GEOLOGY

No high Tibetan Plateau until the Neogene

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The Late Paleogene surface height and paleoenvironment for the core area of the Qinghai-Tibetan Plateau (QTP) remain critically unresolved. Here, we report the discovery of the youngest well-preserved fossil palm leaves from Tibet. They were recovered from the Late Paleogene (Chattian), ca. 25.5 ± 0.5 million years, paleolake sediments within the Lunpola Basin (32.033°N, 89.767°E), central QTP at a present elevation of 4655 m. The anatomy of palms renders them intrinsically susceptible to freezing, imposing upper bounds on their latitudinal and altitudinal distribution. Combined with model-determined paleoterrestrial lapse rates, this shows that a high plateau cannot have existed in the core of Tibet in the Paleogene. Instead, a deep paleovalley, whose floor was <2.3 km above mean sea level bounded by (>4 km) high mountain systems, formed a topographically highly varied landscape. This finding challenges prevailing views on tectonic processes, monsoon dynamics, and the evolution of Asian biodiversity.

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INTRODUCTION

The collision between the Indian and Eurasian plates changed the topography of southern Asia and led to the reshaping of regional and even global atmospheric circulation and climate (1, 2). However, the topographic evolution and paleoclimate history of the Qinghai-Tibetan Plateau (QTP) before and soon after the collision is very poorly constrained (3–7).

The Lunpola Basin is a large Cenozoic sedimentary basin on the QTP and is situated in the Bangong-Nujiang suture zone formed by the Mesozoic collision of the Lhasa and Qiangtang blocks (Fig. 1) (8, 9). The basin has been the focus of several paleoaltimetric studies (3, 5–7, 10, 11) because it offers considerable potential for understanding Tibetan landscape evolution and testing geodynamic models of the QTP formation. The Paleogene Lunpola sediment depocenter is located between the paleo-Gangdese (Eocene elevation, ~4.5 km) (12) and the paleo-Qiangtang (Tangulha) mountain ranges (Middle Oligocene elevation, >5.0 km) (13) to the south and north of the basin, respectively. Unexpectedly, Late Paleogene paleoelevation estimates for this core of Tibet diverge by as much as 4000 m, with some isotope studies suggesting a high (4 to 5 km) Lunpola Basin floor (3, 7), while paleontological data point to much lower elevations of between ~1 and ~3 km (4–6, 11).

Stable isotope quantitative paleoaltimetric proxies are widely used but rely heavily on assumptions regarding the composition of moisture sources, spatially and temporally varying isotopic lapse rates, evapotranspiration, and air parcel trajectories that can change seasonally (14–16). Isotope paleoaltimetry works best where prevailing winds rise normal to a single mountain ridge and where Rayleigh distillation occurs, whereas evaporation/precipitation recycling over plateaus or multiple mountain ranges generates large, poorly quantifiable, uncertainties (17).

Similarly, fossil-based paleoelevations based on organismal temperature constraints and thermal lapse rates can be unreliable because these lapse rates are highly variable in time and space (18), requiring prior knowledge of the topography under investigation. Moreover, all previous fossil-based surface height estimates for the core area of the QTP have used global (4) or regional (5) free-air thermal lapse rates instead of the more appropriate local paleoterrestrial thermal lapse rates (19), which incorporate the surface processes that strongly influence surface temperatures and land-atmosphere energy fluxes.

