First report of leopard fossils from a limestone cave in Kenting area, southern Taiwan (#51992)

First submission

Guidance from your Editor

Please submit by 6 Sep 2020 for the benefit of the authors (and your \$200 publishing discount).



Structure and Criteria

Please read the 'Structure and Criteria' page for general guidance.



Author notes

Have you read the author notes on the guidance page?



Raw data check

Review the raw data.



Image check

Check that figures and images have not been inappropriately manipulated.

Privacy reminder: If uploading an annotated PDF, remove identifiable information to remain anonymous.

Files

Download and review all files from the <u>materials page</u>.

9 Figure file(s)

6 Table file(s)

Structure and Criteria



Structure your review

The review form is divided into 5 sections. Please consider these when composing your review:

- 1. BASIC REPORTING
- 2. EXPERIMENTAL DESIGN
- 3. VALIDITY OF THE FINDINGS
- 4. General comments
- 5. Confidential notes to the editor
- Prou can also annotate this PDF and upload it as part of your review

When ready <u>submit online</u>.

Editorial Criteria

Use these criteria points to structure your review. The full detailed editorial criteria is on your guidance page.

BASIC REPORTING

- Clear, unambiguous, professional English language used throughout.
- Intro & background to show context.
 Literature well referenced & relevant.
- Structure conforms to <u>PeerJ standards</u>, discipline norm, or improved for clarity.
- Figures are relevant, high quality, well labelled & described.
- Raw data supplied (see <u>PeerJ policy</u>).

EXPERIMENTAL DESIGN

- Original primary research within Scope of the journal.
- Research question well defined, relevant & meaningful. It is stated how the research fills an identified knowledge gap.
- Rigorous investigation performed to a high technical & ethical standard.
- Methods described with sufficient detail & information to replicate.

VALIDITY OF THE FINDINGS

- Impact and novelty not assessed.
 Negative/inconclusive results accepted.
 Meaningful replication encouraged where rationale & benefit to literature is clearly stated.
- All underlying data have been provided; they are robust, statistically sound, & controlled.
- Speculation is welcome, but should be identified as such.
- Conclusions are well stated, linked to original research question & limited to supporting results.

Standout reviewing tips



The best reviewers use these techniques

Т	p

Support criticisms with evidence from the text or from other sources

Give specific suggestions on how to improve the manuscript

Comment on language and grammar issues

Organize by importance of the issues, and number your points

Please provide constructive criticism, and avoid personal opinions

Comment on strengths (as well as weaknesses) of the manuscript

Example

Smith et al (J of Methodology, 2005, V3, pp 123) have shown that the analysis you use in Lines 241-250 is not the most appropriate for this situation. Please explain why you used this method.

Your introduction needs more detail. I suggest that you improve the description at lines 57-86 to provide more justification for your study (specifically, you should expand upon the knowledge gap being filled).

The English language should be improved to ensure that an international audience can clearly understand your text. Some examples where the language could be improved include lines 23, 77, 121, 128 - the current phrasing makes comprehension difficult.

- 1. Your most important issue
- 2. The next most important item
- 3. ...
- 4. The least important points

I thank you for providing the raw data, however your supplemental files need more descriptive metadata identifiers to be useful to future readers. Although your results are compelling, the data analysis should be improved in the following ways: AA, BB, CC

I commend the authors for their extensive data set, compiled over many years of detailed fieldwork. In addition, the manuscript is clearly written in professional, unambiguous language. If there is a weakness, it is in the statistical analysis (as I have noted above) which should be improved upon before Acceptance.



First report of leopard fossils from a limestone cave in Kenting area, southern Taiwan

Tzu-Chin Chi Equal first author, 1, 2, Yi Gan Equal first author, 3, Tzu-Ruei Yang Corresp., 1, 3, Chun-Hsiang Chang Corresp., 1

Corresponding Authors: Tzu-Ruei Yang, Chun-Hsiang Chang Email address: lereage@gmail.com, cch@nmns.edu.tw

