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Mio-Pleistocene Zanda Basin biostratigraphy and geochronology, pre-Ice Age fauna, and mammalian evolution in western Himalaya

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The Pliocene (5.3–2.6 Ma) of Tibet witnessed the drying of the northern Tibetan Plateau and the approach to the Pleistocene Ice Age within the background of intensifying Indian and East Asian monsoons. Yet little is known about Pliocene mammals living on the high Tibetan Plateau despite the fact that fossil mammals elsewhere constitute an important knowledge base for terrestrial environments. The late Miocene to Pleistocene Zanda Basin at the northern foothills of the Himalayas affords a welcome opportunity to evaluate the biological response to environmental change at high elevations. Abundant, well-preserved fossil mammals and fish from an 800-m continuous section of fine- to coarse-grained sediments thus open a rare window into a past biological world. For example, the discovery of an ancestral wooly rhino from Zanda Basin that was the precursor of its late Pleistocene megafaunal descendants leads to our "out-of Tibet" hypothesis, suggesting that the high Tibetan Plateau was a Pliocene cradle for Ice Age cold adaptations.

In this paper, we document in detail the mammalian biostratigraphy, chronology, and paleozoogeography based on Zanda Basin fossil mammals. Our high-resolution biostratigraphy and biochronology offer for the first time independent constraints that both support and modify recent magnetostratigraphic correlations. Using characteristic Pliocene and Pleistocene mammals, particularly the small mammal assemblages in the lower part of the section and monodactylid Equus from the upper section, we propose a correlation to C1n to C3An.1r, with an age range of ~400 Ka to 6.4 Ma.

Within the 800-m Zanda section, the lower 0–150 m is of latest Miocene age, spanning 6.4–5.3 Ma. Sparsely fossiliferous, the lower section has produced five taxa so far: Ochotona, Panthera, Ourlianoria, Palaeotragus, and Hipparion—all are consistent with a late Miocene age. The middle 150–620 m section spans the entire Pliocene. This section is by far the most fossiliferous, including such typical Pliocene small mammals as Prosiphneus, Mimomys, Apodemus, and Trischizolagus, as well as large mammals such as Coelodonta thibetana, Hipparion zandaense, Chasmaporthetes, Nyctereutes, Meles, Antilospira, and others. In the upper 620–800 m section the fossils are rare, but do include characteristic Pleistocene taxa such as Equus.

Zoogeographically Zanda Basin mammals are a mixture from two major sources. Taxa such as Mimomys, Prosiphneus, Trischizolagus, Chasmaporthetes, Nyctereutes, Meles, and Xenocyon are commonly found in north China or east Asia. In contrast, several forms, such as unique species of pikas (Ochotona), squirrels (Aepyosciurus), and ancestral Tibetan antelope (Qurliqnoria), seem to belong to an indigenous Tibetan fauna evolved within the plateau. A lack of shared taxa with the Oriental Realm suggests a formidable barrier by the Himalayas despite a short distance (~100 km) between Zanda Basin and the Indian subcontinent.

1. Introduction

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The Tibetan Plateau has been widely recognized as central to our understanding of the evolution of the Indian and Eastern Asian monsoon

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systems, and debates are on-going about the timing of uplift process and its role in initiating changes in climatic regime (e.g., An et al., 2001; Molnar, 2005; Boos and Kuang, 2010; Quade et al., 2011). Until fairly recently direct evidence of the effect of this changing regime on the biological community, especially vertebrates, has been sparse and has played a limited role in the debate (e.g., Ji et al., 1980; Huang and Ji, 1981). To a large extent, this is the result of the recent start of paleontological research efforts and a lack of suitable basins with abundant fossils. Added to this are difficulties in physical access to mountainous terrains and political sensitivity.

Zanda (Zhada) Basin offers an exceptional opportunity to remedy this shortcoming and to begin building an excellent fossil record to test relationships of paleoenvironments and vertebrate evolution in a crucial region of the Tibetan Plateau. The basin's location in the northern foothills of the western Himalaya, at high elevations, and at the headwaters of the Indus River contributes to a unique perspective in the paleozoogeography of its ancient faunas. Nearly four months of field work over five seasons (2006–07, 2009–10, 2012) demonstrate that this is the richest basin on the Tibetan Plateau in terms of abundance of fossil material, areas of potential exposures, and continuity of the fossil record. Archived in well-preserved and extensively exposed late Miocene to Pleistocene sediments, the Zanda fossil assemblage fill a critical void in the late Cenozoic, augmenting heretofore limited knowledge of Pliocene vertebrate faunas in southern Tibet and providing a much needed window into the past paleoenvironment and its effect on biological evolution.

Vertebrate paleontology in Zanda Basin significantly advances our understanding of the following aspects of the history of Tibet: 1, vertebrate communities and diversity in the late Cenozoic; 2, mammalian biochronology and magnetostratigraphy; 3, zoogeographic origins of individual components in Tibetan Plateau faunas; 4, paleoclimate records as preserved in isotopes of vertebrate teeth and bones; 5, paleoenvironments and plant communities as related to the diet of mammalian herbivores; 6, evolution of community structures in mammals and fishes; 7, relationship of cold adaptation in high elevations and for Ice Age megafaunas; and 8, drainage evolution of the Indus River system.