Plant megafossils are important indicators for paleoenvironmental reconstruction because their morphologies and spatial distributions are environmentally restricted and are seldom preserved far from their growth site (20). However, they are rarely recovered from the OTP because of logistical constraints. Here, we describe newly discovered palm fossil (Fig. 2 and fig. S1) leaves from the Lunpola Basin. The leaves are ~1 m in length, with leaf blades radiating from the distal ends of ~70-cm-long petioles. Because the unique anatomical and physiological features of palms render them inherently frost sensitive (21), we can constrain quantitatively, with confidence, the maximum possible elevation of the floor of the Lunpola Basin at the end of the Paleogene and before the rise of Himalaya (22). To do this, we determine paleotemperatures at sea level using coeval proxy data, and, combined with climate model-generated local terrestrial thermal paleo-lapse rates for a range of topographic scenarios, paleoelevations are estimated using the coldest month mean temperature (CMMT) and the following expression

$$Z = (T_{\rm msl} - T_{\rm h})/\gamma$$

where Z is the elevation, $T_{\rm msl}$ is the CMMT at mean sea level, $T_{\rm h}$ is the CMMT at elevation, and γ is the local terrestrial thermal (CMMT) lapse rate (19).

RESULTS

Fossil material and geological context

Palms (family Arecaceae) are diverse with ca. 2585 living species and ca. 188 extant genera. Today, they are all restricted to tropical and subtropical regions and are important environmental indicators because in natural populations, they are highly susceptible to freezing, particularly as seedlings (21). Moreover, the thermophilic requirements of palms have remained relatively stable throughout the Cenozoic (23, 24).

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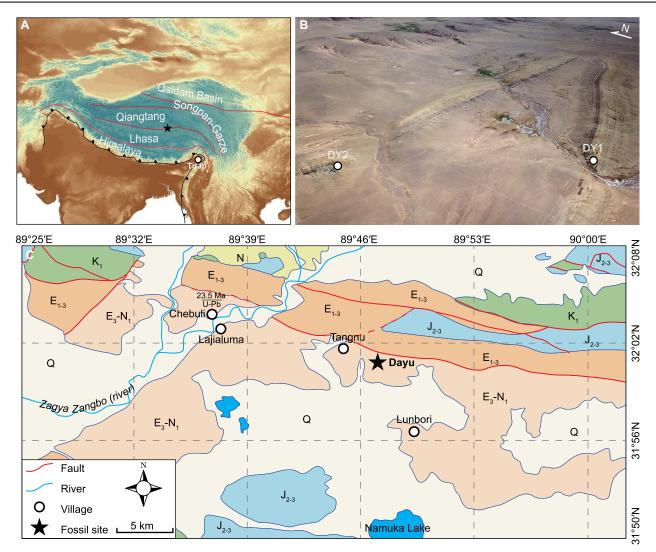


Fig. 1. The palm fossil locality at Dayu, Lunpola Basin, Shuanghu County, central Tibet, China. (A) Main terranes of the Tibetan Plateau and the Himalaya. The star marks the palm fossil location, while the circle marks the Tirap mine site. (B) Fossil sites in Lunpola Basin, both site DY1 and site DY2 bear palm fossils and belong to the same depositional horizon on the limbs of an anticline. (C) Geological map near the fossil locality in Lunpola Basin. J₂₋₃, Middle-Upper Jurassic Jienu Group; K₁, Lower Cretaceous volcanics; E₁₋₃, Paleocene-Oligocene Niubao Fm.; E₃-N₁, Oligocene-Miocene Dingqing Fm.; Q, Quaternary. Photo credit: T. Su, Xishuangbanna Tropical Botanical Garden (XTBG), Chinese Academy of Sciences (CAS).

Newly discovered well-preserved fossil palm leaves, here assigned to a new species *Sabalites tibetensis* T. Su et Z.K. Zhou sp. nov. (Fig. 2), occur in the Lunpola Basin (32.033°N, 89.767°E) within Late Paleogene lake sediments of the lower Dingqing Formation. Ground and drone surveys locate this horizon ~120 m below a U-Pb dated bentonite [23.6 \pm 0.2 million years (Ma)] (Fig. 1 and fig. S2) (10). Magnetostratigraphic constraints (5) point to the palm layer forming at ~25.5 \pm 0.5 Ma (Chattian). This finding, coupled with well-constrained paleoterrestrial lapse rates, allows us to quantify with a high degree of certainty the Late Oligocene elevation of the basin floor.