Longshia-dong Cave, a limestone cave located in the Kenting area within the Kenting National Park of southern Taiwan, yields numerous terrestrial mammalian fossils. Many of them were not reported in historical literature and are neither present in Taiwan. For instance, no historical literature mentioned leopards inhabited in Taiwan, and thus their existence remained unknown. This study describes three fossil leopard (*Panthera pardus*) teeth uncovered from the Longshia-dong Cave. Two isolated lower premolars and one molar, respectively p3, p4 and m1, show a series of progressive increase in size and should have been belonging to the same individual under the subfamily of Pantherinae. Traditional linear measurements and two-dimensional geomorphometric analysis for the occlusal surface outlines were conducted on the fossil teeth and extant pantherines inhabited in Asia such as clouded leopards, leopards, and tigers. Results show that the fossil teeth are similar both in size and morphology to the teeth of extant leopards, suggesting the assignment of the fossil teeth to leopards. This study, for the first time, reported the presence of leopards in the Late Pleistocene of Taiwan. Besides, the smaller size of the fossil teeth than Chinese fossil leopards putatively suggests insular dwarfism, yet more studies are required.

 $^{^{1}}$ Division of Geology, National Museum of Natural Science, Taichung, Taiwan

² Department of Earth and Environmental Sciences, National Chung Cheng University, Chiayi, Taiwan

³ Department of Earth Sciences, National Cheng Kung University, Tainan, Taiwan



1 First report of leopard fossils from a limestone cave in

2 Kenting area, southern Taiwan

```
3
 4
 5
 6
     Tzu-Chin Chi<sup>1,2</sup>, Yi Gan<sup>3</sup>, Tzu-Ruei Yang<sup>1,3</sup>, Chun-Hsiang Chang<sup>1</sup>
 7
 8
 9
     1. Department of Geology, National Museum of Natural Science, 40453 Taichung, Taiwan
10
     2. Department of Earth and Environmental Sciences, National Chung Cheng University, 62102
11
         Chiayi County, Taiwan
     3. Department of Earth Sciences, National Cheng Kung University, 70101 Tainan, Taiwan
12
13
14
     Corresponding Authors:
15
     Tzu-Ruei Yang
16
     No. 1, Guanqian Rd., North Dist., Taichung City 404, Taiwan
17
     Email address: tzurueiyang@nmns.edu.tw
18
     Chun-Hsiang Chang
19
     No. 1, Guanqian Rd., North Dist., Taichung City 40453, Taiwan
20
     Email address: cch@nmns.edu.tw; chang28@gmail.com
21
22
       Keywords: Felidae, Morphology, Panthera pardus, Late Pleistocene, Geomorphometric
23
     analysis
```



Abstract

Longshia-dong Cave, a limestone cave located in the Kenting area within the Kenting National Park of southern Taiwan, yields numerous terrestrial mammalian fossils. Many of them were not reported in historical literature and are neither present in Taiwan. For instance, no historical literature mentioned leopards inhabited in Taiwan, and thus their existence remained unknown. This study describes three fossil leopard (*Panthera pardus*) teeth uncovered from the Longshiadong Cave. Two isolated lower premolars and one molar, respectively p3, p4 and m1, show a series of progressive increase in size and should have been belonging to the same individual under the subfamily of Pantherinae. Traditional linear measurements and two-dimensional geomorphometric analysis for the occlusal surface outlines were conducted on the fossil teeth and extant pantherines inhabited in Asia such as clouded leopards, leopards, and tigers. Results show that the fossil teeth are similar both in size and morphology to the teeth of extant leopards, suggesting the assignment of the fossil teeth to leopards. This study, for the first time, reported the presence of leopards in the Late Pleistocene of Taiwan. Besides, the smaller size of the fossil teeth than Chinese fossil leopards putatively suggests insular dwarfism, yet more studies are required.

Introduction

The Longshia-dong (literally Lobster Cave in Chinese) Cave is located in the Kenting
Forest Recreation Area (KFRA) of the Kenting National Park, southernmost Taiwan (Fig. 1).
The KFRA is covered with thick limestone (Hengchun Limestone) that deposited during the
Middle Pleistocene (Gong & Yui, 1998). A number of caves and fissures formed in the