Our discovery of an ancestral wooly rhino from Zanda Basin made the unexpected link to its Ice Age descendants in northern Eurasia, establishing Tibet as a possible cradle of evolution for the Ice Age megafauna (Deng et al., 2011). The implications of this discovery highlight the importance of understanding major players in the vertebrate faunas and paleoenvironmental conditions during the Pliocene of Tibet, prior to the Ice Age. In this paper, we document in detail the biostratigraphic basis of our "out of Tibet" hypothesis. Our high resolution biostratigraphy lays the foundation for a chronologic framework, on which subsequent descriptions of individual taxa can be built, and will serve as an important reference point in the late Miocene to Pleistocene of Tibet.

2. Material and methods

All vertebrate fossils collected belong to and are housed in the Institute of Vertebrate Paleontology and Paleoanthropology (IVPP), Chinese Academy of Sciences. All Zanda fossil localities begin with an abbreviation of "ZD" followed by the last two digits of the year collected and the field number of that year, such as IVPP locality ZD0604 (04 locality in 2006). For east Asian biochronology, we use the latest proposal of the Chinese land mammal age/stage system by Qiu et al. (in press-b). We adopt the ATNTS2004 Geomagnetic Polarity Time Scale (GPTS) of Lourens et al. (2004) in our magnetic correlations. We define the Pliocene–Pleistocene (Neogene–Quaternary) boundary at C2r–C2An (2.581 Ma), as was recently endorsed by the International Commission on Stratigraphy (Mascarelli, 2009).

We devised a system of using elevation data to correlate individual localities with a master magnetic section in south Zanda. We systematically took GPS readings for all fossil localities, in many cases using the averaging function to achieve greater accuracy. The elevation data are interpolated into individual legs of the nearest measured sections by Saylor and colleagues (Saylor, 2008; Saylor et al., 2010b). We then adopted the lithostratigraphic and sequence stratigraphic correlation scheme by Saylor et al. (2010b) to tie together individual sections to a standard south Zanda magnetic section. The greatest advantage of such a system is that it permits us to closely tie all of our localities with six major measured sections, plus six other smaller partial sections. In most cases, the measured sections are in the same accessible canyons where fossils were collected, and distances of fossil localities to section legs are as small as a few hundred meters to as large as a few km. Because the basin is undeformed and bedding is horizontal, this method of lateral interpolation permits reasonable precision. The biggest source of systemic errors is probably GPS accuracy. In mountainous terranes, such as Zanda Basin, horizontal errors of 6–10 m or less by commercial grade GPS units can be routinely achieved, but errors of vertical measurements can be twice as much, or more. Such systemic errors can translate to an error in age estimates of \pm 0.2 million years (assuming a 1 m/10,000 year sedimentation rate), a tolerable range given the common rate of mammalian speciation of greater than 1 myr (Blois and Hadly, 2009; Uyeda et al., 2011). Our calibration system can be improved by future magnetic studies on individual canyons, in combination with our locality database.

3. Previous paleontologic studies

Physical inaccessibility, political barriers, and the lack of a "dragon bone" hunting tradition by native Tibetans combined to make Zanda Basin practically unknown paleontologically until quite recently, despite the fact that the basin is one of the richest in vertebrate fossils on the Tibetan Plateau. Hints of the existence of fossil mammals first came to the fore in a report on a maxillary of an extinct giraffe Palaeotragus collected from Xiangze Farm area (now Jiade Farm) in 1976 by a group of geologists from a multidisciplinary Qinghai–Tibetan Plateau expedition by the Chinese Academy of Sciences (Zhang et al., 1981). News of vertebrate fossils inspired another geologist, Liang Dingyi, to make a trip himself in 1982, which led to the acquisition of a partial skull and jaws of a three-toed horse near the village of Daba. This specimen was later described as a new species Hipparion zandaense by Li and Li (1990). In the early 2000s, another team of geologists from the Institute of Geomechanics (Chinese Academy of Geological Sciences) made miscellaneous discoveries of a rhinoceros foot bone and a pika cheek tooth (Meng et al., 2004, 2005) that were too poorly preserved to be of significance.

All previous fossil discoveries were chance encounters of isolated specimens by geologists: our 2006 field season was the first documentation of the basin by vertebrate paleontologists. Our paleontologic explorations, spanning five field seasons (in 2006, 2007, 2009, 2010, 2012), thus represent the first systematic effort to assess the vertebrate fossil record in the basin. It soon became clear that Zanda Basin is ideally suited for paleontologic explorations: within our five short field seasons, several hundred specimens have been collected through much of the strata. However, only a new wooly rhinoceros and a partially preserved articulated skeleton of a three-toed horse have so far been adequately described (Deng et al., 2011, 2012), as well as a brief mention of the Zanda fauna in a summary paper on Tibet mammal biochronology (Wang et al., in press-a, in press-b). This report presents, for the first time, a detailed framework of biostratigraphy and geochronology, and full descriptions of individual taxa will follow in separate treatments.

