

1 **Palynological evidence for Late Miocene stepwise**
2 **aridification on the Northeastern Tibetan Plateau**

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4 **J. Liu¹, J. J. Li¹, C. H. Song², H. Yu¹, T. J. Peng¹, Z. C. Hui¹, and X. Y. Ye¹**

5 ¹MOE Key Laboratory of Western China's Environmental Systems & College of
6 Earth and Environmental Sciences, Lanzhou University, Lanzhou 730000, China

7 ²Key Laboratory of Western China's Mineral Resources of Gansu Province & School
8 of Earth Sciences, Lanzhou University, Lanzhou 730000, China

9 Correspondence to: J. J. Li (lijj@lzu.edu.cn)

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1 **Abstract**

2 Holding a climatically and geologically key position both regionally and globally, the
3 northeastern Tibetan Plateau provides a natural laboratory for illustrating the
4 interactions between tectonic activity and the evolution of the Asian interior
5 aridification. Determining when and how the Late Miocene climate evolved on the
6 northeastern Tibetan Plateau may help us better understand the relationships among
7 tectonic uplift, global cooling and ecosystem evolution. Previous paleoenvironmental
8 research has focused on the western Longzhong Basin. Late Miocene aridification
9 data derived from pollen now requires corroborative evidence from the eastern
10 Longzhong Basin. Here, we present a Late Miocene pollen record from the Tianshui
11 Basin in the eastern Longzhong Basin. Our results show that a general trend toward
12 dry climate was superimposed by stepwise aridification: a temperate forest with a
13 rather humid climate existed in the basin between 11.4 and 10.1Ma, followed by a
14 temperate open forest environment with a less humid climate between 10.1 and 7.4Ma,
15 then gave way to an open temperate forest-steppe environment with a relatively arid
16 climate between 7.4 to 6.4Ma. The vegetation succession demonstrates that the
17 aridification of the Asian interior occurred after ~7–8Ma, which is confirmed by other
18 evidence from Asia. Furthermore, the aridification trend on the northeastern Tibetan
19 Plateau parallels the global cooling of the Late Miocene; the stepwise vegetation
20 succession is consistent with the major uplift of the northeastern Tibetan Plateau
21 during this time. These integrated environmental proxies indicate that the long-term
22 global cooling and the Tibetan Plateau uplift caused the Late Miocene aridification of
23 the Asian interior.

24 **1 Introduction**

25 As the latter stage of the global Cenozoic cooling, the Neogene was a critical period
26 for northern hemispheric aridification, especially for the marked aridification of the
27 Asian interior. Establishing when, and how, this process of aridification began and
28 evolved is therefore vital for elucidating the interactions among tectonic uplift, global
29 cooling and ecosystem evolution. Although there is compelling evidence for the

1 aridification of the Asian interior, there is no consensus concerning its evolution and
2 driving mechanisms. For instance, previous researchers have suggested that the
3 aridification of the Asian interior began in the Late Miocene, based particularly on
4 biological and isotopic evidence (Andersson and Werdelin, 2005; Cerling et al., 1997;
5 Dettman et al., 2001; Eronen et al., 2012; Quade et al., 1989; Wang and Deng, 2005;
6 Zhang et al., 2012). However, others have argued that the process of Asian interior
7 aridification may have begun in the Early Miocene (22Ma) or even earlier (in the Late
8 Oligocene), as inferred from Miocene or Oligocene eolian deposition (Guo et al.,
9 2002, 2008; Qiang et al., 2011; Sun et al., 2010). The particular driving mechanisms
10 of such aridification also remain enigmatic. Up until now, the tectonic uplift of the
11 Tibetan Plateau (TP), global cooling and land–sea distributions have been suggested
12 as the major drivers (An et al., 2001; Gupta et al., 2004; Kutzbach et al., 1993; Liu
13 and Yin, 2002; Miao et al., 2012; Molnar et al., 2010). However, there is little
14 consensus about which one is the most important driver. We focused on the region of
15 the northeastern TP to explore the nature of the interactions between tectonics and
16 climate.

17 The geographically-extensive Longzhong Basin, consisting of a series of sub-basins,
18 is located in the northeastern TP. These sub-basins present a continuous record of
19 mammalian fossil-rich Cenozoic sediments, recording the effect of TP uplift on
20 regional climates (Fang et al., 2003, 2005; GRGST, 1984; Li et al., 2006, 2014), as
21 well as the effect of the global cooling. On the other hand, it lies in the so-called
22 monsoonal triangle, a transition zone from a warm-humid Asian monsoonal climate to
23 a dry-cold inland climate and to the alpine climate of the TP (Li et al., 1988, 2014)
24 (Fig. 1a). Its particular geological and geographical characteristics make it sensitive to
25 document the aridification history of northern China. As a field laboratory for
26 studying tectonic-climate interactions (Molnar et al., 2010; Tapponnier et al., 2001),
27 the Longzhong Basin might be the most promising for distinguishing TP uplift and
28 associated environmental change.

29 As a reliable paleoenvironmental proxy, pollen has been used to reconstruct past

1 climates because of its abundance and excellent preservation within sediments.
2 Previous research has demonstrated that the Tianshui Basin, as a sub-basin of the
3 Longzhong Basin, exhibits a typical Late Miocene lacustrine-fluvial sedimentary
4 succession containing abundant pollen (Li et al., 2006). Here we reconstruct a
5 high-resolution palynological record from the well-dated Yaodian Section, located in
6 the southern part of the Tianshui Basin. Our results not only provide new evidence for
7 the evolution of vegetation in the Late Miocene and climate change on northeastern
8 margin of the TP, but also shed new light on the aridification of the Asian interior.