47 Hengchun Limestone, some of which, such as the Longshia-dong Cave, accumulated numerous 48 terrestrial mammal fossils. This cave opens at ca. 240 m above the present sea level and is a 49 small tunnel inclined gently toward its inner part (Kawamura et al., 2016). The investigation of the Longshia-dong Cave was initiated by Prof. Ai Kawamura from the 50 Aichi University of Education, Japan and Dr. Chun-Hsiang Chang from the National Museum of 51 52 Nature Science, Taiwan (Kawamura et al., 2016). To date, fossils of Cervidae, Rodentia (e.g., 53 Microtus and Hystrix), Carnivora (three teeth), Macaca sp., and Rhinolophus sp. uncovered from this cave are identified (Kawamura et al., 2016). However, the Microtus (Rodentia) is now 54 55 restricted to high mountains in Taiwan and the *Hystrix* sp. (Rodentia) is no longer present in 56 Taiwan. Besides, leopard teeth were not reported in Taiwan previously. These lines of evidence 57 indicate a very different faunal setting in comparison with the present one. 58 In previous excavations scientists have discovered putative Middle-Late Pleistocene big cat 59 fossils, such as *Panthera* cf. tigris and *Panthera* sp. from the Chochen area in southern Taiwan (Fig. 1; (Otsuka, 1984; Chen, 2000a; Chen, 2000b; Wei, 2007). Besides, fossil remains of 60 Panthera tigris have also been collected from the Penghu Channel, a N-S striking submarine 61 62 valley off the west coast of Taiwan (Fig. 1) (Ho et al., 1997; Asahara et al., 2015). While both of 63 these studies indicate a rich fossil record of felids in the Middle-Late Pleistocene of Taiwan, the modern Felidae in Taiwan are only featured by two species, including clouded leopard (Neofelis 64 nebulosa) and leopard cat (Prionailurus bengalensis). Even more recently, Chiang et al. (2015) 65 66 presumed that the former is extinct in Taiwan, thus leaving only the latter present to our 67 knowledge. 68 Previous studies indicated the similarities of Taiwan's fauna to the Early Pleistocene fauna of southern China (e.g., Otsuka and Lin, 1984a, 1984; Otsuka and Lin, 1984b; Lai, 1989; Qi et 69 al., 1999; Chen, 2000a; Fooden and Wu, 2001). The fauna of southern China probably entered 70



71 Taiwan in the Late Pliocene to Early Pleistocene, when Taiwan was connected with China (Lai, 72 1989); Chen (2000a); Fooden and Wu, 2001); however, only clouded leopard was reported in 73 Taiwan's historical record. While Swinhoe (1862) had never seen a living individual, he 74 described it as a small, short-tailed, small-footed animal based on the fur specimen and named it as Leopardus brachyurus. Later, Swinhoe (1870) replaced L. brachyurus with Felis macrocelis, 75 76 but *L. brachyurus* is still the most commonly used name. 77 In the excavation to the Longshia-dong Cave in 2014, three teeth were collected from the same horizon. A preliminary study has indicated their affinity to feline remains, though further 78 79 investigations are needed (Gan, 2016). This study thus aims to reveal their taxonomic affinity 80 based on traditional linear and geometric measurements. Besides, the comparison between the 81 studied material and leopard fossils from various sites of the Pleistocene in China, including the 82 Zhoukoudian site (Teilhard & Pei, 1941) and Mentougou Bull Eye Cave (Deng et al., 1999) of 83 Beijing, Lantian (Gongwangling) of Shanxi (Hu, 1978), Anyang (Yinxu) of Henan Province (de Chardin & Young, 1936), Liucheng Cave of Guangxi Province(Pei, 1987) (Fig. 1), also allows 84 us to reveal more details. 85 Geological setting 86 87 The Kenting National Park is located in the Hengchun Peninsula, southernmost Taiwan (Fig. 1). The Hengchun Peninsula represents the earliest stage of the Taiwanese orogeny (Huang 88 et al., 1985), and therefore many incipient thrust faults are observed. A major boundary fault, the 89 90 N-S striking Hengchun Fault (Fig. 1) (Chen et al., 2005), divides the Hengchun Peninsula into 91 two terranes, including the Central Range in the East and the Western Foothills (Hengchun 92 Valley and West Hengchun Hill) in the West (Yen & Wu, 1986). Since the Late Pleistocene, the 93 Hengchun Peninsula was uplifted at a rate of 2-6 mm/yr and thus gave rise to the development of