Biostratigraphically, the lack of previous paleontologic explorations in Zanda Basin has a positive effect. Lacking historic "baggage" (such as in numerous Chinese basins with rich "dragon bone" hunting traditions, where fossils were acquired with questionable or no stratigraphic context) we are able to place every collected specimen in precise stratigraphic context. The only exceptions are two previously described specimens of Palaeotragus microdon (Zhang et al., 1981) and Hipparion zandaense (Li and Li, 1990) that lack detailed locality information. We are thus able to place all fossils in a modern stratigraphic context with a high resolution and consistency rarely seen in other Chinese basins.

4. Geologic and environmental setting

The Zanda Basin is a late Miocene through Pleistocene pull-apart sag basin located just north of the high Himalayan ridgecrest in the westcentral part of the orogen (Fig. 1). Flanked by the Himalaya Range to the southwest and Ayilariju Mountain to the northeast, basin sediments can be seen as far southeast as 31°06′ N 80°35′ E near the village of Qulong in the upper reach of the Langqên Zangbo (Sutlej) River and as far northwest as the Qusong area; current outcrop extent of the basin fill is $>$ 9000 km².

The basin occupies a region with an extended tectonic history. To the northeast of the basin, the Oligo-Miocene Great Counter Thrust (GCT), a south-dipping, top-to-the-north thrust system, modifies the Indus Suture (e.g., Ganser, 1964; Yin et al., 1999; Murphy and Yin, 2003). The South Tibetan Detachment System (STDS), southwest of the basin, is an earlymid Miocene (Hodges et al., 1992, 1996; Searle et al., 1997; Murphy and Harrison, 1999; Hodges, 2000; Murphy and Yin, 2003; Searle et al., 2003; Yin, 2006; Cottle et al., 2007) series of north-dipping, low-angle, top-to-the-north normal faults. Zanda Basin sediments onlap, and hence post-date movement on, both the STDS and the GCT. Ongoing exhumation of the Gurla Mandhata metamorphic core complex to the southeast of the basin began at ~9 Ma (Murphy et al., 2002). Qusum (Leo Pargil) metamorphic core complex to the northwest basin began at 14–16 Ma (Thiede et al., 2006) and these faults cut Zanda Basin sediments (Saylor et al., 2010a). The axis of the basin is approximately northwest–southeast, parallel to the general arc of the Himalaya. Tectonic control, basin geometry, basin sedimentology, depositional environment, drainage evolution, stable isotopes, paleoclimates, and elevation history have been explored during the recent decade (Saylor, 2007; Saylor et al., 2007a, 2007b; Saylor, 2008; Saylor et al., 2008; Wang et al., 2008a, 2008b; Kempf et al., 2009; Saylor et al., 2009, 2010a, 2010b; Quade et al., 2011; Y. Wang et al., 2011, 2012).

With an elevation of 3700–4500 m, modern Zanda Basin has an annual mean temperature of close to 0 °C (based on data from nearby Shiquanhe in National Climatic Data Center, 2012). Located at the western end of the Himalaya and on its northern flank (~31–32° N, 79– 80° E), this region has a relatively dry summer created by long distance from and degradation of the combined effects of Indian and Southeast Asian monsoons coming from the east and southeast (Böhner, 2006; Bolch et al., 2012), although summer rains account for ~80% of the annual precipitation in the area (Li, 2006; Tian et al., 2007). Winter, however, is relatively wet compared to eastern Himalaya because of the stronger influence of the winter Westerlies. If a similar monsoon system operated in western Himalaya during the Pliocene, then relatively heavier winter snow and a drier summer, compared to the eastern Himalaya, would be expected. Such a climatic regime seems to be a major factor on modern erosion, causing steeply incised canyons, and would have an impact on the evolution of pre-Ice Age Tibetan faunas.

5. Depositional setting and fossil preservation

Zanda Basin sediments show cyclical changes that may be attributed to Milankovitch forcing (Saylor et al., 2010b). The sedimentary basin fill is undisturbed and lies in angular or buttress unconformity with the underlying deformed Tethyan Sedimentary Sequence (TSS) strata. After deposition, incision by the Langqên Zangbo River exposed the entire basin fill. The basin fill consists of ~800 m of fluvial, lacustrine, eolian and alluvial fan deposits. The lower part of the section consists of ~200 m of trough cross-bedded sands and well-organized, imbricated, pebble to cobble conglomerate. Associated sedimentary structures include stacked, 3–4 m sand–gravel filled channels and longitudinal bars. We interpret these features as fluvial deposits laid down by largescale rivers ancestral to the Sutlej or Indus based on provenance and

Fig. 1. Map of known basins producing fossil vertebrates in the Tibetan Plateau. Political boundaries in the maps do not imply any opinion concerning the legal status of any country or territory.

paleocurrent orientation data (Saylor et al., 2010a). Interbedded finegrained sand and silt horizons showing extensive soft-sediment deformation and containing abundant mammal, gastropod and plant macrofossils are interpreted as marshy bog or overbank deposits within a low-gradient fluvial setting. The middle unit (approximately 250 m) consists of an upward coarsening succession of lacustrine progradational parasequences. Individual parasequences are up to 17 m thick and range between profundal lacustrine claystones and deltaic and wave-worked sediments, including evidence of occasional desiccation. The top 350 m continues the upward coarsening progression displayed in the middle portion but becomes much coarser. The profundal lacustrine facies disappears and is replaced by deltaic or lake-margin deposits. Individual parasequences vary between deltaic or lake-margin and alluvial-fan and fan-delta conglomerates.