9 **2 Geological and geographical settings**

10 The rhomboid-shaped Longzhong Basin, which is one of the largest intermountain
11 and fault-controlled sedimentary basins on the northeastern TP, is geographically
12 delineated by the left-lateral strike-slip Haiyuan Fault to the north, the Liupan Shan
13 Fault to the east and northeast, the Laji Shan Fault to the southwest, and the Western
14 Qinling Fault to the south (Fig. 1b). The Tianshui Basin, one of its sub-basins, is
15 located in the southeastern part of the Longzhong Basin (Fig. 1b). It has witnessed the
16 continuous deposition of mammalian fossil-rich Cenozoic sediments from the
17 surrounding mountains; these sediments record the interactions between mountain
18 uplift, erosion and climate change (Alonso-Zarza et al., 2009; Li et al., 2006; Liu et al.,
19 2015; Peng et al., 2012, 2015). At present, the East Asian Monsoon influences this
20 region, engendering a semi-humid, warm temperate, continental monsoon climate,
21 characterized by relatively hot, humid summers and cold, dry winters. The mean
22 annual temperature and mean annual precipitation of this area are ~11 °C and 492mm,
23 respectively, with rainfall concentrated mainly in summer and autumn (Fig. 1c). The
24 modern natural vegetation in this region is warm-temperature forest-grassland. Warm
25 grasslands are distributed in the valleys, and consist mainly of *Arundinella hirta*,
26 *Spodiopogon sibiricus* and *Themeda triandran*. Shrubs such as *Zizyphus jujube*,
27 *Sophora viciifolia* and *Ostryopsis davidiana* are found on the hillsides. Trees,
28 including *Quercus liaotungensis*, *Pinus tabulaeformis*, *P. armandi* and *Platycladus*
29 *orientalis*, grow in the mountains (Huang, 1997).

1 The selected Yaodian Section (105°55' E, 34°38' N) is located in the southern part of
2 the Tianshui Basin (Fig. 1d). The Neogene sequence in section is capped by loess and
3 lies unconformably on top of the Paleogene Guyuan Group. It has been divided into
4 the Ganquan Formation (Fm), the Yaodian Fm and the Yangjizhai Fm, in sequence
5 upwards (Li et al., 2006). In this study, our research mainly focuses on the Late
6 Miocene Yaodian Fm and Yangjizhai Fm. Based on a determination of lithology and
7 sedimentology, the Yaodian Fm can be divided into three principal strata. The lower
8 stratum consists of massive fine gravel sandstone, sandstone and brown silty
9 mudstone, occasionally with thin brown mudstone or interbedded paleosols, which
10 can be considered fluvial channel deposits (Fig. 2e). Abundant teeth of *Hipparion*
11 *weihoense*, *Cervavitus novorossiae*, *Ictitherium* sp. and their bone fragments were
12 excavated from this stratum. The middle stratum of the Yaodian Fm consists of the
13 interbedding of siltstone or fine sandstone with mudstone intercalated with paleosols,
14 overlying the fluvial channel deposits. The assemblage's characteristics are typical of
15 floodplain deposition (Fig. 2d). The upper stratum of the Yaodian Fm is characterized
16 by rhythmic cycles composed of grey or brown mudstone or sandy marlite and
17 intraclastic marl intercalated with brown siltstone and mudstone, and contains fossil
18 algae and gastropods; this section is representative of shallow lake deposition (Fig. 2a
19 and c). The upper stratum is common throughout the basin, and is analogous to the
20 "Zebra Bed" stratum found in the Linxia Basin in the western Longzhong Basin (Li et
21 al., 1995). The Yangjizhai Fm is principally composed of reddish brown mudstone or
22 silty mudstone and yellowish brown calcrete or calcareous mudstone, with scattered
23 sandstone or grey mudstone and marlite. These sediments were deposited under
24 strong evaporative conditions in distal floodplain to palustrine environments (Fig. 2b).
25 Previous paleomagnetic investigations have indicated that the Yaodian Fm ranges
26 from 11.67 to 7.43Ma in age, and that the Yangjizhai Fm dates from 7.43 to 6.40Ma,
27 both these ranges being consistent with the formations' biostratigraphic ages (Li et al.,
28 2006).

29 **3 Materials and methods**

1 Most of the samples came from lacustrine mud deposits and fine grain size
2 intercalations found in floodplain and fluvial channel deposits. Because the lower
3 10m of the Yaodian Fm consists of coarse gravel sandstone, and it was difficult to
4 find fine-grained sediments therein, this part of the formation was not sampled. A
5 total of 200 samples were processed for palynological analysis. For each
6 sample, >100g of sediment was washed in 20% HCl, soaked in 39% HF and then
7 treated with 10% HCl solution to enable fluoride dissolution. We then concentrated
8 pollen by physical enrichment procedures, using ZnCl₂ separation and ultrasound
9 sieving over a 10µm filter. Samples were stored in glycerin. Identifications were
10 based on atlas of pollen and spores (Wang, 1995; Song, 1999), as well as modern
11 reference slides from the collection of the Laboratory of Sporopollen Analysis of the
12 Geography Department of Lanzhou University. Palynological diagram was plotted
13 using Tilia v2.0.b.4 (Grimm, 1993) and pollen-assemblage zones were constructed
14 using Stratigraphically-constrained cluster analysis (CONISS) (Grimm, 1987).

15 **4 Results**

16 Only 126 of the 200 samples contained enough palynomorphs to provide reliable data;
17 the remaining 74 possessed fewer than 300 identifiable grains and have not been
18 included in the analysis. Most of the latter samples had been preserved under
19 oxidizing conditions, or had high carbonate content. Approximately 80 different
20 palynomorphs were identified at family or genus level. Percentages were expressed on
21 the total number of recognized taxa. Tree pollen consists mainly of *Pinus*,
22 Cupressaceae and *Ulmus*, along with *Quercus* and *Betula*. Additionally, a number of
23 subtropical plants pollen, such as *Liquidambar*, *Pterocarya* and *Carya* (which are no
24 longer found in this area today), appear often in low abundance. Herbaceous pollen is
25 mainly from *Artemisia*, Chenopodioideae, Poaceae and Asteraceae. Pollen from
26 extremely drought-tolerant plants, such as *Ephedra* and *Nitraria*, only appear
27 sporadically in single samples. In addition, the section also contains fern spores and
28 *Pediastrum* colonies. A selection of the more important taxa is given in Fig. 3.
29 CONISS (Grimm, 1987) yields three distinct zones, described from the bottom up as

1 follows:

2 **4.1 Zone 1 (195.5–158.5m, 11.4–10.1Ma)**

3 Samples from this zone exhibit high percentages of tree pollen, averaging 75%.
4 Coniferous taxa are mainly *Pinus* (19%) and Cupressaceae (18%), with smaller
5 amounts of *Picea* and *Cedrus*. *Ulmus* (20%) is the most common broadleaf tree pollen,
6 accompanied by pollen of *Betula* (3%), *Quercus* (2%) and *Salix* (2%). Other arboreal
7 taxa are *Juglans* and *Castanea*, with <2% respectively. Herbaceous taxa mainly
8 include *Artemisia* (7%), Chenopodioideae (6%) and Poaceae (2%), along with small
9 amounts of Asteraceae, Ranunculaceae and Rosaceae, with amounts <2%
10 respectively. Aquatic plants, algae and some subtropical taxa are also represented in
11 this zone with low abundance.

