptibility, loss on ignition, charcoal, stable isotope, and phytolith data are used to investigate long-term ecosystem processes in the cerrádo savanna. There are three primary hypotheses investigated in this study:

1. Edaphic conditions are the dominant control on the presence of savanna versus forest vegetation on the Huanchaca Mesetta.
2. Climate is the dominant control on savanna structure and floristic composition.
3. The Late Holocene rise in *Mauritia flexuosa* was driven by climate rather than a change in human land-use.

1.1 Study site

Noel Kempff Mercado National Park (NKMNP), a 15 230 km² biological reserve in northeastern Bolivia, is located on the Precambrian Shield near the southwestern margin of the Amazon Basin, adjacent to the Brazilian States of Rondônia and Mato Grosso (Burbridge et al., 2004). It is a UNESCO World Heritage Site, in recognition of its globally important biodiversity and largely undisturbed ecosystems, including *terra firme* (non-flooded) evergreen rainforest, riparian and seasonally-flooded humid evergreen forest, seasonally flooded savanna, wetlands, upland cerrádo savannas, and semi-deciduous dry forests (Mayle et al., 2007). NKMNP occupies an ecotone between Amazon rainforest to the north and dry forests and savannas to the south, containing 22 plant communities (Fig. 1) (Burn et al., 2010). Huanchaca Mesetta palm swamp

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(14°32'10.66" S, 60°43'55.92" W, elevation: 800 m.a.s.l.) is located within NKMNP on the Huanchaca Mesetta – an 800–900 m elevation table mountain. The palm swamp is approximately 200 by 50 m, comprised entirely of a mono-specific stand of the palm *Mauritia flexuosa*.

1.2 Climate

The climate of NKMNP is characterized by a tropical wet and dry climate (Da Silva Meneses and Bates, 2002). The mean annual precipitation at NKMNP derived from nearby weather stations (Concepción, Magdalena, San Ignacio) is ca. 1400–1500 mm per year, with mean annual temperatures between 25 and 26 °C (Hanagarth, 1993; Montes de Oca, 1982; Roche and Rocha, 1985). There is a three to five month dry season during the Southern Hemisphere winter (May–September–October), when the mean monthly precipitation is less than 30 mm (Killeen, 1990). Precipitation falls mainly during the austral summer (December–March), originating from a combination of deep-cell convective activity in the Amazon Basin from the South American Summer Monsoon (SASM) and the Intertropical Convergence Zone (ITCZ) (Vuille et al., 2012). The SASM transports Atlantic moisture into the basin and corresponds to the southern extension of the ITCZ. The ITCZ is driven by seasonal variation in insolation; thus, maximum Southern Hemisphere insolation and precipitation occur in the austral summer (Bush and Silman, 2004; Vuille et al., 2012). During winter (June–August), cold, dry polar advections from Patagonia, locally known as *surazos*, can cause short-term cold temperatures to frequently decrease down to 10 °C for several days at a time (Latrubesse et al., 2012; Mayle and Whitney, 2012). These abrupt decreases in temperature may potentially influence the distribution of temperature-limited species on the Huanchaca Mesetta.

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1.3 Geomorphology

The Huanchaca Mesetta table mountain is near the western limit of the Brazilian Shield and dominates the eastern half of NKMNP. It is composed of Precambrian sandstone and quartzite (Litherland and Power, 1989). The top of the mesetta is flat, with a gently rolling surface and at elevations ranging from 500–900 m a.s.l. (Da Silva Meneses and Bates, 2002). The substrate of the mesetta is rocky, and soils are thin and low in organic material (Litherland and Power, 1989). Continuity of the crystalline or sedimentary blocks of the mesetta is broken by an extensive network of peripheral or inter-mesetta depressions formed from a combination of erosion, dolerite dike intrusions and faulting on the mesetta (Litherland and Power, 1989; Da Silva Meneses and Bates, 2002). These depressions act as catchments for sediment and water, resulting in sediment accumulation, which supports more complex vegetation communities. High species diversity exhibited on the Huanchaca Mesetta, compared with other savanna regions of South America, is attributed to the long history of isolation of this edaphically-controlled table-mountain savanna (Mayle et al., 2007).

1.4 Vegetation

The cerrádo savanna on Huanchaca Mesetta is dominated by a continuous grass cover with sparsely scattered small trees and shrubs that grows on the thin, well-drained, nutrient-poor soils (Killeen, 1998b). Woody species include *Byrsonima coccolobifolia*, *Caryocar brasiliensis*, *Erythroxylum suberosum*, *Vochysia haenkeana*, and *Callisthene fasciculata*. Trees and shrubs include *Qualea multiflora*, *Emmotum nitens*, *Myrcia amazonica*, *Pouteria ramiflora*, *Diptychandra aurantiaca*, *Kielmeyera coriacea*, *Ouratea spectabilis*, and *Alibertia edulis*. Sub-shrubs include *Eugenia punctifolia*, *Senna velutina*, and herbaceous species include *Chamaecrista desvauxii*, and *Borreria* sp. Grass families include the Rapataceae (C₃) (*Cephalostemon microglochis*), Orchidaceae (*Cleistes paranaensis*) (CAM, C₃), Iridaceae (*Sisyrinchium* spp.) (C₄), Xyridaceae (*Xyris* spp.) (C₄), and Eriocalaceae (*Eriocalon* spp., *Paepalanthus* spp.,