Since the initial establishment of the Zanda Formation as a lithological unit (Zhang et al., 1981), additional formation (such as the Tuolin and Xiangze formations) or even group (Zanda Group) names were proposed (Zhu et al., 2005), often based on a perceived depositional hiatus that later proved to be false (Wang et al., 2008b; Saylor et al., 2009). Here we follow Saylor et al. (2009, 2010a, 2010b) and use a single unit, Zanda Formation, for the entire basin sequence.

Vertebrate fossils are present in most fine- to medium-grained sediments in most accessible exposures that are not too steeply incised (Fig. 2). Fossil fishes are found throughout the middle unit, particularly in fine-grained sediments (Figs. 3, 4). Fossil mammals, especially large mammals, are most abundant in near-shore facies and often closely associated with exposures of basement rocks, which often formed islands or the shore line of the Zanda paleo-lake (Fig. 2). We have systematically searched for fossils in all exposures that are approachable by off-road vehicles. Nonetheless our limited efforts were mostly concentrated in areas of high promise, based on satellite images, and are far from an exhaustive search.

The high-energy conglomerate units from the upper and lower portions of the Zanda Formation are mostly unfossiliferous except for a handful of Pleistocene localities producing large mammal fragments. Our prospecting was thus mainly focused in the median to fine-grained unit in the middle section. However, the very fine-grained 220–320 m section mainly produces fish (Fig. 4). With the exception of a partially articulated wooly rhinoceros skeleton from IVPP locality ZD1055 and occasional partial fish skeletons, the vast majority of mammal and fish fossils occur as single elements or a few associated bones or teeth. Partially articulated specimens are uncommon. In most cases the distance of transport is probably short. Concentrated fossil occurrences are exceedingly rare; we found only two sites (IVPP locality ZD1001 and ZD1208), by far our richest, in our five years of exploration that are worthy of extensive quarrying or dry-screening and IVPP locality ZD1001 has a concentration of both large and small mammals, apparently by a fluvial process.

6. Biostratigraphy and biochronology

Despite being in a tectonically active area, Zanda Basin strata underwent minimal post-depositional deformation. Bedding is flat-lying throughout most of the basin, and because of the dominance of shallow lake sediments, most bedding planes are perfectly horizontal and can be traced laterally for long distances. With deep incisions in the majority of exposures, visual correlation is possible in much of the basin. We took advantage of this unique preservation and devised a system of correlations using GPS elevations to tie all fossil localities to a master paleomagnetic section (see Section 2, Materials and methods).

However, such steeply incised canyons in the majority of exposures are not ideal for preservation and collection of fossils. Near-vertical cliffs not only are inaccessible for prospecting but also do not accumulate fossils when they are exposed and erode out. A total of 240 vertebrate fossil localities have been discovered during the past five seasons (Table 1). Of these, three are small mammal sites (IVPP localities ZD0609, 0904, 1001) yielding multiple taxa. Most of the rest of the localities are large

mammals or fish bones, often yielding one or at most a few taxa. Preservation of different size classes of vertebrates is dependent on water energy and closeness to the paleo-lake shore. In addition to the preservational constraints, discovery of fossil sites is also dependent on access to relatively flat or gently sloping exposures.

Fossil mammals are generally best preserved in the middle part of the Zanda Formation at about the 170–600 m level (Fig. 4). With relatively high diversity, mammals from this segment also offer the best chronologic constraints, and the following discussions are mostly based on this assemblage. The lowest stratigraphic small mammal locality is IVPP locality ZD0609 at 174 m (from the base of the section). Of biochronologic significance are Nannocricetus, Mimomys, Apodemus, and Trischizolagus mirificus. Nannocricetus is present in early late Miocene through Pliocene of Inner Mongolia. Zanda Basin specimens of Nannocricetus are too rare to be precisely compared at the moment. Teeth of Mimomys are also rare. However, they are unlikely to be ancestral forms such as Promimomys or lophodont taxon such as Microtodon, the latter occurring in late Miocene. Represented by three upper first molars and one lower molar, crown heights in Zanda Basin Mimomys are most comparable to Mimomys (Aratomys) bilikeensis from early Pliocene Bilike locality of Inner Mongolia, which is the earliest representative of arvicoline rodents in China (Qiu and Storch, 2000). Arvicoline rodents first appeared in the early Pliocene of western Siberia (Repenning, 2003) and shortly afterward dispersed to northern Asia (Qiu and Storch, 2000), Europe (Fejfar et al., 1997; Chaline et al., 1999), and North America (Repenning, 1987; Lindsay et al., 2002; Bell et al., 2004). Apodemus has a long stratigraphic span, ranging from late Miocene to present day of Zanda Basin. Finally, T. mirificus is a common element at Bilike, Inner Mongolia, which is conventionally regarded as one of the earliest Pliocene faunas in north China (Qiu and Wang, 1999; Qiu et al., 2006; Z.-D. Qiu et al., in press). Zanda T. mirificus is indistinguishable from the Bilike form both in size and morphology, and this species is presently known in Bilike and Zanda only. Therefore, of the above four small mammals, Mimomys and T. mirificus offer the strongest chronologic constraint. Although it is possible that the Zanda Mimomys predates all other records of the genus in Eurasia, we think this is unlikely given a well-established record elsewhere. We thus prefer an age of earliest Pliocene to latest Miocene (but see additional discussion in Section 7, Magnetostratigraphy).