12 **4.2 Zone 2 (158.5–63.5m, 10.1–7.4Ma)**

13 In this zone, total tree pollen percentage decreases, averaging 54%. Coniferous taxa
14 are principally represented by *Pinus* (14%), Cupressaceae (7%), *Picea* (2%) and
15 *Cedrus* (1%). Among broadleaf trees, the dominant taxa are *Ulmus* (8%), *Quercus*
16 (2%), *Betula* (2%), *Salix* (2%) and *Juglans* (1%). Herbaceous taxa are dominated by
17 *Artemisia* (14%) and Chenopodioideae (9%), along with Poaceae (5%), Asteraceae
18 (3%) and Ranunculaceae (3%). Aquatic vegetation reaches the highest value found in
19 the entire profile. Subtropical taxa, such as *Liquidambar*, *Pterocarya*, *Carya* and
20 Rutaceae, are represented with low abundance. The zone is divided into two subzones,
21 Zone 2-1 (158.5–106.5m, 10.1–8.6Ma) and Zone 2-2 (106.5–63.5m, 8.6–7.4Ma).
22 Herbaceous pollen percentages are slightly higher in Zone 2-2 than in Zone 2-1.

23 **4.3 Zone 3 (63.5–30m, 7.4–6.4 Ma)**

24 The samples from this zone record a further decrease in tree pollen to an average
25 value of 39%. Coniferous taxa are characterized by *Pinus* (7%) and Cupressaceae
26 (5%). *Ulmus* (5%) dominates the broadleaf tree pollen, with *Quercus* and *Betula*
27 accounting for 2%, respectively. Herbaceous taxa are composed of *Artemisia* (19%),
28 Chenopodioideae (11%) and Poaceae (9%), together with Asteraceae (5%),

1 Ranunculaceae (3%), Brassicaceae (3%) and Polygonaceae (2%). Aquatic plants and
2 thermophilic species almost disappear.

3 **5 Discussion**

4 **5.1 Vegetation and climate reconstruction**

5 The sedimentary facies of the Yaodian Section indicate four successive depositional
6 stages: fluvial channel; floodplain; shallow lake; and distal floodplain to palustrine.
7 Transitionals can be dated to 10.4, 9.23 and 7.43Ma, respectively (Li et al., 2006) (Fig.
8 2). Our palynological record shows stepwise changes at 10.1 and 7.4Ma, lagging
9 slightly behind those evinced by the sedimentary facies. Another distinctive feature of
10 the palynological record is that the green lacustrine deposits of fine grain size exhibit
11 dense palynomorph concentrations, with higher tree pollen percentages. In contrast,
12 the reddish floodplain deposits with coarse grain sizes possess sparse palynomorph
13 concentrations, with higher herbaceous pollen percentages (Fig. 3). However, in the
14 same pollen zones, we find that the palynomorph concentration clearly changes
15 between different sedimentary facies, but that percentage fluctuations are minor.
16 Between different pollen zones, the palynomorph percentages change strongly within
17 the same sedimentary facies. We can therefore conclude that the changes in the
18 palynological record are caused by changes in regional vegetation, rather than
19 different preservation conditions. The paleoecological information inferred from the
20 percentage change of pollen record can thus be considered reliable.

21 According to modern surface pollen studies, *Pinus* is often overrepresented in pollen
22 records because of its abundant pollen production and the ease with which this pollen
23 is transported over long distances. As a general rule, it can be assumed that there
24 is/was no proximate pine forest if less than 25 to 30% of *Pinus* pollen occurs in
25 samples (Li and Yao, 1990). Higher percentages of Cupressaceae and Taxodiaceae
26 coexistent with temperate tree, shrub and herbaceous pollen may reflect a warmer,
27 wetter and more humid climate (Song, 1978). Nowadays, *Ulmus* is commonly
28 distributed in the sub-humid temperate and warm temperate mountain foothills of

1 northern China, but percentages of its pollen collected from the Chinese Loess Plateau
2 surface soils never exceed 1%, even under broadleaved forests containing elm (Liu et
3 al., 1999). In general, when their abundance exceeds 3–5% of the arboreal pollen total,
4 birch and oak can be considered to be/have been present in woodland (Liu et al.,
5 1999). *Salix* produces very little pollen, and most of this pollen falls near the tree
6 itself (Li et al., 2000). Modern *Artemisia* and Chenopodioideae are extensively
7 distributed throughout the arid and semi-arid regions of China. Chenopodioideae are
8 more drought-resistant than *Artemisia*. Higher percentages of *Artemisia* pollen may
9 reflect a semi-arid grassland environment, while higher percentages of
10 Chenopodioideae pollen may reflect an arid desert environment. Surface pollen
11 analysis shows that *Artemisia* and Chenopodioideae are greatly overrepresented in the
12 pollen rain. Only when Chenopodioideae and *Artemisia* pollen abundance exceeds 30%
13 of the total should their presence be considered as primarily local (Herzschuh et al.,
14 2003; Ma et al., 2008). Poaceae pollen abundance is sparse, usually only 3–6%, even
15 when it represents the dominant modern species (Tong et al., 1995).