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Syngonanthus spp.) (C₄) (Killeen, 1998b). In the inter-fluvial depressions organic rich soil is sufficiently deep to support humid evergreen forests islands which are typically dominated by mono-specific stands of *Mauritia flexuosa* (Mayle and Whitney, 2012; Da Silva Meneses and Bates, 2002). *Mauritia flexuosa* is a monocaulous, aborescent palm, averaging 20–30 m tall which is typically associated with a low, dense understory (da Silva and Bates, 2002; Furley and Ratter, 1988; Kahn, 1988;). *Mauritia flexuosa* is confined to lower elevations (< ca. 1000 m elevation) in warm/wet climates (Rull and Montoya, 2014). *Mauritia flexuosa* swamps favor inter-fluvial depressions that remain flooded during the dry season, when the surrounding terrains dry out (Huber, 1995a, b; Kahn and de Granville, 1992). The abundance of *M. flexuosa* in permanently flooded, poorly drained soils is the result of pneumatophores (aerial roots) which enable its growth in anaerobic conditions (Kahn, 1988; Rull and Montoya, 2014). Seasonal water deficits saturate the soil profile in the wet season and desiccate soil during the dry season resulting in a dominance of herbaceous versus woody plants surrounding the inter-fluvial depressions (Killeen, 1998b). The seasonal dryness leads to drought, plant water stress, and frequent fire activity resulting in the development of xeromorphic and sclerophyllous plant characteristics on the open mesetta (Killeen, 1998b). The spatial distribution of evergreen forest versus drought-tolerant savanna vegetation is additionally constrained by edaphic conditions limiting the expansion of forest vegetation because of the heavily weathered sandstone soils dominant outside the inter-fluvial depressions (Killeen and Schulenberg, 1998). Limited soil development precludes rainforest from developing on the large, rocky expanses of the mesetta (Killeen and Schulenberg, 1998). The essentially treeless campo cerrado that grows around Huanchaca Mesetta palm swamp is edaphically constrained and has likely grown on this mesetta for millions of years (Mayle and Whitney, 2012). Thus, the vegetation of the Huanchaca Mesetta is influenced by both climatic and non-climatic controls including seasonal hydrologic conditions, edaphic soil constraints and frequent fire activity (Killeen and Schulenberg, 1998).

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2 Materials and methods

2.1 Sediment core

A 5.48 m-long sediment core from Huanchaca Mesetta palm swamp was collected in 1995 using a Livingstone modified square-rod piston corer from the center of the swamp. The uppermost 15 cm, containing a dense root mat, was discarded because of the presence of fibrous roots and potential for sediment mixing. Huanchaca Mesetta sediment cores were transported to the Utah Museum of Natural History for analysis. They were photographed and described using a Munsell soil color chart. Visual descriptions, including sediment type, structure, texture, and organic content were undertaken to assist interpretation of the palaeo-environmental data.

2.2 Chronology

The chronological framework for Huanchaca Mesetta was based on eight AMS radiocarbon dates from non-calcareous bulk sediment and wood macrofossils analyzed at the University of Georgia Center for Applied Isotope Studies (Table 1). The uncalibrated radiometric ages are given in radiocarbon years before AD 1950 (years “before present”, yrBP). The errors are quoted at one SD and reflect both statistical and experimental errors. Radiocarbon ages were calibrated using CALIB 7.0 and the IntCal13 calibration dataset (McCormac et al., 2004). IntCal13 was selected in place of the SHcal13 calibration curve because of the latitudinal location (14° S) of Huanchaca Mesetta and the proximal hydrologic connection with the origin of the South American Monsoon in the Northern Hemisphere. The seasonal migration of the Intertropical Convergence Zone (ITCZ) is thought to introduce a Northern Hemisphere ¹⁴C signal to the low latitude Southern Hemisphere (McCormac et al., 2004). This study area is located in the low latitudes (14° S) and within the range of the ITCZ migration; thus, the IntCal13 calibration curve was selected for the radiocarbon calibrations. Following calibration, the mean age value of the largest probability at 2 sigma was used to create

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the smoothing spline age model using classical age-depth modeling, in the package CLAM (Blaauw, 2010) within the open-source statistical software R (Fig. 2).

2.3 Loss on ignition

The variability in the organic and carbonate content of sediments is used, in conjunction with magnetic susceptibility, to identify periods of variability in sediment composition and organic content throughout the Holocene. Organic and carbonate sediment composition was determined by loss on ignition, conducted at contiguous 1 cm increments throughout the cores. For each sample, 1 cm³ of sediment was dried in an oven at 100 °C for 24 h. The samples underwent a series of 2 h burns in a muffle furnace at 550 and 1000 °C to determine the relative percentage of the sample composed of organics and carbonates. Concentration was determined by weight following Dean (1974) standard methodology.

2.4 Magnetic susceptibility

Magnetic susceptibility (MS) was measured to identify mineralogical variation in the sediments (Nowaczyk, 2001). The MS of sediments is reflective of the relative concentration of ferromagnetic (high positive MS), paramagnetic (low positive MS), and diamagnetic (weak negative MS) minerals or materials. Typically, sediment derived from freshly eroded rock has a relatively high MS, whereas sediments that are dominated by organic debris, evaporites, or sediments that have undergone significant diagenetic alteration typically have a low or even negative MS (Reynolds, 2000). Shifts in the magnetic signature of the sediment can be diagnostic of a disturbance event (Gedye et al., 2000). Sediment cores were scanned horizontally, end to end through the ring sensor. MS was conducted at 1 cm intervals using a Barington ring sensor equipped with a 75 mm aperture.