Large mammals are also mostly characterized by Pliocene elements. Carnivorans, such as Chasmaporthetes (IVPP localities ZD0636, 0908, 1029), Pliohyaena (=Pliocrocuta) (IVPP locality ZD1208), Vulpes (IVPP localities ZD1001, 1055), Nyctereutes (IVPP localities ZD0624, 1208), and Meles (IVPP localities ZD1001, 1004, 1208), are all typical Yushean taxa, even though most of these are stratigraphically 30 to more than 200 m higher than the lowest Mimomys horizons (IVPP locality ZD0609). The Asiatic first occurrences of these genera, some of them (such as canids) as immigrants from North America, are mostly confined in Pliocene, although occasional late Miocene records have been suggested elsewhere (Qiu, 1987; Qiu and Tedford, 1990; Tedford and Qiu, 1991; Qiu et al., 2004; Tedford et al., 2009). Other than some Miocene left overs, such as Qurliqnoria, ungulates collectively have a Pliocene characteristic as well.

Stratigraphically below IVPP locality ZD0609 (174 m), there are only a handful of localities producing five mammal taxa: Ochotona, Panthera, Qurliqnoria, Palaeotragus, and Hipparion. These are known in late Miocene of either north China (Ochotona, Palaeotragus, Hipparion) (Z.-X. Qiu et al., in press) or Tibetan Plateau (Qurliqnoria) (Wang et al., in press-b), or there are no prior record (Panthera). Mammals from the lower section thus offer no strong age constraint, except being consistent with a late Miocene age.

Fossil localities above the 620 m level are even fewer. Only two identifiable taxa are present: Equus and an advanced cervid. Equus, in particular, is a North American immigrant first appearing at the beginning of the Pleistocene and widely seen in northern China (Qiu, 2006) and in south Asia and Europe (Lindsay et al., 1980). Despite the paucity of fossils, the upper 180 m of Zanda Formation must thus be Pleistocene in age.

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Fig. 2. Map showing important fossil localities and geographic locations discussed in the text. Vertebrate fossil sites are frequently adjacent to exposures of Tethyan basement rock (here colored in brown).

Fossil fishes occur throughout the Zanda section in fine-grained fluvial or lacustrine sediments. Studies of Tibetan fossil fishes in the Pliocene are just beginning (Chang et al., 2008; Wang and Chang, 2010, 2012). They may be of great value in paleoenvironmental, evolutionary, and zoogeographic pursuits, but so far cannot offer any insight into the geochronology.

7. Magnetostratigraphy

There have been at least three independent paleomagnetic studies of the Zanda Basin strata during the past 13 years (Qian, 1999; Saylor, 2008; Wang et al., 2008b). In addition, Zhu et al. (2005) and Meng et al. (2004) alluded to a fourth paleomagnetic section of their own, but no 86 X. Wang et al. / Palaeogeography, Palaeoclimatology, Palaeoecology 374 (2013) 81–95

Fig. 3. Histogram showing relative abundance of vertebrate fossil localities along the 800 m section. Individual fossil localities are interpolated into six nearest measured sections, which are in turn correlated to the main south Zanda section by lithostratigraphic and sequence stratigraphic means (Saylor et al., 2010b).

documentation was provided. All three published studies obtained an $800 + m$ section for the total thickness of Zanda Basin sediments, and arrived at roughly similar magnetic reversal patterns.

Qian's (1999) first attempt simply correlated his magnetic section at the Xiangze area, with 12 normal and 13 reversed magnetozones, to then "Chron 6" through Jaramillo Chron (6.15–1.5 Ma). Qian's selection of the Xiangze area may have been influenced by the presence of the fossil giraffe reported by Zhang et al. (1981), although he did not explicitly use this fossil as a constraint.

Wang et al. (2008b) listed four fossil sites in their more densely sampled magnetic section revealing 15 normal and 15 reversed magnetozones that are correlated to 2An–4Ar (9.5–2.6 Ma). Of their four fossil constraints, three are mammals (Hipparion zandaense, Ochotona sp., and Dicerorhininae) and a fourth included four species of gastropods. However, of the three mammals they mentioned, only one site yielded identifiable mammals (four cheek teeth of H. zandaense) collected by the authors, and the other two are inferred occurrences of Ochotona sp. (a single cheek tooth in Meng et al., 2005) and a rhinoceros (a metapodial in Meng et al., 2004) purportedly collected from the Dingdingka area, which is across the Langqên Zangbo River many km north of Wang et al.'s (2008b) magnetic section. More importantly, the latter two fossils are not well-preserved enough to offer meaningful constraints within their proposed age range of late Miocene through Pliocene. H. zandaense, on the other hand, occurs only in Zanda Basin and its age is thus not externally referenced. In fact, as our own biostratigraphy shows (Figs. 4, 5), H. zandaense has a long range that spans from late Miocene through Pliocene. In general, the three fossil mammals (two of them lacking precise stratigraphic occurrence) cited by Wang et al. (2008b) can only serve as a weak age constraint in the range of late Miocene to Pliocene, too wide to be of much utility in a basin that straddles the entire age range. As discussed below, Wang et al.'s (2008b) correlation misplaced the Pleistocene part of the section and yielded age estimates that are about 2 Ma too old.