16 Our record therefore indicates that, during the period when the Yaodian Fm was being
17 deposited, the study area was covered by temperate forests and a warm and humid
18 climate. Mixed deciduous forests, characterized by the dominance of *Pinus*,
19 Cupressaceae, *Ulmus* and *Quercus*, were distributed within the basin and the low
20 altitude hills surrounding it. Mid- and high-altitude forests with *Abies*, *Picea* and
21 *Cedrus* existed in the surrounding uplands. The river banks or lake margins were
22 colonized by *Salix*, *Alnus*, *Fraxinus* and Taxodiaceae. Cyperaceae, *Typha* and
23 *Myriophyllum* grew along the lake shores or in shallow water areas. Ranunculaceae,
24 Poaceae, Chenopodioideae and *Artemisia*, principally occupied the forest understory,
25 or were distributed in forest clearings. However, as indicated by our record, the
26 environment was not static. During 11.4–10.1Ma, temperate forest grew in the basin
27 indicating a rather humid climate. The growth of fluvial channel deposits and the
28 presentation of a large number of mammalian fossils (Li et al., 2006) also support the
29 theory that much denser vegetation capable of supporting large mammals such as

1 rhinoceroses developed during this interval. Moreover, we know that the northern
2 Tianshui Basin was dominated by temperate and warm-temperate deciduous broadleaf
3 forest (Hui et al., 2011). Our result is also consistent with research into the climatic
4 evolution of the Qaidam Basin, which found that the presence of $\delta^{18}\text{O}$ values
5 characteristic of large mammals indicated a warmer, wetter, and perhaps
6 lower-altitude Qaidam Basin (Zhang et al., 2012). The early Late Miocene mammal
7 fauna discovered in the Qaidam Basin also reflects a wooded environment, in which
8 many streams with aquatic plants such as *Trapa* and *Typha* developed (Wang et al.,
9 2007). From 10.1–7.4Ma, the study area was dominated by a warm-temperate open
10 forest environment and a less humid climate, relative to the previous interval.
11 Sedimentary facies become characteristic of shallow lake deposits (Li et al., 2006).
12 Mammal fauna identified in the eastern Qaidam Basin also indicates that a mixed
13 habitat of open and wooded environments, with abundant freshwater streams, was
14 predominant at that time (Wang et al., 2007). In particular, herbaceous plants also
15 increased their presence in the Tianshui Basin after ~8.6Ma, as confirmed by
16 mammalian fossil records. In the northern Tianshui Basin at ~9.5Ma, there is
17 evidence of a sizeable rhinoceros population, which would have required a relatively
18 moist woodland environment to sustain itself. However, the typical *Hipparion* fauna
19 at ~8.0Ma probably represents a relatively temperate climate with more mixed
20 vegetation, i.e. an open forest environment rather than a vast, open landscape. Large
21 mammals would still have been able to survive in such an environment (Zhang et al.,
22 2013).

23 An open temperate forest-steppe environment developed in the study region,
24 indicating significant aridification after ~7.4Ma. Grassland, composed principally of
25 Poaceae, *Artemisia* and Chenopodioideae, developed in most of the basin, while
26 shrinking areas of open forest, dominated by Cupressaceae, *Ulmus* and *Quercus*,
27 existed in the surrounding mountains. *Salix* continued to grow in relatively humid
28 environments such as riverbanks. Distal floodplain to palustrine deposits now
29 characterized the study area (Li et al., 2006). A sudden increase in magnetic

1 susceptibility after ~7.4Ma may indicate an arid environment (Zhang, 2013) (Fig. 4b).
2 In the northern part of the Tianshui Basin, drought-tolerant *Artemisia* predominated
3 after 7.4Ma, further confirming the presence of a drier climate (Hui et al., 2011) (Fig.
4 4c). Additionally, the growing presence of grazer mammalian species at the end of the
5 Miocene in the Tianshui Basin suggests that the local environment was principally
6 occupied by grassland, with some woodland, and even some deserts (L. P. Liu et al.,
7 2011) (Fig. 4d). Furthermore, the gradual increase in eolian sediments after 7.4Ma in
8 the Linxia Basin would indicate a period of intense desertification in central China
9 (Fan et al., 2006) (Fig. 4e). Biomarker evidence from the Linxia Basin also indicates a
10 distinct change in climate toward arid-cold conditions at ~8Ma (Y. L. Wang et al.,
11 2012). The isotopic compositions of herbivorous fossil teeth and paleosols from the
12 Linxia Basin (Wang and Deng, 2005) and southwestern China (Biasatti et al., 2012)
13 also indicate a shift to a drier, or seasonally drier, local climate. In the Qaidam Basin,
14 *Hipparion teilhardi* fossils are characterized by slenderer distal limbs, and dated to
15 the end of the Miocene, implying an adaptation by this animal to the open steppe
16 environment (Deng and Wang, 2004). Marine sediments also indicate that the climate
17 changed at this time. For example, local seawater $\delta^{18}\text{O}$ reconstructions from ODP Site
18 1146 in the northern South China Sea suggest that the climate of east and south Asia
19 shifted toward more arid conditions after ~7.5Ma (Steinke et al., 2010) (Fig. 4f).

20 **5.2 More arid condition at the end of the Miocene and possible causes**

21 Based on the Late Neogene Chinese mammalian fossils data, Zhang (2006) suggested
22 that mammal communities in northern China were rather stable and uniform from
23 ~13Ma to the end of the Miocene (~7–8Ma), and that differentiation between the
24 humid fauna communities prevalent in eastern China and the dry fauna communities
25 identified in western China occurred after the end of the Miocene. The diversity in
26 Bovidae fossils also increases significantly toward the end of the Miocene, with some
27 genera appearing in southwestern China (Chen and Zhang, 2009), indicating an
28 expansion of grasslands and aridification. Using macro- and microfloral quantitative
29 recovery techniques to reconstruct the climate in northern China at the time, Y.-S. C.