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2.5 Charcoal

Sediment samples were analyzed for charcoal pieces greater than 125 μm using a modified macroscopic sieving method (Whitlock and Larsen, 2001) to reconstruct the history of local and extra-local fires. Charcoal was analyzed in contiguous 0.5 cm intervals for the entire length of the sediment core at 1 cc volume. Samples were treated with 5% potassium hydroxide in a hot water bath for 15 min. The residue was gently sieved through a 125 μm sieve. Macroscopic charcoal (particles > 125 μm in minimum diameter) was counted in a gridded petri dish at 40 \times on a dissecting microscope. Non-arboreal charcoal was characterized by two morphotypes: (1) cellular “graminoid” (thin rectangular pieces; one cell layer thick with pores and visible vessels and cell wall separations) and (2) fibrous (collections or bundles of this filamentous charcoal clumped together). Arboreal charcoal was characterized by three morphotypes: (1) dark (opaque, thick, solid, geometric in shape, some luster, and straight edges), (2) lattice (cross-hatched forming rectangular ladder-like structure with spaces between) and (3) branched (dendroidal, generally cylindrical with successively smaller jutting arms) (Jensen et al., 2007; Mueller et al., 2014; Tweiten et al., 2009). Charcoal pieces were grouped into non-arboreal and arboreal categories based on their morphology, which enabled the characterization of fuel sources in the charcoal record (Mueller et al., 2014). Charcoal counts were converted to charcoal concentration (number of charcoal particles cm^{-3}) and charcoal accumulation rates by dividing by the deposition time (yr cm^{-1}) using CHAR statistical software (Higuera et al., 2009). In CHAR, charcoal data was decomposed to identify distinct charcoal peaks based on a standard set of threshold criteria. Low frequency variation is considered background charcoal which reflect changes in the rate of total charcoal production, secondary charcoal transport and sediment mixing (Higuera et al., 2007). If the charcoal data exceed that background threshold, it is considered a peak and interpreted here as a fire episode. Background was calculated using a 700 year moving average.

and further developed in both North America (Fredlund and Tieszen, 1994) and the Neotropics (Bertoli de Pomar, 1971; Iriarte and Paz, 2009; Iriarte, 2003; Piperno and Pearsall, 1998; Piperno, 2006; Sendulsky and Labouriau, 1966; Söndahl and Labouriau, 1970; Teixeira da Silva and Labouriau, 1970; Zucol, 1999, 2000, 1996, 1998). The phytolith percentage diagrams were plotted using Tilia and Tilia Graphing software (Grimm, 1987).

3 Results

Four distinct vegetation types were identified based on constrained cluster analysis (CCA) of phytolith assemblages. These four zones are: the Late Glacial (14 500–11 500 cal yr BP), the Early Holocene (11 500–9000 cal yr BP), the Middle Holocene (8000–3500 cal yr BP), and the Late Holocene (3500 cal yr BP to present).

3.1 Zone 1: 14 500–11 500 cal yr BP Late glacial

The Late Glacial vegetation on Huanchaca Mesetta was dominated by arboreal taxa, grasses and Asteraceae (Opaque Perforated platelets) phytoliths (Fig. 3). The phytolith assemblage likely contains both in situ vegetation production and wind-blown vegetation from the surrounding rocky savanna. Both C_4 Panicoideae and C_3 Pooideae grass phytoliths were present during the Late Glacial. The presence of C_3 Pooideae grasses is interpreted as cooler Late Glacial conditions compared to present. The Late Glacial vegetation community at Huanchaca Mesetta lacks a modern analogue plant community in NKMNP. The presence of both of C_3 Pooideae and C_4 Panicoideae grasses suggest some degree of landscape heterogeneity. A consistent layer of very dark sandy silt dominated the lithology of Huanchaca Mesetta during the Late Glacial. The magnetic susceptibility and bulk density values were low and exhibit minimum variability compared to the rest of the record (Fig. 4). Coupled with LOI organic values below 10%, the sediment lithology was summarized as a low-energy depositional

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3.2 Zone 2: 11 500–9000 cal yr BP Early Holocene

Decreased C₄ Panicoideae grasses and consistent C₃ Pooideae grasses, aboreal, and Asteraceae (Opaque perforated platelets) phytoliths dominated the assemblage. The presence of C₃ grasses, and the absence of *M. flexuosa*, the dominant component of the modern basin vegetation, suggest temperatures cooler than present. The lithology, magnetic susceptibility, bulk density, and LOI values indicate minimal shift during the vegetation transition. Organic geochemistry reflected a change in organic matter source, with $\delta^{13}\text{C}$ values becoming more negative, indicating an increase in the contribution of C₃ vegetation ca. 11 000 cal yr BP. The $\delta^{13}\text{C}$ contribution of C₄ grasses decreased dramatically from 60 to 20 % during this period. These data correspond to a decrease in C₄ Panicoideae grass phytoliths and an increase in arboreal phytoliths. Low levels of terrestrial organic input into the system were indicated by low carbon concentrations and C:N values ranging between 25 and 30. N cycling changed during this zone, with $\delta^{15}\text{N}$ values exhibiting greater amplitude and higher frequency variability. The $\delta^{15}\text{N}$ values ranged between 4 and 8 ‰ indicating increased variability in denitrification rates associated with increasing wet (anaerobic) to dry (aerobic) conditions. The N concentrations were low, between 0.05 and 0.01 %, indicating minimal nitrogen availability in the system.