Most recently, Saylor et al. (2009) presented two more magnetic sections from the south and southeast Zanda Basin. A composite magnetic column was derived by combining parts of the sections with more magnetozones, and resulted in a total of 12 normal and 12 reversed zones. Saylor et al.'s south Zanda magnetic section was sampled from the same Zharang Canyon as that by Wang et al. (2008b), i.e., Zanda– Bolin section in Kempf et al. (2009), although individual measured segments (legs) differ from each other. It is thus reassuring that these two magnetic sections complement each other in many features. Where Saylor et al. (2009) under-sampled at their 100–150 m level, Wang et al.'s much denser sampling revealed a relatively long reversed zone missed by Saylor et al. (2009) (Fig. 5).

Both Qian (1999) and Saylor et al. (2009, 2010b) arrived at similar magnetic correlations in their upper sections. Saylor et al. started their top normal magnetic zone at C1n, whereas Qian, missing a normal at his top section, begins the long reversed at C1r. This is in contrast to Wang et al. (2008b) who placed his top normal (N1) in 2An. Both Wang et al. (2008b) and Saylor et al. (2009, 2010b), however, arrived at a similar correlation in the lower part of their sections (the onset of sedimentation), despite their differences in the upper sections. For chronologic constraints, Saylor et al. (2009) also used past reports of vertebrate fossils (Hipparion, Palaeotragus, etc.), as well as invoking a regional C3/C4 vegetation transition at ~7 Ma (e.g., Quade et al., 1989; France-Lanord and Derry, 1994; Quade and Cerling, 1995; Quade et al., 1995; Garzione et al., 2000; Ojha et al., 2000; Behrensmeyer et al., 2007; Ojha et al., 2009), to argue their case of a late Miocene through Pleistocene correlation of \sim 9.2 to \leq 1 Ma.

As discussed above, our own mammalian assemblages offer an independent set of age constraints. Importantly, in the lower part of the sequence, we have recovered a small mammal assemblage (IVPP localities ZD0609 and 0904) that falls in the 174–186 m level of Saylor et al.'s (Saylor, 2008; Saylor et al., 2009) south Zanda section within the top part of an alternating greenish sandstone and silt unit and just below the fine-grained lacustrine mudstones with fine laminations. Furthermore, two localities in the top section that produce *Equus* also help to anchor that part of the section to the Pleistocene.

Overall, the fossiliferous middle Zanda sequence yields characteristic Pliocene faunas with few Miocene leftovers, although the upper alluvial conglomerates and lower fluvial sandstones, from which few vertebrate fossils are found thus far, must have ranged into the Pleistocene and late Miocene, respectively. Based on our paleontologic perspective, we re-interpreted previously published paleomagnetic columns from various parts of the basin (Table 2; Fig. 5). The top 180 m (620–800 m) section with Equus is securely placed in the Pleistocene part (C1n–C2r) of the GPTS, as had been similarly correlated by Qian (1999) and Saylor et al. (2009). Such a correlation implies that both Wang et al. (2008b) and Saylor et al. (2009) had missed the very short normal chrons (C1r.1n, Jaramillo, and C1r.2n, Cobb Mountain) in the 667–744 m range, something not unexpected because of the extremely coarse-grained nature of that part of the section. Qian (1999), on the other hand, appears to have sampled at least one of the short chrons (Fig. 5). Our correlations in the next two major magnetochrons (C2An and C2Ar) below the Pleistocene section also agree with that by Saylor et al. (2009), making the upper half (400 m and up) of the Zanda section relatively uncontroversial in chronology, although a brief reversed zone (R3) in Wang et al. (2008b) must be left unexplained.

The lower half of the composite, however, is less satisfactorily resolved. Significantly, the most severe constraint is placed on the normal interval that contains IVPP locality ZD0609, which produced a small mammal fauna that is Pliocene in character, and we have elevated it to

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Fig. 4. Biostratigraphy of Zanda vertebrates, based on data from Tables 1 and 3. Individual fossil localities (solid circles) are interpolated into six nearest measured sections, which are in turn correlated to the main south Zanda section by lithostratigraphic and sequence stratigraphic means (Saylor et al., 2010b). For additional explanations, see caption of Table 3.