1 Liu et al. (2011) proposed that the west–east temperature and precipitation gradient
2 pattern did not develop in northern China until the end of the Miocene. This
3 corroborates the quantitative results gained from using mammalian fossils as a proxy
4 for paleoprecipitation (Liu et al., 2009). A semi-quantitative reconstruction of Chinese
5 Neogene vegetation also indicated that the aridification of western, central and
6 northern China occurred during the Miocene–Pliocene transition (Jacques et al., 2013).
7 Indeed, in order to adapt to the arid climate of northern China during the end of the
8 Miocene, some plants and arthropods also evolved more arid-tolerant species, such as
9 *Frutescentes* (Fabaceae) (Zhang and Fritsch, 2010), *Ephedra* (Ephedraceae) (Qin et
10 al., 2013) and *Mesobuthus* (Buthidae) (Shi et al., 2013). This marked aridification has
11 been well documented in other parts of Asia. For example, dramatic changes in the
12 carbon isotopic ratio of leaf waxes at ODP Site 722 indicate an increasing aridity at
13 the end of the Miocene in continental source regions, including Pakistan, Iran,
14 Afghanistan, and the Arabian Peninsula (Huang et al., 2007) (Fig. 4g). The isotopic
15 compositions of herbivorous fossil teeth and paleosol carbonates also suggest that the
16 climate became drier over the Indian Subcontinent, China, and Central Asia toward
17 the end of the Miocene (Badgley et al., 2008; Barry et al., 2002; Biasatti et al., 2012;
18 Cerling et al., 1997; Quade et al., 1989; Wang and Deng, 2005; Zhang et al., 2009).
19 The evidential synchronicity of these climatic events in Asia strongly suggests that the
20 aridification of the Asian interior began at the end of the Miocene (~7–8Ma). The
21 onset of such a marked aridification is further corroborated by the presence of red clay
22 across much of the Chinese Loess Plateau (An et al., 2001).

23 Precipitation in arid northwestern China is primarily caused by the Asian Summer
24 Monsoon, whereas the Asian Winter Monsoon promotes a cold and dry climate.
25 Besides the monsoon source, the westerlies also bring precipitation into China.
26 During the Neogene, Eurasia was influenced by global cooling, land-sea redistribution
27 and regional tectonic uplift (Lease et al., 2007; Li et al., 2014; Guo et al., 2008; Miao
28 et al., 2013, 2015; Molnar et al., 2010; Mudelsee et al., 2014; Zachos et al., 2001;
29 Zhang et al., 2007), and these three factors are considered as the major drivers for the

1 formation and evolution of the Asian monsoon and inland arid climate.

2 During the Late Neogene, the most significant global cooling event occurred at
3 ~14Ma (Mudelsee et al., 2014; Zachos et al., 2001), followed by a longer-term but
4 minor cooling trend (4–10Ma, Mudelsee et al., 2014) (Fig. 4h). Although the global
5 cooling should somehow lead to net aridification on the planet, cooling and
6 aridification trends do not seem to run parallel (van Dam, 2006). The complexity of
7 the atmospheric and oceanic circulation systems ensures that general cooling may
8 result in precipitation decrease in some regions and increase in others (van Dam,
9 2006). However, integrated studies showed that the global cooling during the
10 Neogene had significant influences on driving the Asian monsoon and inland arid
11 climate (e.g. Lu et al. 2010; Lu and Guo, 2014; Tang and Ding, 2013), especially
12 since the Late Miocene (Lu and Guo, 2014). The possible mechanism lies in two
13 aspects. Firstly, it is clear that the global cooling has strengthened the Siberia High,
14 which dominates winter monsoon circulation and aridity in eastern Asia (Lu and Guo,
15 2014). This would result in enhanced and more frequent cold surges in the
16 mid-latitudes of Northern Hemisphere. Secondly, the global cooling caused the
17 weakening of hydrological cycle, expanding of ice sheets, lowering of sea level and
18 increasing of continental surface (Lu and Guo, 2014; Tang and Ding, 2013). This
19 would reduce the moisture mass transported into the continental interior (Tang and
20 Ding, 2013). Therefore, we speculate that the global cooling could intensify aridity of
21 the Asian interior.

22 Besides the above focusing on the climate effects of the global cooling, model
23 simulations have paid special attention to the climatic effects of the land-sea
24 redistribution. For example, model simulations suggest that the westward retreat of
25 the Paratethys from central Asian has contributed significantly to Asian climates (e.g.
26 Guo et al., 2008; Ramstein et al., 1997; Zhang et al., 2007). However, a large number
27 of geological evidences suggest that the vast majority/even all Paratethys regression
28 from the Tarim Basin (northwest China) occurred at the Oligocene (e.g. Bershaw et
29 al., 2012; Bosboom et al., 2014). Meanwhile, numerical simulation also indicates that

1 the spreading of the South China Sea may enhance the south-north contrast of
2 humidity in China (Guo et al., 2008), and brings more precipitation into Asia.
3 Nevertheless, many studies indicate that western and northern China became drier
4 during the Neogene (e.g. Guo et al., 2008; Tang and Ding, 2013; Sun and Wang,
5 2005). Therefore, although the land-sea redistribution had a significant impact on the
6 major climate reorganization in Asia during the Late Oligocene/Early Miocene (Guo
7 et al., 2008; Zhang et al., 2007), it should have a limited effect on the formation and
8 development of the Asian inland arid climate during the Late Miocene.

9 Model simulations have also paid attention to the climate effects of the TP uplift. The
10 scenarios of whole-plateau uplift (e.g. Kutzbach et al., 1993), phased uplift (e.g. An et
11 al., 2001; Kitoh, 2004; Liu and Yin, 2002) and sub-regional uplift (e.g. Boos and
12 Kuang, 2010, 2013; Chen et al., 2014; Tang et al., 2011, 2013; Wu et al., 2012), with
13 increasing complexity, are usually designed for discovering the cause-effect relations
14 between the plateau uplift and paleoclimate change (Liu and Yin, 2011). The different
15 models conclude that the uplift of the TP played an essential role in affecting the
16 atmospheric circulation and forming the monsoon and arid climate when the
17 whole/sub-regional plateau exceed a critical height (An et al., 2001; Boos and Kuang,
18 2010, 2013; Chen et al., 2014; Kutzbach et al., 1993; Liu and Yin, 2002; Tang et al.,
19 2011, 2013; Wu et al., 2012). However, because of the different model setups and
20 boundary conditions, there still exist many uncertainties in the different forms of the
21 plateau uplift forcing and regional climatic responses (Liu and Yin, 2011). The
22 geological/proxy research can provide the constraints for the model boundary
23 conditions, whereas numerical simulation can test the geological/proxy result.
24 Therefore, it is useful to compare the geological/proxy results and the numerical
25 simulations (Micheels et al., 2007, 2011). Many geological studies have suggested
26 that the TP experienced rapid uplift during the interval ~8–10Ma (e.g. Enkelmann et
27 al., 2006; Fang et al., 2003, 2005; Lease et al., 2007; Li et al., 2014; Molnar et al.,
28 2010; Wang et al., 2006; X. X. Wang et al., 2012; Zheng et al., 2006, 2010) (Fig. 4i),
29 but the timing and degree of the uplift are still debated. The Late Miocene uplift