Numerous palm fossils have been reported worldwide (table S1), and fossils in this study can be assigned to the subfamily Coryphoideae, which, although it includes the most cold-tolerant tribe Trachycarpeae, also encompasses the tribe Sabaleae. Sabaleae contains genera to which *Sabalites*, in having costapalmate leaves (25), are most similar. Fossils from the Lunpola Basin are characterized by their prominent spine-like

structures at the bases of the leaf blades (Fig. 2 and table S2), which are different from any other previously reported palm fossils.

S. tibetensis is morphologically most similar to living palms that belong to the tribe Sabaleae, which can tolerate CMMTs no lower than 5.2°C (21), while our own survey of the morphologically closest extant relatives suggests that S. tibetensis may not have tolerated a CMMT lower than 7.7°C and a mean annual temperature (MAT) below 14.2°C (table S3). However, MAT is not a meaningful measure of cold tolerance because any given MAT can reflect a wide range of growth-limiting thermal extremes, depending on the annual thermal range. Most critical is the lowest temperature reached and the duration of freezing encapsulated as the CMMT and the frequency of freezing conditions. For a conservative approach that yields the highest possible paleoelevations of the Lunpola Basin floor, we use the lowest CMMT that can be tolerated by all living palms (5.2°C) (21), rather than that of the morphologically nearest living genera (7.7°C).

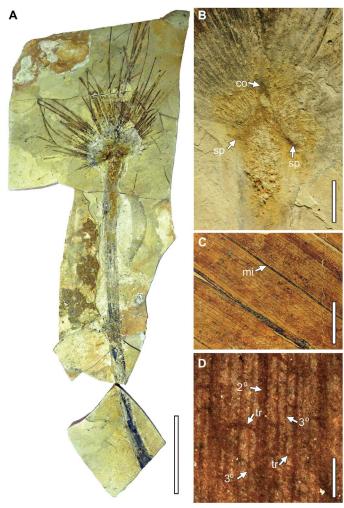


Fig. 2. S. tibetensis T. Su et Z.K. Zhou sp. nov. (A) Holotype, specimen no. XZDY2-0200. (B) Enlargement of the basal part of leaf blade. co, costa; sp, spine-like structures. (**C** and **D**) Specimen no. XZDY1-0071, venation pattern. mi, mid-vein; 2°, secondary vein; 3°, tertiary vein; tr, transverse vein. Scale bars, 10 cm (A), 3 cm (B), 1 cm (C), and 1 cm (D). Photo credit: T. Su, XTBG, CAS.

To obtain the maximum paleoelevation at which *S. tibetensis* could have grown requires knowledge of paleothermal lapse rates at the growth location and the temperature regime at sea level at that location. Past attempts to quantify paleosurface height using this approach have used global (4) or regional (5) free-air thermal lapse rates, but these are highly variable in time and space, particularly inland, over plateaus and in mountainous landscapes (18). Moreover, free-air thermal lapse rates are very different from those that plants experience within a few meters of the land surface (terrestrial thermal lapse rates) (18) and are therefore inappropriate for phytopaleoaltimetry (19). Because terrestrial lapse rates are highly dependent on local climate and, thus, topography, justifying any particular lapse rate for Tibetan paleolocations without integrating climate modeling is problematic and unconstrained.

Experimental design

To determine surface paleoelevation from our palm fossils, we modeled Late Paleogene local terrestrial thermal lapse rates (LTTLRs) at the Lunpola paleolocation (31.73°N, 83.11°E) using the HadCM3L (specifically HadCM3LB-M2.1) General Circulation Model, a version

of the U.K. Met Office HadCM3 v. 4.5 family (26) with a resolution of $3.75^{\circ} \times 2.5^{\circ}$ longitude × latitude for both atmosphere and ocean.