C3n.4n. Such a correlation has a cascading effect of moving the next few magnetozones upward, compressing the lower 200–400 m section into a relatively short interval around C3n. Our new age estimates of the Zanda section (Fig. 5) thus span ~400 Ka to 6.4 Ma using the GPTS of ATNTS2004 (Lourens et al., 2004). Our correlation has the undesirable effect of leaving unexplained several short reversals [333–345 m (N8), \sim 225 m (R11), and 110–120 m (N14)] observed by Saylor et al. (2009) and Wang et al. (2008b).

If we allow a more relaxed interpretation of the small mammal fauna from IVPP locality ZD0609, an alternative correlation is possible for the lower section (dashed lines in Fig. 5). This alternative correlation, more consistent with an even depositional rate, would imply that taxa such as Mimomys and Trischizolagus must have occurred ~1 Ma earlier in Zanda Basin than their known records elsewhere. Furthermore, the alternative also carries its own unexplained short magnetozones (N11 and N12 in Wang et al., 2008b). Such an alternative magnetic interpretation of C1n–C4r for Zanda strata (~ 400 Ka to 8.3 Ma) is closest, but not identical, to that proposed by Qian (1999) although the latter lacks adequate documentation and does not permit placement of our fossils in his section. If the alternative correlation is correct, a number of short magnetochrons (C3B) that are not seen in all three known magnetic sections may have been missed in the coarse sediments at the base of the section.

8. Zanda faunal characteristics and zoogeography

The Zanda Pliocene mammal fauna has two major components. First, many of the large and small mammals have clearly a north China affinity. Typical large mammals from north China or eastern Asian arid or semiarid regions include Chasmaporthetes, Pliohyaena ($=$ Pliocrocuta), Meles, Nyctereutes, Xenocyon, Antilospira, and Hipparion. Small mammals that can find their counterparts in north China include Nannocricetus, Prosiphneus, Apodemus, Mimomys, and Trischizolagus; these are especially abundant in the Pliocene of Inner Mongolia, located at much higher latitudes than Zanda. This is perhaps not surprising given that high

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Table 1

List of vertebrate fossil localities, elevation, stratigraphic level as interpolated to nearest measured sections (Saylor et al., 2010a, 2010b), stratigraphic level relative to south Zanda measured section (m from bottom), and calibrated paleomagnetic ages (Ma). See Sections 2, 6, and 7 (Material and methods, Biostratigraphy, and Magnetostratigraphy) for explanations of correlation scheme. Ages for magnetic chrons in the Geomagnetic Polarity Time Scale (GPTS) are based on ATNTS2004 in Lourens et al. (2004).

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Table 1 (continued)

(continued on next page)

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Table 1 (continued)

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Table 1 (continued)

altitudes in Tibet translate to a climate much colder than its equivalent latitudes elsewhere.

The second group consists of native Tibetan components in the fauna. These include Qurliqnoria, Pseudois, and possibly Panthera for large mammals and Aepyosciurus and several species of Ochotona for small mammals. Qurliqnoria, mostly confined to the Tibetan Plateau, first appeared in the late Miocene of Qaidam Basin (Bohlin, 1937; Wang et al., 2007; X. Wang et al., 2011) and may have given rise

Fig. 5. Correlation of three published paleomagnetic sections. Red circle indicate key fossil sites (ZD0609, 0904) for biochronologic constraints. Ages for magnetic chrons in the Geomagnetic Polarity Time Scale (GPTS) are based on ATNTS2004 in Lourens et al. (2004).

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Table 2

Alternative correlations of south Zanda magnetic sections with magnetic chrons in the Geomagnetic Polarity Time Scale (GPTS) based on ATNTS2004 in Lourens et al. (2004).

to the modern Tibetan antelope (Chiru) Pantholops hodgsonii (Gentry, 1968). The Blue Sheep, Pseudois, is another native Tibetan basal caprine possibly related to pantholopines (Fernández and Vrba, 2005).

We chose to not subdivide the Zanda Pliocene fauna for the moment. There is an apparent small mammal faunal turnover from IVPP localities ZD0609/0904 (5.12–5.05 Ma) to ZD1001 (4.42 Ma), recording a change from Trischizolagus mirificus to T. dumitrescuae and various species of Ochotona, but because of the scarcity of small mammal localities, it is not clear how much of this is due to a sampling effect.

Additional collecting in Zanda Basin will surely extend the ranges of many taxa, and our local first appearance and last appearance datum (Table 3) should only be considered as tentative. For example, we consider it likely that Qurliqnoria should extend down through the late Miocene because it has been known in early late Miocene in Qaidam Basin (Wang et al., 2007; X. Wang et al., 2011). Similarly, Palaeotragus likely had a deeper history because it has been reported in Gyirong Basin (Huang et al., 1980; Wang et al., in press-b), whose Hipparion fauna has been dated to 7.0–6.7 Ma (Yue et al., 2004).

In the Pleistocene part of the section (the upper 180 m), an Equus fauna is emerging even though the very coarse-grained sediments reveal no more than a handful of localities producing very fragmentary materials. Of these, we can recognize some Equus limb bones and a deer antler fragment that appears to be more advanced than those seen in the Pliocene part of the section. Nonetheless, the local first appearance of Equus (IVPP locality ZD0915, 625 m, 2.48 Ma) is close to the beginning of the Pleistocene, indicating a rapid spread of the monodactylid horse as soon as it arrived in the Old World (Lindsay et al., 1980), highlighting its usefulness as a biochron. Therefore, despite the scarcity of fossil sites in the upper Zanda section, there is definitively observable faunal turnover at the Plio-Pleistocene boundary around the 620 m level.