1 would have achieved an altitude sufficient to block the penetration of moisture from
2 the source region into western China (Dettman et al., 2001, 2003). There are also
3 increasing proxy evidences that the Asian Summer Monsoon weakened after ~10Ma
4 (e.g. Clift et al., 2008; Wan et al., 2010), while the Asian Winter Monsoon
5 strengthened, particularly toward the end of the Miocene (e.g. An et al., 2001; Clift et
6 al., 2008; Jacques et al., 2013; Jia et al., 2003; Sun and Wang, 2005), implicating the
7 intensified Asian inland aridification. It is consistent with the most model simulations
8 that aridity of the Asian interior will be intensified along with the uplift of the TP.
9 However, it should be noted that there is no doubt regarding the effects of the global
10 cooling on the general trend toward a dry climate in the Asian interior.

11 **6 Conclusion**

12 The Late Cenozoic basins, located at the northeast TP, document the environmental
13 changes associated with tectonic uplift and global cooling. We investigate a Late
14 Miocene pollen record from the Tianshui Basin. Our results indicate that a temperate
15 forest with a rather humid climate regime (11.4–10.1Ma), gave way to a temperate
16 open forest environment with a less humid climate (10.1–7.4Ma); this was in turn
17 replaced by an open temperate forest-steppe landscape, accompanied by a relatively
18 arid climate (7.4–6.4Ma). The vegetation succession demonstrates that the
19 aridification of the Asian interior occurred after ~7–8Ma, as corroborated by other
20 studies of Asia. Our findings support the idea that the long-term global cooling and
21 the TP uplift caused the Late Miocene aridification of the Asian interior.

22 *Acknowledgements.* We thank Q. Y. Cui and Y. Z. Ma for their early pollen work, and
23 Dr. L. Dupont and an anonymous reviewer for their valuable comments and
24 suggestions. This work was co-supported by the State Key Program of National
25 Natural Sciences of China (grant no. 41330745), the (973) National Basic Research
26 Program of China (grant no. 2013CB956403) and the National Natural Science
27 Foundation of China (grant nos. 41301216, 41272128 and 41201005).