Thirteen topographic sensitivity studies (scenarios V1 to V8 and P0 to P4) for Tibet were performed (fig. S3). Scenarios V1, V3, V5, and V7 comprise a topographic valley system where a Gangdese mountain range is set to 4.5 km, the Qiangtang mountain range is set to 4 km, and a valley floor height of 2 km is set between them. Scenarios V2, V4, V6, and V8 comprise a 3.3-km valley floor between the Gangdese and Qiangtang mountain system, each set to 4 km. Because the paleorotated proxy site (31.73°N, 83.11°E) is located on the boundary between two model grid cells, each valley topographic configuration was run for two model latitudinal locations: 32.5°N (V1 to V4) and 30.0°N (V5 to V8). These valley systems were further modified by opening (V1, V2, V5, and V6) and closing the eastern end of the valley into Yunnan (V3, V4, V7, and V8) based on recent data showing an elevated eastern Tibet at 33 Ma (27). Scenarios P0 to P4 represent Tibet as a series of simple plateaus at elevations of 0 to 4 km, respectively, with no mountain ranges.

Because pCO_2 is uncertain for the Late Paleogene, with proxy estimates ranging from 280 parts per million (ppm) to as high as 1120 ppm (although likely not >840 ppm) (28), we also performed two CO_2 sensitivity studies to assess the influence of CO_2 on lapse rates. Our modeling automatically accounts for secular climate change. The Lunpola Late Paleogene LTTLRs vary, depending on topography ranging from 4.1° to 7.64°C/km (Table 1), but doubling CO_2 made a little difference (+0.1° \pm 0.159°C/km). These LTTLRs differed markedly from the Late Paleogene global average terrestrial lapse rates (4.61° to 4.73°C/km).

An essential component of the surface paleoelevation estimates using thermal lapse rates is knowledge of terrestrial temperatures at mean sea level ($T_{\rm msl}$) proximal to the location whose elevation is being sought. Model-derived $T_{\rm msl}$ values are topographically dependent (Table 1), so here, we use real-world proxy data to estimate conservative (warm) $T_{\rm msl}$ values for the Lunpola site. The closest coeval terrestrial proxy data is a Climate-Leaf Analysis Multivariate Program (CLAMP)–derived CMMT of 20.4° \pm 3.6°C from Tirap, Assam, Northeast India (27.289°N, 95.771°E) (29). Inconsistencies between proxy temperatures at Tirap (MAT and CMMT) and model predictions for that location highlight poorly performing model scenarios (Table 1).

The use of the lowest CMMT threshold for most extant palms (5.2°C) and the Late Paleogene local LTTLR at the palm locality shows in all but two scenarios (V5 and V7) that the fossil palm could not have lived higher than $\sim\!\!2.3$ km above sea level (Table 1). This elevation bound rises by only 400 m if we use the CMMT threshold (2.2°C) of the most cold-tolerant extant palms belonging to the Trachycarpeae, to which our fossil material shows only superficial resemblance and to which it is unlikely to be related.

Scenarios V1 to V4 all fail because the predicted CMMTs are too low at the prescribed land surface, as do scenarios V2, V4, V6, and V8, which also fail because the maximum survivable elevation (\sim 2.1 km) is below the model land surface (3.2 km). Model failure to reproduce the Tirap proxy conditions contributes to the failure of scenarios V2 and V6.

All plateau scenarios fail because either the CMMT is below the minimum 5.2°C for palm survivorship (P3 and P4), the predicted MAT at the Tirap site does not match the proxy data (P0 to P2), or the maximum elevation for palm survivorship is below the prescribed land surface (P3 and P4).