94	coral reef and limestone (Hengchun Limestone). Many karst caves were afterward formed and
95	harbor various fossils. A thin layer of reddish sand and gravels (Xu, 1989; Hseu et al., 2004)
96	overlying the Hengchun Limestone was named as Eluanpi Bed in the Southeast, or Taiping
97	Formation in the West, to the Longshia-dong Cave (Fig. 1).
98	Our studied area, the Longshia-dong Cave, is located in the southeastern part of the Kenting
99	National Park, a national park that is featured by the karst landscape mostly contributed by the
100	Hengchun Limestone (Fig. 1). The cave opens to the East, measuring 30~40 m long, 8 m wide,
101	and about 5 m deep, and a puddle was found in the end of the cave (Kawamura et al., 2016).
102	Most of the limestone in the cave is covered by reddish sediment that is contributed by limestone
103	pebbles and fossils, though the boundary between the reddish sediment and Hengchun Limestone
104	is unclear. A flow stone made of carbonate calcite was found 7 m away from the cave entry, and
105	the fossil-bearing sediments are found behind it. The fossil-bearing sediments are characterized
105106	the fossil-bearing sediments are found behind it. The fossil-bearing sediments are characterized by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great
106	by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great
106 107	by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great similarity to the Eluanbi Bed. Besides, a layer of blackish mud and light yellow silt overlying the
106107108	by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great similarity to the Eluanbi Bed. Besides, a layer of blackish mud and light yellow silt overlying the fossil-bearing sediments, based on a previous study, is composed of recent alluvial deposits since the Holocene (Wang, 2015).
106107108109	by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great similarity to the Eluanbi Bed. Besides, a layer of blackish mud and light yellow silt overlying the fossil-bearing sediments, based on a previous study, is composed of recent alluvial deposits since the Holocene (Wang, 2015).
106 107 108 109	by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great similarity to the Eluanbi Bed. Besides, a layer of blackish mud and light yellow silt overlying the fossil-bearing sediments, based on a previous study, is composed of recent alluvial deposits since the Holocene (Wang, 2015). While the age of the fossil-bearing sediments is unlikely estimated, previous studies have put
106107108109110111	by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great similarity to the Eluanbi Bed. Besides, a layer of blackish mud and light yellow silt overlying the fossil-bearing sediments, based on a previous study, is composed of recent alluvial deposits since the Holocene (Wang, 2015). While the age of the fossil-bearing sediments is unlikely estimated, previous studies have put emphasis on the age of the Hengchun Limestone, which gave a maximum estimation for the
106 107 108 109 110 111 112	by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great similarity to the Eluanbi Bed. Besides, a layer of blackish mud and light yellow silt overlying the fossil-bearing sediments, based on a previous study, is composed of recent alluvial deposits since the Holocene (Wang, 2015). While the age of the fossil-bearing sediments is unlikely estimated, previous studies have put emphasis on the age of the Hengchun Limestone, which gave a maximum estimation for the leopard fossils uncovered from the cave. The Hengchun Limestone was either considered two-
106 107 108 109 110 111 112 113	by a mixture of reddish sand and mud, as well as limestone pebbles, which shows a great similarity to the Eluanbi Bed. Besides, a layer of blackish mud and light yellow silt overlying the fossil-bearing sediments, based on a previous study, is composed of recent alluvial deposits since the Holocene (Wang, 2015). While the age of the fossil-bearing sediments is unlikely estimated, previous studies have put emphasis on the age of the Hengchun Limestone, which gave a maximum estimation for the leopard fossils uncovered from the cave. The Hengchun Limestone was either considered two-stage (Gong, 1982) or three-stage (Shih et al., 1989) reef formation, but both studies have



suggested a date of 325-125 ka (Gong, 1994), this study will discuss further details of the leopard remains based on the 500 ka.