Charcoal accumulation at Huanchaca Mesetta increased ca. 11 200 cal yr BP coupled with an increase in the fire frequency to 5 episodes (periods of increased burning) per 1000 years. The peak magnitude values indicated two substantial fire episodes (periods of increased burning) ca. 10 200 and 9100 cal yr BP. The lack of significant change in the lithology suggests that taphonomic conditions were consistent during this interval. The increase in grass phytoliths during this period coupled with the increase in charcoal accumulation and fire episodes suggest that the Early Holocene vegetation community was becoming increasingly more fire dependent and vegetation was likely adapting to the increase in fire frequency associated with the period.

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6500 cal yr BP suggesting a change in the fire ecology on the mesetta. From 5000 to 3750 cal yr BP, grass charcoal increased. This is coincident with the establishment of *M. flexuosa* palm swamp and increased C₄ grasses in the surrounding watershed. After 3900 cal yr BP, charcoal accumulation and fire frequency increased. Significant increases in grass charcoal reflected a change in the fuel composition in the watershed. Phytolith, isotope and charcoal data suggest that after 3900 cal yr BP, the *M. flexuosa* within the basin became increasingly fire-sensitive and the occurrence of a fire within the palm stand would have had consequences for the vegetation not adapted to fire. The fire adapted C₄ grass dominated watershed continued to be fire-dependent.

3.4 Zone 4: 2800 cal yr BP–Present: Late Holocene

During the Late Holocene arboreal taxa was replaced by a pure stand of *M. flexuosa*. C₄ Panicoideae (warm, wet) grasses continued to dominate the surrounding watershed. *Mauritia flexuosa* values reached the highest levels during the Holocene from 2000–1200 cal yr BP. The hat shaped phytoliths indicate very low concentrations of other palms during this time. There was a gradual decrease in *M. flexuosa* towards present coupled with the highest levels of C₄ Panicoideae grasses ca. 200 cal yr BP and a decrease in C₄ Chloridoideae (warm, dry) grasses in the surrounding watershed. The lithology consisted of black detrital peat ca. 2450–2050 cal yr BP associated with high LOI (ca. 22) and magnetic susceptibility values (ca. 1000). After 2500 cal yr BP the %C, %N, and δ¹⁵N increased suggesting moist, anoxic conditions that enabled moderate denitrification from the swamp. These lithologic and isotopic data represented the establishment of modern palm swamp characterized by increased autochthonous organic accumulation. The δ¹³C values reached modern levels by 2800 cal yr BP although, values exhibit increased variability, fluctuating between –19 and –24‰ co-varying with the C₄ grass contribution between 10–20%. After ca. 800 cal yr BP δ¹³C values were ca. –18‰ and the % C₄ contribution was ca. 50%. These data corresponded to the highest levels of C₄ Panicoideae grass phytoliths in the record. The dark detrital peat lithology was interrupted by two coarse sand layers

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ca. 1550 cal yr BP and ca. 300–200 cal yr BP, followed by a shift back to black detrital peat ca. 200 cal yr BP to present. These sand layers were characterized by a decrease in LOI from ca. 22 to 2, C : N ratios from ca. 25 to 0, and $\delta^{15}\text{N}$ from ca. 5 to 0‰ coupled with increased magnetic susceptibility and bulk density values suggesting clastic flood events associated with sandy sediments low in organic material. From 300 cal yr BP %C values increased from ca. 1 % to > 20 % reached the highest values in the record. The %N values increased from ca. 1 to the peak Holocene values of 1.2 near at present. The dramatic increases in both %C and %N were likely the result of in situ carbon cycling and nitrogen fixation.

Charcoal accumulation at Huanchaca Mesetta remained low 2800 to 1800 cal yr BP with a FRI of 5 episodes (periods of increased burning) per 1000 years from 2800 to 1800 cal yr BP Grass charcoal reached the highest continuous levels ca. 2800 to 2000 corresponding to high levels of fire adapted C_4 grass phytoliths. Increased grass charcoal coupled with low peak magnitude values and high fire frequency indicated that the vegetation surrounding the palm swamp was fire dependent and fire adapted. However within the moist *M. flexuosa* palm stand, the vegetation remained fire sensitive. Charcoal accumulation increased ca. 1400 to 1200 cal yr BP and 700 cal yr BP, and reached peak Holocene values ca. 500–400 cal yr BP. Increased charcoal was coupled with the lowest FRI values in the record. Peak magnitude increased significantly around 1200 cal yr BP and the largest peak magnitude values ca. 200 cal yr BP. These charcoal values were cropped for plotting and visualization purposes. Raw counts exceed 1200 thus the values are also provided as log transformed (Fig. 7). Peak frequency increased after ca. 400 cal yr BP to ca. 4 fire episodes (periods of increased burning) per 1000 years towards present. There was a decrease in grass charcoal indicating increased woody biomass burned. The increased charcoal accumulation coupled with low FRI and more woody charcoal was interpreted as fire episodes that infrequently penetrated the fire sensitive palm stand and burned the *M. flexuosa* woody biomass. The charcoal, phytolith, and isotope data collectively suggest that the

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Middle Holocene dry period suggests a common mechanism affecting Amazonian moisture and precipitation. During this period, weakened SASM convective activity was attributed to an orbitally-driven minimum in January insolation at 15–10° S (Berger and Loutre, 1991). Reduced insolation would have restricted the southerly penetration of the ITCZ and deep cell convective activity over the Amazon Basin, thus decreasing the length of the summer rainy season resulting in longer, more severe dry seasons (Berger and Loutre, 1991).