The only scenarios compatible with both the sea level proxy constraint and palm survivorship above model land surface elevation are

Table 1. Model-predicted climate and maximum survivable paleoelevations for an array of topographic scenarios (V1 to V8 and P0 to P4) for Tibet in the Late Paleogene. Thermal regimes regarded as fatal to palms (21) or failing to reproduce proxy values or occurring below prescribed land surfaces are shown in red. Blue shading indicates unconstrained (by proxy data) model-predicted CMMTs fatal to palms, and gray shading indicates model failure to reproduce proxy thermal conditions. Green shading indicates palm-survivable scenarios. The total number of frost days, number of 5-day cold events <0°C, and number of 5-day cold events <0°C

Topography scenario		MAT (°C)	CMMT (°C)	Total frost days (<0°C)	No. 5-day events <0°C	No. 5-day events <-5°C	Tirap MAT (°C) msl (proxy, 26.2° ± 2.3°C)	Tirap CMMT (°C) msl (proxy, 20.4° ± 3.6°C)	Lunpola CMMT lapse rate (γ) (°C/km)	Predicted highest survivable elevation (m)
V1	Valley (2 km), 4 km uplands, open	22.4	4.0	2504	161	68	28	19.3	5.3	2867
V2	Valley (3.2 km), 4 km uplands, open	15.9	0.4	3662	165	140	26.5	16.6	7.1	2132*
V3	Valley (2 km), 4 km uplands, closed	21.2	1.2	2854	160	91	27.8	19.1	4.1	3719
V4	Valley (3.2 km), 4 km uplands, closed	16.3	0.1	3695	185	146	27.8	19.1	6.8	2233*
V5	V1 latitude sensitivity	29.4	10.1	962	80	2	27.8	19.5	6.9	2194
V6		18.1	4.3	2590	161	81	26.4	17.0	7.2	2100*
V7	V3 latitude sensitivity	24.7	9.6	1034	81	6	27.8	19.1	6.7	2246
V8		17.8	3.5	2716	183	76	26.4	16.8	7.2	2094*
P0	0-km plateau	29.4	10.2	674	51	1	31.5	17.7	4.5	3380
P1	1-km plateau	26.1	8.9	1084	86	10	29.0	18.1	6.3	2396
P2	2-km plateau	23.2	8.0	1280	107	20	28.6	20.3	6.7	2260
P3	3-km plateau	18.6	4.9	2346	172	47	26.9	18.2	6.6	2304*
P4	4-km plateau	8.9	-1.7	4718	196	177	27.0	19.7	7.6	1982*

those with a valley floor at a height of 2 km bounded by mountains (V1, V3, V5, and V7), with either an open or closed eastern end. However, if the eastern end is closed (elevated) and the valley is deep (2-km valley floor) as in scenarios V3 and V7, this leads to lower lapse rates where palms can potentially survive to slightly higher elevations (52 m; Table 1), but eastern closure can make conditions in the valley highly susceptible to wind directions, resulting in slightly higher frequencies of potentially seedling-fatal frost days (Table 1).

Not all 2-km valley scenarios are plausible if model surface CMMT values are accepted. Model predictions independent of the Tirap proxy data (Table 1) show that all the more northerly valley scenarios (V1 to V4, valley located at 32.5°N) yield a CMMT at the valley floor surface below the 5.2°C palm-survivable threshold. The more southerly location of the Lunpola valley (30.0°N) with a 2-km valley floor and a closed eastern end (scenario V7) yields the most benign conditions in terms of potential survivability to well above the valley floor, and the second fewest frost days (Table 1), but even then, survivable elevations do

not exceed 2.3 km. Newly dated plant fossil evidence suggests just such a highland in eastern Tibet in the Paleogene (27).

DISCUSSION

Our finding that the Lunpola paleovalley floor was at an elevation of no more than 2.3 km at \sim 25 Ma supports other low elevations derived from paleontological data (4–6) that have largely been overlooked or dismissed possibly because the surface height estimates were based on organisms less well environmentally constrained than palms, the inappropriate use of free-air lapse rates, and the loosely constrained adjustments for secular climate change. These approaches have led to surface height estimates ranging from \sim 1 km (6) to \sim 3.5 km (5), of which the \sim 1-km elevation is the most compatible with the outcome of our work.