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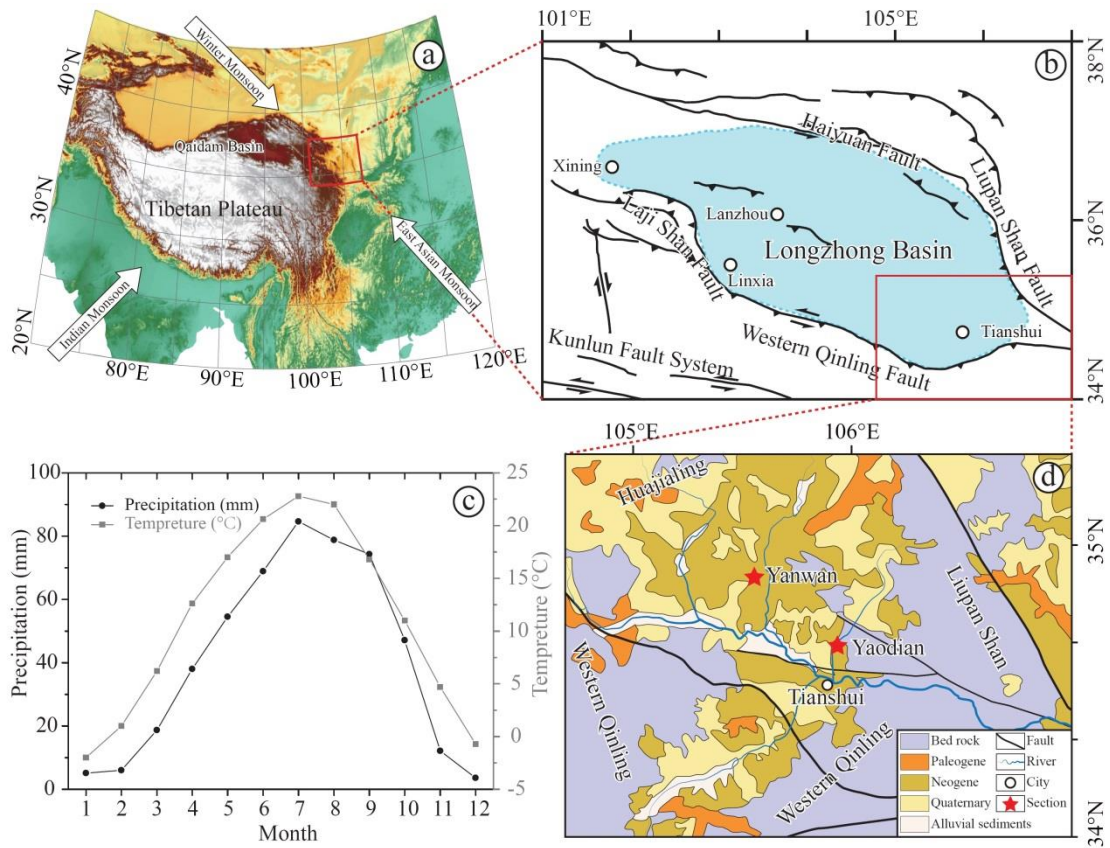
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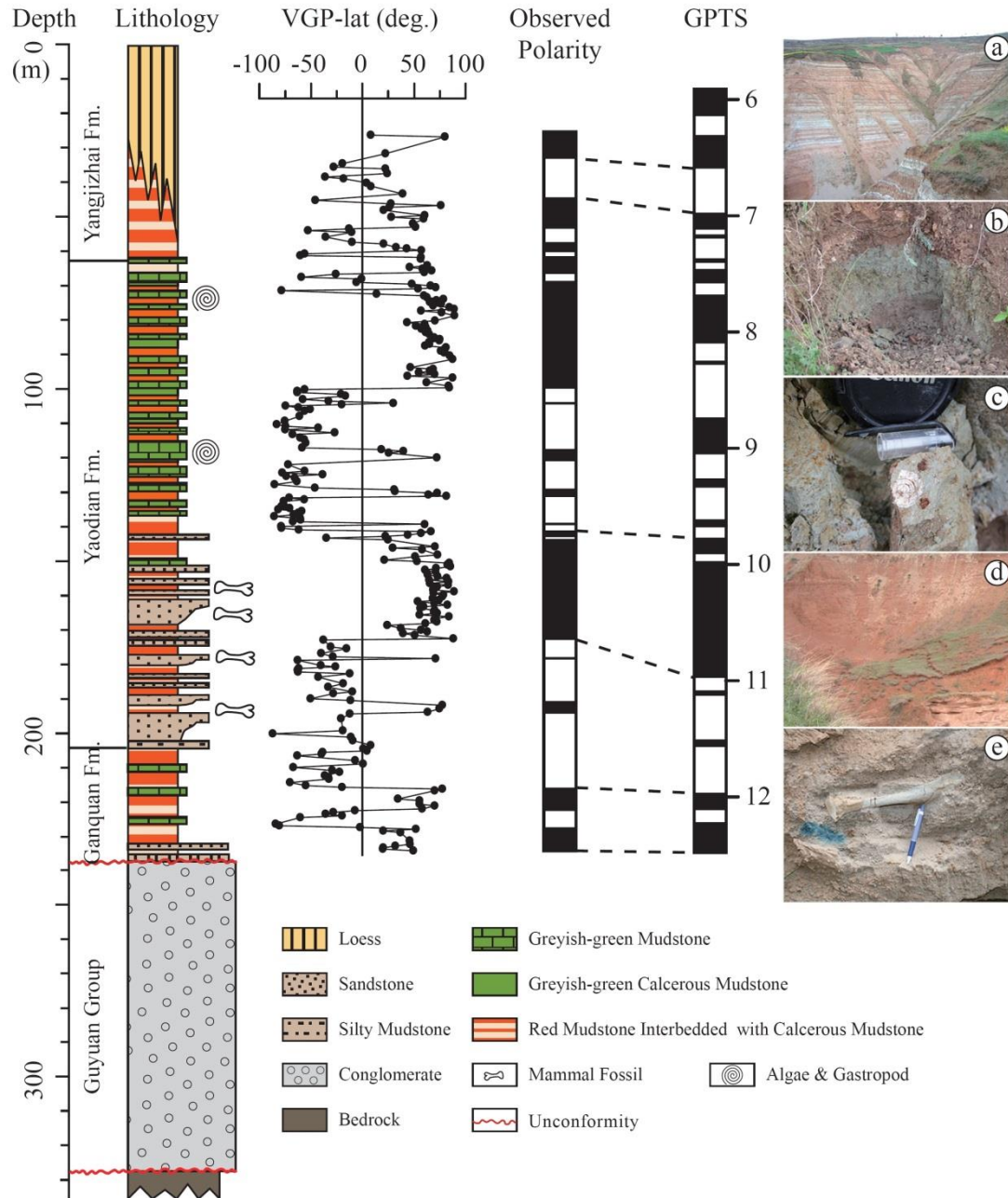
2 Figure 1. Geographic setting of Yaodian Section. (a) Location of the Longzhong Basin.

3 (b) Major tectonic faults of the Longzhong Basin. (c) Mean monthly temperature and

4 mean monthly precipitation in the Tianshui area, 1971-2000. (d) Geological map of

5 the Tianshui Basin.

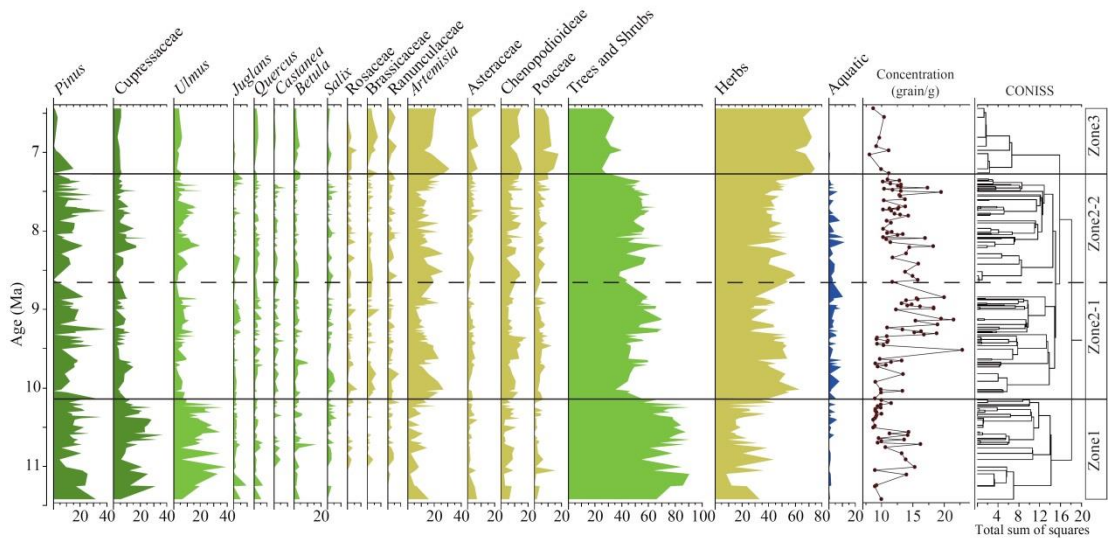
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2 Figure 2. Lithology and magnetic stratigraphy of the Yaodian Section (according to Li
 3 et al., 2006). (a) The entire Yaodian Fm. (b) Yangjizhai Fm distal floodplain to
 4 palustrine deposits. (c) Yaodian Fm upper stratum lacustrine deposits, containing
 5 gastropod fossil fragments. (d) Yaodian Fm middle stratum floodplain deposits, with
 6 paleosols. (e) Yaodian Fm lower stratum fluvial channel deposits, containing
 7 fossilized animal bones. GPTS, standard geomagnetic polarity timescale in million
 8 years (Ma).