The discrepancy in increased fire activity in the lowlands sites and decreased fire activity on the mesetta is attributed to fuel connectivity. In the lowland sites of Laguna Bella Vista, Laguna Chapin, and Laguna Orícore, dry forest–savanna vegetation provided sufficient fuel and increased fire activity during the Middle Holocene. At Huanchaca Mesetta decreased available moisture limited vegetation growth and fuel availability, particularly in the edaphically constrained rocky mesetta surrounding the basin. The lack of fine C₄ grass connective fuels resulted in decreased burning on the mesetta.

In the Late Holocene (3550 cal yr BP to present) the pollen assemblages of Laguna Bella Vista, Laguna Chaplin and Laguna Orícore, indicate an expansion of humid evergreen closed-canopy rainforest vegetation coupled with significant decreases in charcoal concentrations (Burbridge et al., 2004; Burn et al., 2010; Carson et al., 2014). Additionally, Lake Titicaca reach modern water levels during this time (Rowe et al., 2003) indicating wetter regional conditions with less severe dry seasons. The rainforest–savanna ecotone is currently at its most southerly extent over at least the last 50 000 years (Mayle et al., 2000; Mayle and Whitney, 2012; Burbridge et al., 2004). The progressive succession through the Holocene in the lowlands of NKMNP from savanna/semi-deciduous forest to semi-deciduous/evergreen forest to evergreen rainforest is part of a long-term uni-directional trend of climate-driven rainforest expansion associated with the regional increase in precipitation associated with a stronger South American Summer Monsoon (Mayle et al., 2004). The basin wide increase in mean annual precipitation and reduction in the length/severity of

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hypothesis suggests that the increase in *M. flexuosa* and charcoal accumulation is attributed to increased precipitation and wet season lightning fires driven by strengthened SASM activity (Kahn and de Castro, 1985; Kahn and de Granville, 1992; Kahn, 1987). The second hypothesis suggest that the simultaneous rise in *M. flexuosa* and charcoal was linked to intentional planting or semi-domestication of *M. flexuosa* for human use (Behling and Hooghiemstra, 1998, 1999; Montoya et al., 2011a; Rull and Montoya, 2014). Currently there is insufficient archaeological evidence from any of these savanna sites to support a robust anthropogenic signal (Rull and Montoya, 2014). Previous paleoecological studies in the lowlands demonstrate humans were the dominant driver of local-scale forest–savanna ecotonal change in those areas (e.g. Bolivian *Llanos de Moxos*) dominated by complex earth-moving pre-Columbian cultures (Carson et al., 2014; Whitney et al., 2014). These studies suggest that even in areas with extensive geometric earthworks, inhabitants likely exploited naturally open savanna landscapes that they maintained around their settlement, rather than practicing labor-intensive deforestation of dense rainforest (Carson et al., 2014). Evidence for human occupation of the lowlands has been found with ceramics from soil pits in an interfluve ca. 25 km NW of Laguna Chaplin and abundant ceramics and charcoal dating to ca. 470 cal yr BP recovered from anthosols (terra preta) throughout La Chonta ca. 150 km W of NKMNP (Burbridge et al., 2004). Implementing a new methodology to concentrate and isolate cultigen pollen (Whitney et al., 2012), the re-analysis of pollen data from Laguna Bella Vista and Laguna Chaplin revealed *Zea mays* pollen was present around 1700 to 940 cal yr BP, approximately 2000 years after the initial increase in *M. flexuosa* at these sites (B. Whitney, personal communication, 2014). Although humans were present in NKMNP, there is no evidence that they drove regionally significant ecotonal changes in forest–savanna boundaries. The patterns of forest–savanna shifts exhibited at these sites are consistent with climate forcing (Burbridge et al., 2004). The absence of archaeological data on Huanchaca Mesetta coupled with the inhospitable, nutrient poor, rocky soil, and limited access to the mesetta would have made human habitation unlikely. Although the *M. flexuosa* swamps

may have been used for hunting and gathering purposes, these data do not suggest humans were the driving mechanism behind the initial establishment or proliferation of *M. flexuosa* in the interfluvial depressions of the Mesetta.

The comparison of the Huanchaca Mesetta record to previous studies coupled with the absence of archaeological remains on the mesetta support the third hypothesis, that expansion of *M. flexuosa* at this site was largely controlled by natural drivers (edaphic, climate, lightning caused fires) opposed to anthropogenic drivers. In contrast to the conclusions from other studies, this record provides no evidence for an anthropogenically-driven fire regime, deforestation, soil erosion, or cultivation on the mesetta. These data suggest that natural drivers control the continued presence of savanna vegetation and fire activity on the Huanchaca Mesetta for the past 14 500 years.