Our result contrasts with an earlier pollen-derived surface height estimate of 3090 to 3290 m (5), but this difference can be explained by (i) downslope transport of pollen from taxa living on the bounding

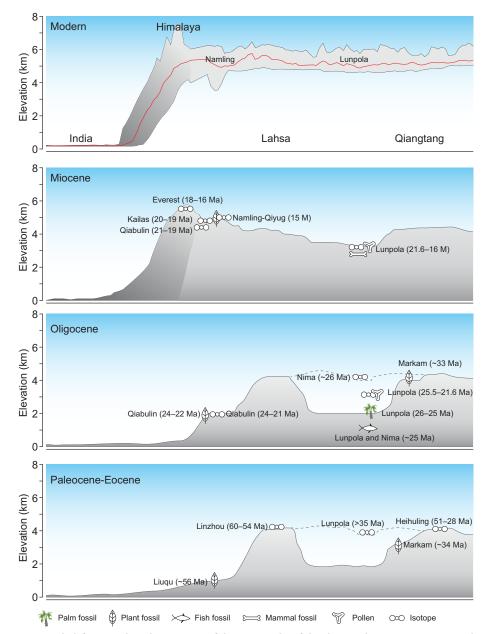


Fig. 3. Schematics representing south (left) to north (right) transects of the topography of the Lhasa and Qiangtang terranes, Tibet, and the Himalaya from the Paleocene to the present. Data sources: Paleocene—Liuqu (22), Linzhou (12), Lunpola (3, 13), Heihuling (13), and Markam (27); Oligocene—Qiabulin (22, 42), Nima (6, 43), Lunpola (6, 43), and Markam (27); Miocene—Qiabulin (22, 42), Kailas (33), Everest (44), Namling (45), Qiyug (46), and Lunpola (4, 5). Note that Markam is located in eastern Tibet.

mountain slopes and (ii) lapse rate values. First, pollen representing montane conifers (*Pinuspollenites*, *Piceapollenites*, and *Abiespollenites*) and temperate broad-leaved species (*Betulaepollenites*, *Quercoidites*, and *Ulmipollenites*) occur within the Dingqing sediments admixed with typically subtropical evergreen pollen types (*Castanopsispollenites* and *Meliaceoidites*) (5) that presumably represent the valley floor paleovegetation. This mixing of high and low elevation taxa would falsely raise the height estimate for the valley floor by an indeterminate amount, but it does point to a pronounced topographic relief surrounding the Dingqing lake catchment. Second, the pollen study (5) used a regional free-air thermal lapse rate of 5.79°C/km rather than the more

appropriate (18, 19) within-valley terrestrial thermal lapse rates of 6.74° to 6.9°C/km derived and applied here (Table 1). This difference in lapse rates alone raises the pollen-based valley floor elevation estimate by 308 to 367 m compared with ours. If only the subtropical components of the pollen flora were used, in combination with a more appropriate terrestrial lapse rate, the estimated valley floor elevation would be much lower than 3 km.

Free-air lapse rate and organismal tolerance issues apart, the estimate of ~3 km derived from rhinocerid remains (4) may not be entirely incompatible with our result in that the deposits yielding the rhinocerid are dated as 18 to 16 Ma and so are several million years

younger than the palm-bearing layer. There is some evidence to suggest that regional uplift and drying began at 23.7 Ma (11), although the rate and duration of this uplift require further investigation.

The valley system that we envisage is not the one produced by river incision, but a foreland basin-like depression resulting from the Mesozoic-Paleogene underthrusting of the Lhasa Terrane beneath the Qiangtang Terrane (30, 31) and, as such, must have extended along the length of the Bangong-Nujiang suture zone. Within this depression, a series of long-lasting interconnected deep and shallow lakes formed fed by inflowing streams and rivers (11, 30, 32), but with little evidence of significant fluvial outflows. In the Nima Basin, also within the Bangong-Nujiang suture zone, the Late Oligocene sedimentological evidence for gravity flows reaching the lake shoreline indicates that relief was relatively steep and that the source terrane was within a few kilometers (30). Moreover, the Lunpola lake system seems to have been saline and hyper stratified (32), which indicates strongly a closed lake system, even though it had inflows from the south and the nearby freshwater Baingoin Basin (32). The existence of a large accommodation space and lack of significant drainage from the valley system is evidenced by the accumulation of ~4000 m of Cenozoic sediments within several Bangong-Nujiang suture zone basins (11, 30, 33, 34). The documented lacustrine deposits of the Dingqing Formation alone are at least 1100 m thick based on borehole data (11).