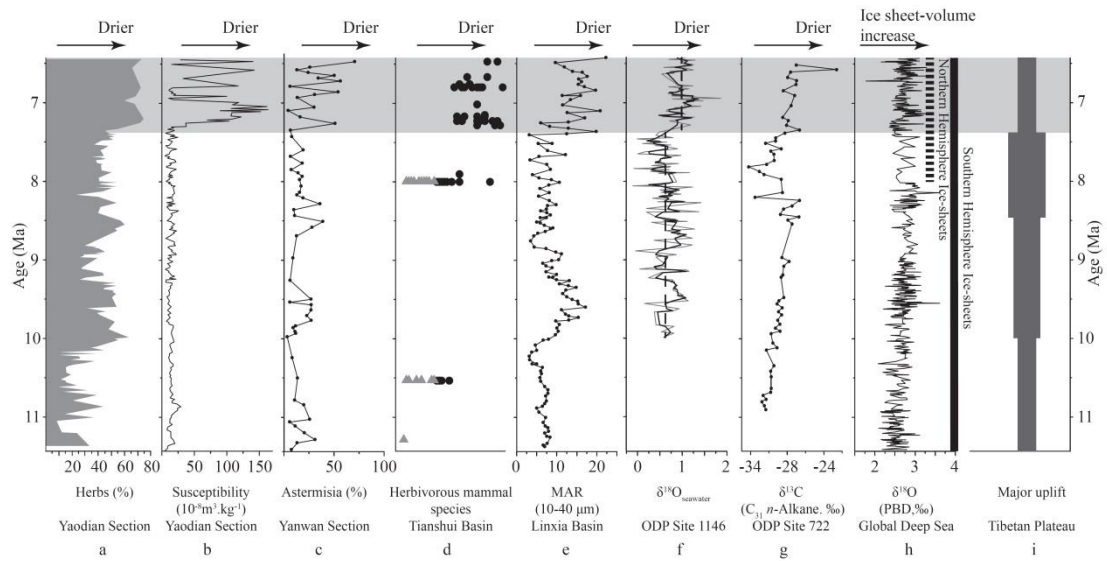
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2 Figure 3. Histogram showing pollen percentages for the most significant angiosperms
 3 and gymnosperms.

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2 Figure 4. Proxy records of aridification for East Asia during the Late Miocene. (a)

3 Herbaceous pollen percentage for the Yaodian Section (this study). (b) The magnetic

4 susceptibility of the Yaodian Section (Zhang, 2013). (c) Drought-tolerant *Artemisia*

5 pollen percentage in the Yanwan Section, northern Tianshui Basin (Hui et al., 2011).

6 (d) Herbivorous mammal species in the Tianshui Basin (Guo et al., 2002; L. P. Liu et

7 al., 2011; Li et al., 2006; Zhang et al., 2013). Black circles represent species which

8 adapted to relatively arid environments, including *Sinocricetus zdanskyi*, *Kowalskia*

9 *sp.*, *Hansdebminjia pusillus*, *Lophocricetus grabau*, *Mesosiphneus praetingi*, *M. sp.*,

10 *Paralactaga anderssoni*, *Parasoriculus sp.*, *Prosiphneus eriksoni*, *P. licenti*, *P.*

11 *tianzuensis*, *Prospermophilus orientalis*, *Pseudomeriones abbrevuuuus*, *P.*

12 *complicidens*, *Sicista sp.*, *Alilepus annectens*, *Allorattus sp.*, *Apodemus sp.*, *Chardina*

13 *sp.*, *C. sinensis*, *C. truncatus*, *Chardinomys nihowanicus*, *C. sp.*, *C. yusheensis*,

14 *Pliosiphneus lyratus*, *Cricetinus mesolophidus*, *Mimomys teilhardi*, *Sinotamias sp.*,

15 *Ochotona gracilis*, *O. lagreli*, *O. lingtaica*, *O. minor*, *O. plicodenta*, *O. sp.*,

16 *Ochotonoma sp.*, *O. primitiva*, *Trischizolagus mirificus*, *Hipparion chiai*, *H.*

17 *dermatorhinum*, *H. fossatum*, *H. plocodus*, *H. sp.*, *H. weihoense*, and *Gazella sp.*; grey

18 triangles represent species which adapted to relatively humid environments, including

19 *Chleuastochoerus stehlini*, *Cervavitus novorossiae*, Cervidae gen. et sp. Indet.,

20 *Palaeotragus microdon*, *P. sp.*, *Samotherium sinense*, *S. sp.*, Rhinocerotidae indet.,

21 *Acerorhinus fuguebsis*, *Chilotherium habereri*, *C. sp.*, *C. wimani* and *Protanancus*

1 *tobieni*. (e) Eolian sediment mass accumulation rates in the Linxia Basin, northeastern
2 TP (Fan et al., 2006). (f) South China Sea $\delta^{18}\text{O}_{\text{seawater}}$ estimate from ODP Site 1146
3 (Steinke et al., 2010). (g) Carbon isotope ratios of leaf wax C_{31} *n*-alkane extract from
4 ODP Site 722 (Huang et al., 2007). (h) Compiled global deep-sea $\delta^{18}\text{O}$ values (Zachos
5 et al., 2001, the data available online
6 <http://www.es.ucsc.edu/~jzachos/Publications.html>. A new compilation has been
7 published by Mudelsee et al. (2014), which is congruent with the Zachos' curve for
8 the Miocene part). (i) Schematic model showing the major periods of TP uplift
9 (Enkelmann et al., 2006; Fang et al., 2003, 2005; Lease et al., 2007; Li et al., 2014;
10 Molnar et al., 2010; Wang et al., 2006; X. X. Wang et al., 2012; Zheng et al., 2006,
11 2010).