5 Implications for savanna ecology and conservation

The presence of savanna vegetation for the past 14 500 years at Huanchaca Mesetta has significant implications for understanding modern savanna ecology and for the implementation of conservation strategies in the 21st century. Previous research on the evolution and development of savanna ecosystems has attributed much of the development of savannas to anthropogenic origins driven by the intentional use of fire (Arroyo-Kalin, 2012; Behling and Hooghiemstra, 1999; Behling, 2002; Berrío et al., 2002a; Hooghiemstra et al., 1998; Ramos-Neto and Pivello, 2000; Rull and Montoya, 2014). The results from this study demonstrate that the continued presence of the savanna ecosystem at Huanchaca Mesetta is attributable to edaphic and climatic controls. The presence of fire in this system for the past 14 500 years indicates that naturally occurring, lightning-caused fire is an integral part of the ecology of the savanna ecosystem. Despite changes in floristic composition and tree density within the drainage basin, the savanna ecosystem has been resilient to major climatic changes in both temperature and precipitation since the Late Glacial period. These

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data suggest that savanna ecosystems will continue to be resilient to future climate change associated with global warming. The long history of ecosystem stability in the face of dramatic climate variability attests to the fact that the Huanchaca Mesetta savanna is one of the most floristically diverse savannas anywhere in the Neotropics (Da Silva Meneses and Bates, 2002). The continued protection of the Huanchaca Mesetta savanna as a UNESCO world heritage site, coupled with the savannas natural resilience to climatic change exhibited over at least the past 14 500 years, indicates that despite significant global warming predicted for the 21st century (IPCC, 2014), the future is optimistic for the conservation and preservation of biological diversity in the Huanchaca Mesetta savanna ecosystem.

Acknowledgements. Funding to S. Y. Maezumi was provided by Global Change and Sustainability Center, the Graduate Research Fellowship, the Don Currey Graduate Research Fellowship, and the PAGES Graduate Research Fellowship. We thank Mary McIntyre and Daniel Harris for their help in sample preparation and analysis. Jennifer Watling and the Archaeobotany Lab at the University of Exeter assisted in phytolith training. Lee Grismer provided support for this research. The University of Leicester provided funding to F. E. Mayle. We thank Tim Killeen and the Museo de Historia Natural “Noel Kempff Mercado”, Santa Cruz, Bolivia for providing logistical support, and in particular Rene Guillen and local guides from the village of Florida (e.g. Juan Surubi) for assistance with coring the site.

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Table 1. AMS radiocarbon dates from Huanchaca Mesetta.

Lab number	Material	Depth (cm)	^{14}C age (yr BP)	$\delta^{13}\text{C}$ ratio	Intcal 13 BP 2 sigma
UGAMS 15158	Macrofossil	17	190 ± 20	-28.8	0–289
UGAMS 17252	Bulk Sediment	58	2310 ± 25	-18.8	2211–2356
UGAMS 15264	Bulk Sediment	118	1360 ± 20	-22.9	1272–1305
UGAMS 12023	Bulk Sediment	190	2480 ± 20	-22.62	2473–2715
UGAMS 17253	Bulk Sediment	225	3365 ± 25	-20.7	3561–3689
UGAMS 17254	Bulk Sediment	277	6545 ± 30	-22.6	7422–9622
UGAMS 15159	Bulk Sediment	320	8600 ± 30	-22.8	9524–9622
UGAMS 17255	Bulk Sediment	380	$11\,905 \pm 35$	-16.3	13\,577–13\,789

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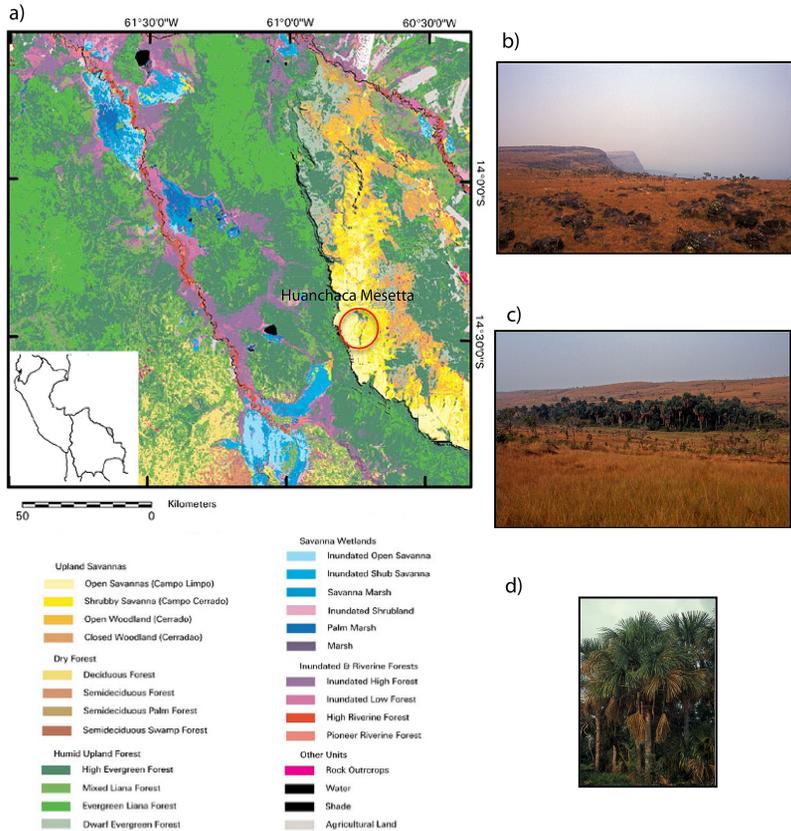


Figure 1. Huanchaca Mesetta study site, (a) vegetation map of Noel Kempff Mercado National Park modified from Killeen et al. (1998), (b) view from atop Huanchaca Mesetta, (c) Huanchaca Mesetta palm swamp, (d) mono specific stand of *Mauritia flexuosa*.