118

119

117

116

Materials & Methods

120	Three well-preserved whitish fossil teeth were collected from the Longshia-dong cave in the
121	2014 excursion led by one of the authors (CH. Chang). All of them (F056584, F056585, and
122	F056586) are housed at National Museum of Natural Science, Taichung, Taiwan (NMNS) under
123	the accession numbers provided. The three teeth were discovered in a very small grid (50 cm*50
124	cm) and present a series of progressive increase in size; thus, we believed they should have been
125	belonging to a felid individual (Fig. 2). The felid fossil teeth, based on their shape (Hillson,
126	2005), are assignable to p3 (F056584), p4 (F056585) and m1 (F056586) from the right lower jaw
127	(Fig. 2). In addition to the three felid teeth, the skulls and mandibles of 30 extant specimens,
128	including nine clouded leopards (Neofelis nebulosi or Neofelis diardi), six leopards (Panthera)
129	pardus), and 15 tigers (Panthera tigris), which are housed at the NMNS, Endemic Species
130	Research Institute, and Taipei Zoo, respectively, were also included in this study (see the details
131	in the Supplement I). Prior to our qualitative studies of the fossil and extant felid teeth
132	(Kawamura, 1992; Fukawa, 2000), we compare their morphological features to determine the
133	assignment of the fossil felid teeth (Fig. 3). Photos of all specimens were taken with a Panasonic
134	Lumix DMC-GF1 camera and a Panasonic Lumix GF1 14-45mm/F3.5-5.6 lens. These photos
135	were afterwards imported into the TPS software for the traditional linear measurements and
136	geomorphometric studies (Rohlf, 2005).
137	Traditional linear measurements were taken point-to-point; a total of 22 dental dimensions
138	including antero-posterior crown length (1, 5, 15 in Fig. 3), dorsoventral crown height (3, 7, 10,



139 13, 17, 20 in Fig. 4), width of each cusp (4, 8, 11, 14, 18, 21 in Fig. 4), anteroposterior length of 140 cusps (2, 6, 9, 12, 16, 19 in Fig. 4), and carnassial length (22 in Fig. 4) (Christiansen, 2008), were obtained from all specimens (a fossil individual and 30 extant specimens, see the details in 141 the Supplement I). The fossil felid individual only preserves the lower right jaw and therefore 142 143 only 22 data were obtained. The 30 extant specimens, on the other hand, permit the measurement 144 of both lower right and left jaws, thus contributing to a 30*22*2 data matrix (note that some data 145 are not available due to the poor preservation, see the details in the Supplement I). 146 The data from the aforementioned traditional linear measurements were introduced into two 147 rounds of principal component analysis (PCA) (Morrison, 1976; Dunteman, 1994), which plots 148 the data to a new coordinate system contributed by two principal components (Jolliffe, 2002; Hsu, 2003). All PCA in this study were performed with the R package "stats" (R Core Team, 149 150 2013). In the first round of PCA, three teeth (p3, p4, and m1) are seen as a dataset, and thus any 151 missing of them leads to the removal of the specimen from the first PCA. Besides, two datasets are available from an extant specimen since the lower right and lower left jaws are both 152 153 measured. Ultimately, one dataset from the felid fossil, 14 datasets from the nine clouded 154 leopards (supposedly 18), five datasets from the six leopards (supposedly 12), and ten datasets 155 from the 15 tigers (supposedly 30), were used in the first round of PCA (Fig. 5a). 156 To avoid the effect of the interspecific difference in tooth size on the first round of PCA, we 157 performed the second round of PCA with the R package "stats" for p3, p4, and m1, separately. 158 For instance, we obtained three ratios, including of protoconid length (2 in Fig. 4 and Tab. 1) to 159 crown length (1 in Fig. 4 and Tab. 1), of protoconid height (3 in Fig. 4 and Tab. 1) to crown 160 length (1 in Fig. 4 and Tab. 1), and protoconid length (2 in Fig. 4 and Tab. 1) to protoconid height (3 in Fig. 4 and Tab. 1) based on the data from all available p3 (a fossil felid, 13 clouded 161 leopards, five leopards, and nine tigers). The three ratios were input into R program for PCA 162