The disparity between the isotope-derived paleoelevations and those based on fossils is likely due to downslope groundwater/stream transport of high-altitude meteoric isotope signatures. By contrast, leaves and animals have minimal capacity for downslope transport unscathed, and their environmental adaptations and tolerances reflect conditions close to their depositional location, i.e., surrounding the valley bottom lake margins. Together, the isotopic and biotic paleoaltimeters provide insights into the landscape diversity that are missed using only one form of elevation proxy.

Our discovery of *S. tibetensis* challenges the concept of a "proto-Tibetan Plateau," a term that suggests that the central part of Tibet was flat, if not as high as today (3, 7), or with moderate topographic lows of just 1 to 1.5 km below the adjacent Gangdese and Qiangtang highlands (5). Instead, we infer that a large valley system with a floor lower than 2.3 km AMSL existed in central Tibet as late as 25 Ma and that the topography of Tibet during the Late Paleogene was far more complex than often envisaged (Fig. 3), a complexity that may be fundamental to the evolution of modern Asian biodiversity "hot spots" (35). The deep paleovalley system between the Gangdese and Qiangtang highlands now accommodates ~4000 m of Cenozoic sediments (5) and suggests that a combination of valley fill and predominantly compressional tectonics raised the surface elevation after ~25 Ma to form the present high plateau.

MATERIALS AND METHODS

Morphological description

Family: Arecaceae Schultz Sch. Subfamily: Coryphoideae Burnett

Genus: Sabalites G. Saporta emended Read and Hickey **Species:** S. tibetensis T. Su et Z.K. Zhou sp. nov. **Holotype:** XZDY2-0200 (designated here) (Fig. 2A)

Paratypes: XZDY1-0071, XZDY2-0201, XZDY2-0202, and

XZDY3-0001 (designated here) (fig. S1)

Repository: Palaeoecology Collections, XTBG, CAS

Type locality: Dayu Village, Lunpola Basin, central Tibetan Plateau

Age: Late Oligocene (Chattian)

Etymology: The specific epithet "*tibetensis*" refers to the fossils being found in central Tibet.

Diagnosis: Leaves costapalmate. Costa/petiole thick at the base and gradually tapers toward apex, petiole robust, unarmed; longitudinal striations present on the petiole/costa. One pair of spine-like structures at the base of leaf blade. Leaf segment plicate, emerging from the costa an acute angle; fused at emerging point. Each leaf segment bears midveins; two order of veins present on either side of the mid-vein, transverse vein present either perpendicularly or oblique.

Description: The description is based on two specimens, which are part (Fig. 2A) and counterpart. Leaves are costapalmate (Fig. 2A and fig. S1); maximum preserved length of the leaf blade is ~310 mm, and maximum preserved width of the leaf blade is ~325 mm; leaf blade slightly asymmetrical at base; petiole present, unarmed, maximum preserved length is ~703 mm (including costa), maximum preserved width is ~30 mm, flattened, nearly same width from proximal to distal end; costa maximum length is 53 mm, wide at base (~34 mm) and gradually tapers distally; distinct spine-like structure present on the base of the costa, which is ~12 mm in length having a broad base and tapers gradually at the distal end; leaf segment plicate, fused only on the lower portion of the leaf blade; each leaf segment bears a mid-vein in the middle, numerous 2° veins run parallel on either side of the mid-vein, 3° veins present, running parallel in between 2° veins, three to five in number; transverse veins present either perpendicularly or oblique to 3° veins (Fig. 2, C and D).