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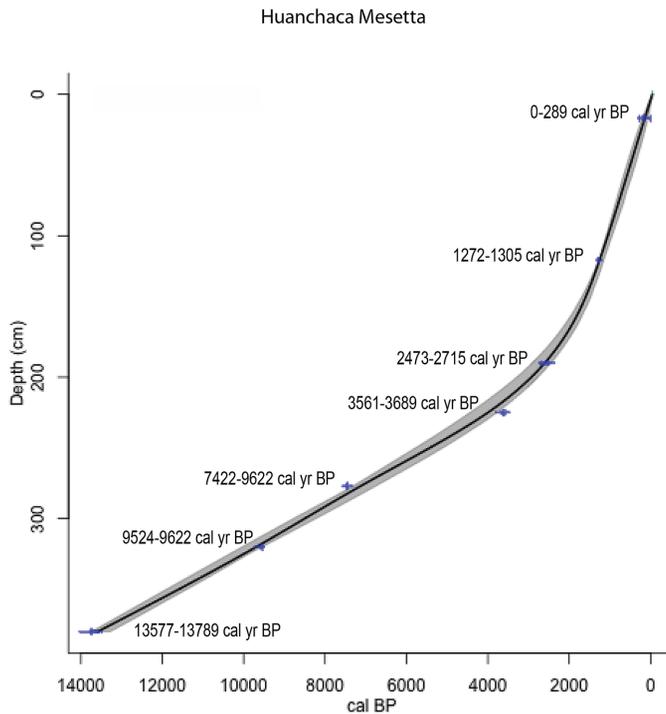


Figure 2. Clam age-depth model for Huanchaca Mesetta.

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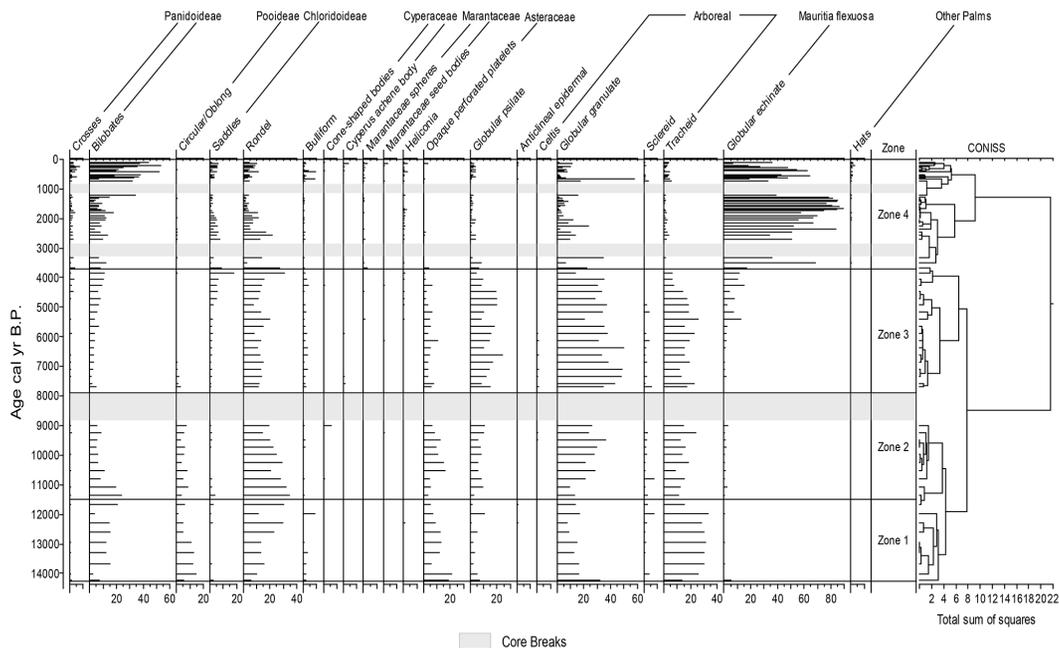


Figure 3. Huanchaca phytolith data separated by zones created by constrained cluster analysis (CONISS). Grey bars indicate core breaks.

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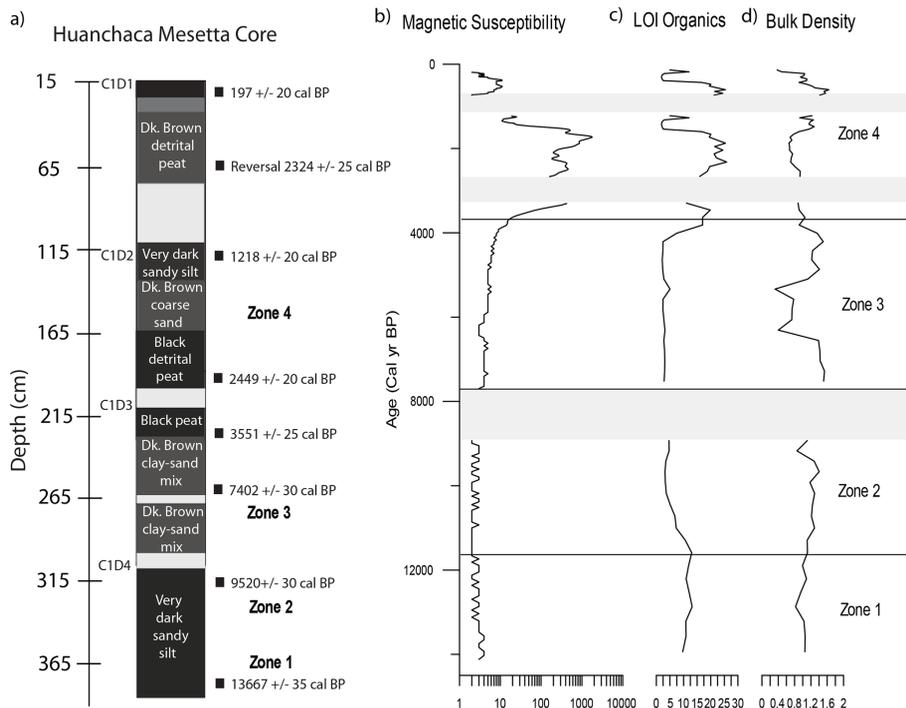