The three-toed horse Hipparion zandaense, on the other hand, is the most densely sampled of any mammal species in Zanda (55 localities; Table 3). Between its last appearance datum (IVPP locality ZD1232, 530 m, 3.36 Ma) and the first appearance of Equus, however, there is a long hiatus with poor sampling. This 0.8 myr gap is clearly a result of unfossiliferous coarse-grained sediments, and consequently we are unable to address the issue of how closely Equus locally replaced Hipparion.

If a more relaxed interpretation of the biochronology of Zanda small mammals is taken, allowing taxa such as Mimomys and Trischizolagus to extend their ranges into the late Miocene (dashed lines in Fig. 5), then our "out of Tibet" hypothesis can also be applied to the small mammals. However, such a tantalizing prospect will have to await the full study of the small mammal fauna.

9. Conclusion

The late Miocene to Pleistocene Zanda Basin is uniquely situated at the northern foothills of the Himalayas and preserves an excellent record of fossil mammals and fishes. This is the best record so far known in Tibet, especially in its exceptionally preserved Pliocene mammal assemblages. The 800 m sedimentary sequence is essentially continuous, often fine-grained, and is ideal for archiving both fishes and terrestrial mammals.

Despite the current preliminary stage of exploration, the Zanda fossil assemblage has revealed an ancestral wooly rhino that was the precursor of its late Pleistocene megafaunal descendants. Such a discovery highlights our "out of Tibet" hypothesis of Pliocene high Tibet as a "training ground" for cold adaptations.

A detailed biostratigraphy based on fossil mammals is presented. Individual localities are correlated to an 800 m master lithostratigraphic and magnetic section, offering a unified, basin-wide framework of superposition. Correlations of individual fossil sites have an error range of about \pm 20 m or less, or \pm 0.2 myr in age range. This high-resolution biostratigraphy thus provides, for the first time, a window into the past biological world.

The new Zanda mammal assemblage provides a far stronger biochronologic constraint than has so far been possible. Using its

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Table 3

Local first and last appearances (FAD and LAD) of Zanda vertebrates and individual Zanda (ZD) localities. In cases where there are more than two localities, locality numbers for the first and last appearances are indicated by bold type. See Fig. 4 and Table 1 for complete listing of all localities. For two of the more densely sampled taxa, Hipparion zandaense and Coelodonta thibetana, many of the localities produced postcranial or poorly preserved cranial and dental materials and identifications to species level is often not possible. Since only one three-toed horse and one rhinoceros are so far known, we tentatively include all localities with less identifiable materials within these two species, which are treated in the same way in Fig. 4. Ages for magnetic chrons in the Geomagnetic Polarity Time Scale (GPTS) are based on ATNTS2004 in Lourens et al. (2004).

^a Indicates an estimate due to reworked status of the locality or a previous (1976) collection without precise locality information. Age resolution to less than 0.2 myr apart may not be meaningful due to systemic errors (see Section 2, Material and methods).

characteristic Pliocene mammals, particularly the small mammal assemblage from IVPP locality ZD0609 in the lower part of the section, we are able to reinterpret three previously published magnetic sections, and propose a revised correlation of the magnetic section to C1n to C3An.1r, spanning ~400 Ka to 6.4 Ma.

The lower 0–150 m is the latest Miocene part of the section. Dominated by conglomerates and other coarse-grained sediments, the lower section is sparse in fossil sites. Only five taxa are available so far: Ochotona, Panthera, Qurliqnoria, Palaeotragus, and Hipparion. All are consistent with a late Miocene age.

Spanning the entire Pliocene is the middle 150–620 m section. By far the most fossiliferous, this section includes a rich variety of large and small mammals. Typical for this group are such small mammals as Nannocricetus, Prosiphneus, Mimomys, Apodemus, and Trischizolagus, and such large mammals as Coelodonta thibetana, Hipparion zandaense, Chasmaporthetes, Nyctereutes, Meles, Antilospira, and others.

The upper 620–800 m is Pleistocene in age. This section is the least fossiliferous because of its abundant coarse gravels and conglomerates. Despite the shortage of fossil localities, we did find unambiguous Pleistocene mammals, such as Equus. Our lowest Equus locality, ZD0915, is very close to the 620 m mark, offering another excellent chronologic constraint.

Zoogeographically late Cenozoic Tibetan mammals seem to come from two major sources. A north China or east Asia component includes common elements such as Mimomys, Prosiphneus, Trischizolagus, Chasmaporthetes, Nyctereutes, Meles, and Xenocyon. The age relationships of these taxa outside Tibet thus offer important biochronologic constraints. The second component is the native Tibetan fauna, and this group includes pikas (Ochotona), squirrels (Aepyosciurus), and ancestral Tibetan antelope (Qurliqnoria). This latter assemblage does not provide age constraints by themselves, but offers important insights into indigenously evolved mammals adapted to high Tibet.

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