Remarks: The characteristic features of the fossil leaf include palmate, plicate leaves with costa (costapalmate), and an unarmed long petiole. These characters suggest that the fossil leaves have affinities with the subfamily Coryphoideae in the Arecaceae. A large number of modern palm leaves were observed using herbaria at XTBG and the Kunming Institute of Botany, Kunming, the website of the Royal Botanic Garden, Kew, and Palmweb (www.palmweb.org).

As Read and Hickey (25) stated that "Since it is very difficult to identify specimens of modern palms accurately from their leaves alone, no attempt should be made to place fossil palm fragments in genera of modern palms unless unquestionably identifiable with them." Under these circumstances, the fossils were placed in the organ genus *Sabalites* G. Saporta proposed for costapalmate fossil leaves (36).

Numerous palm leaf fossils have been reported from various parts of the world, and it is not possible to compare all of them with the present fossil; however, we compared those that are most similar. In China, several costapalmate leaves have been described from Eocene sediments, such as Sabalites asymmetricus, Sabalites robustus, Sabalites tenuifolius, Sabalites szei, and Sabalites changchangensis (37), and Oligocene sediments, namely, Chuniophoenix slenderifolia, Livistona rotundifolia, and Trachycarpus formosa (38), based on some morphological features and preserved cuticular structures. In all the aforesaid fossils, the leaf blades are incomplete, the leaf segments are fused, and the preserved petioles are very short. These features are significantly different from those of the Tibetan fossils.

Similarly, a coryphoid palm leaf named *Sabalites longirachis* has been described from the early Campanian of Austria and Maastrichtian of the Pyrénées (23), and although it looks similar to our fossil, it has fused leaf segments, which is a major difference. Srivastava *et al.* (39) reported a similar type of coryphoid palm from the late Maastrichtian-Danian sediments of Deccan Intertrappean beds of central India, but their fossil bears a long costa and fused leaf segments as distinct from our Tibetan fossil leaf. Therefore, in the absence of any similar leaves, we proposed the new species *S. tibetensis* T. Su et Z.K. Zhou sp. nov.

Climate model

HadCM3L is a fully coupled atmosphere-ocean model consisting of 19 vertical levels in the atmosphere and 20 ocean depth levels and has an integrative vegetation model. We ran HadCM3L until ocean circulation equilibrated fully with no trend in the integrated ocean temperature. HadAM3, the atmospheric model of HadCM3L using the equilibrated sea surface temperatures (SSTs) from HadCM3L simulation, was used to perform the topography sensitivity studies. The HadCM3 family has seen extensive use as a principal model of the Inter-governmental Panel on Climate Change (IPCC) AR3-5 experiments. Crucially, the HadCM3 family shows skill in representing not only the mean climate state compared with observations globally (26) but also the monsoon climate of East Asia (40), where it outperforms most other models.

Simulations of the Chattian paleogeography use the Getech Plc. palaeogeographic reconstructions (0.5° \times 0.5° longitude \times latitude) (40) upscaled to the HadCM3L resolution (41). An ice sheet over Antarctica is also prescribed in these sets of simulations. Each simulation was run >6000 model years to achieve a steady state in the deep ocean as well as a radiative balance in the atmosphere.

SUPPLEMENTARY MATERIALS

Supplementary material for this article is available at http://advances.sciencemag.org/cgi/content/full/5/3/eaav2189/DC1

Fig. S1. S. tibetensis T. Su et Z.K. Zhou sp. nov.

Fig. S2. Stratigraphy of the Dingqing Formation at Dayu, Lunpola Basin.

Fig. S3. Late Paleogene model elevations (meters) showing the different topographic scenarios for the QTP region.

Table S1. Fossil records of palms around the world.

Table S2. Morphological comparisons between fossils from Lunpola Basin and modern palm genera. Table S3. Climate ranges of 12 living genera that show the closest morphological similarity to *S. tibetensis* T. Su et Z.K. Zhou sp. nov.

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