Figure 4. Huanchaca Mesetta lithology, **(a)** lithological description of the core profile, **(b)** magnetic susceptibility, **(c)** loss on ignition (LOI), **(d)** bulk density. Grey bars represent core breaks.

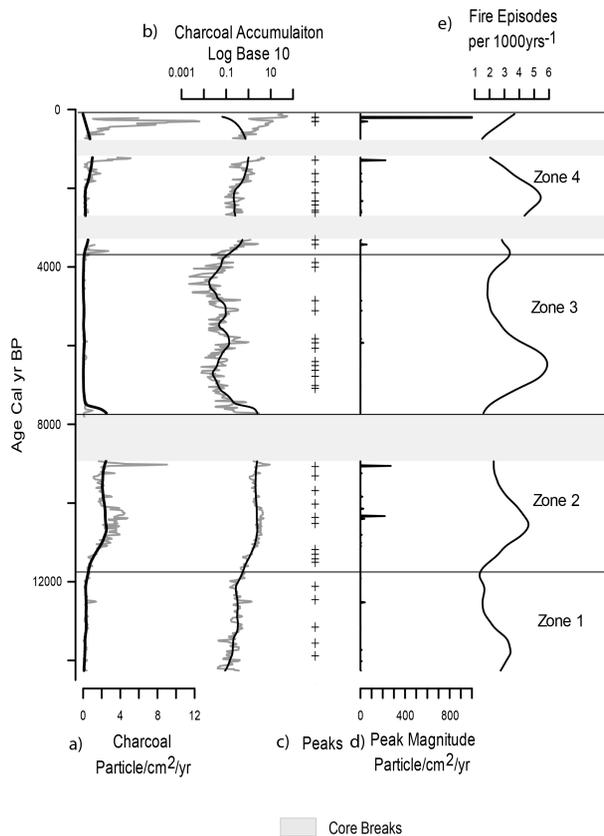


Figure 6. Huanchaca Mesetta charcoal data, **(a)** charcoal accumulation in grey, black background, **(b)** charcoal accumulation log base 10 in grey, black background, **(c)** peaks indicated by crosses, **(d)** peak magnitude, **(e)** fire episodes per 1000 years. Grey bars indicate core breaks.

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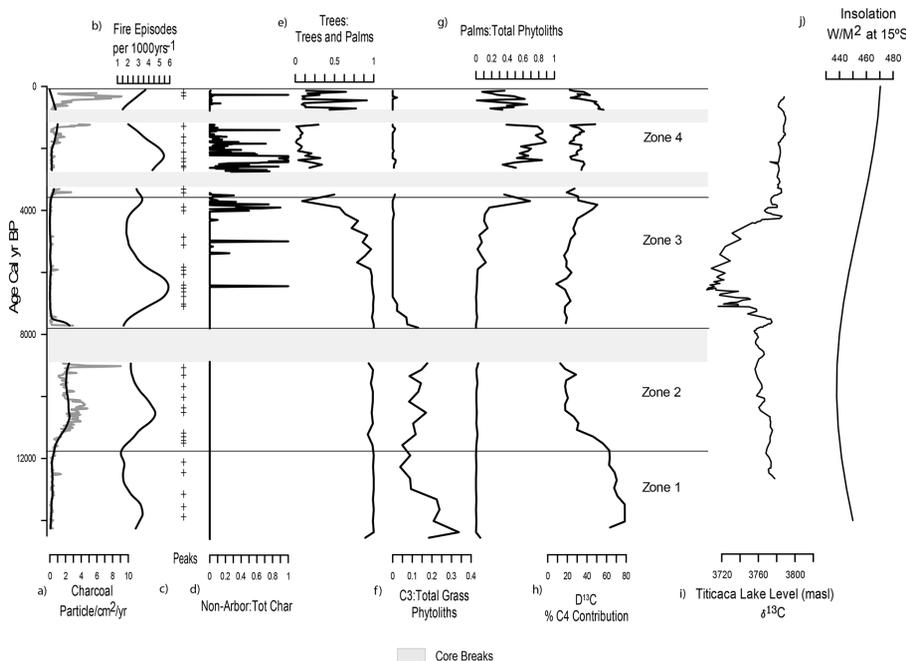


Figure 7. (a) Charcoal accumulation in grey with background in black, (b) fire frequency per 1000 years, (c) peaks indicated by plus signs, (d) ratio of non-arbooreal to total charcoal, (e) ratio of trees and palms, (f) ratio of C₃ to total grasses, (g) ratio of palms to total phytoliths, (h) percent C₄ contribution, (i) lake level of Titicaca in m.a.s.l., and (j) insolation at 15° S. Grey bars indicate core breaks.