analysis with the package "stats." For PCA analysis (Fig. 5b), on the other hand, six ratios and
seven ratios were obtained based on the data from all available p4 and m1, respectively (see the
details in Fig. 5) and were introduced into R program for PCA analysis with the package "stats"
(Figs. 5c and 5d).
In addition to the PCA analyses based on traditional linear measurements, we also performed
geomorphometric analysis (Slice, 1996) because of its utility of revealing the morphological
similarities between different groups (Zelditch et al., 1995) and the ability to excluded the factor
of allometric growth. All photographs were input into the program tpsUtil for building up a tps
file. To access the morphology of the occlusal surface of each tooth in the absence of apparent
landmarks, we used the "curve mode" in the program tpsDig 2.05 (accessed on Dec 1, 2014 from
http://life.bio.sunysb.edu/morph/; (Rohlf, 2005) to place evenly distributed 150 semi-landmarks
around the occlusal surface on each photo (Gunz & Mitteroecker, 2013). The 150 semi-
landmarks were then digitized from photographs using tpsDig 2.05, which converted points
marked on the photographs into Cartesian x, y coordinates. After scaling and alignment of the
digitized semi-landmarks, a relative warp analysis (RWA) was then performed on the set of
specimen semi-landmarks in tpsRelw (Rohlf, 2007) to unravel the morphological variation
between the fossil and extant teeth. We then visualize the morphological variation from RWA by
plotting the relative warp axes as a PCA. Relative differences are presented in the form of thin
plate spline deformation grids (Fig. 6; (Zelditch et al., 2004; Tseng et al., 2010).

Results

- Description and morphological comparison
- Family Felidae Fischer de Waldheim, (1817).



186	Subfamily Pantherinae Pocock, 1917.
187	Genus Panthera Oken, 1816.
188	Panthera pardus (1758) (Fig. 2)
189	Based on our observation of all extant felid specimens and previous studies (Gray, 1867;
190	Christiansen & Kitchener, 2011; King, 2012), we concluded the following common dental
191	characters (Fig. 3): (1) two-rooted p3, p4 and m1; (2) p3 is smaller than p4 and only has
<mark>192</mark>	hypoconid; (3) p4 is in a similar size to m1; (4) p3, p4 has three cusps (paraconid, protoconid)
193	and hypoconid) and its paraconid and hypoconid are well-developed (5) m1 has two well-
194	developed cusps (paraconid and protoconid) and undeveloped talonid. All of the aforementioned
195	features are present in the fossil teeth and thus indicate the assignment of the fossil teeth to
196	Felidae. Moreover, the fossil teeth can be further assigned to Pantherinae based on their similar
197	size to extant pantherines. Pumas (Puma concolor), which belong to Felinae, a sister group to
198	Pantherinae, are the only group of felines that present similar dental size; however, their current
199	distribution (only in Americas) makes the attribution of the fossil teeth to puma unlikely.
1 <mark>99</mark> 200	distribution (only in Americas) makes the attribution of the fossil teeth to puma unlikely. In previous studies (Meachen-Samuels & Van Valkenburgh, 2009), dental size is the only
200	In previous studies (Meachen-Samuels & Van Valkenburgh, 2009), dental size is the only
200 201	In previous studies (Meachen-Samuels & Van Valkenburgh, 2009), dental size is the only character that was used for distinguishing different pantherines, such as lions (<i>Panthera leo</i>),
200 201 202	In previous studies (Meachen-Samuels & Van Valkenburgh, 2009), dental size is the only character that was used for distinguishing different pantherines, such as lions (<i>Panthera leo</i>), snow leopards (<i>Panthera uncia</i>), tigers, clouded leopards, and leopards. For instance, the dental
200 201 202 203	In previous studies (Meachen-Samuels & Van Valkenburgh, 2009), dental size is the only character that was used for distinguishing different pantherines, such as lions (<i>Panthera leo</i>), snow leopards (<i>Panthera uncia</i>), tigers, clouded leopards, and leopards. For instance, the dental size of lions are significantly larger than the one of the fossil teeth. Moreover, the attribution of
200 201 202 203 204	In previous studies (Meachen-Samuels & Van Valkenburgh, 2009), dental size is the only character that was used for distinguishing different pantherines, such as lions (<i>Panthera leo</i>), snow leopards (<i>Panthera uncia</i>), tigers, clouded leopards, and leopards. For instance, the dental size of lions are significantly larger than the one of the fossil teeth. Moreover, the attribution of the fossil teeth to snow leopards is unlikely since snow leopards are only present in high
200 201 202 203 204 205	In previous studies (Meachen-Samuels & Van Valkenburgh, 2009), dental size is the only character that was used for distinguishing different pantherines, such as lions (<i>Panthera leo</i>), snow leopards (<i>Panthera uncia</i>), tigers, clouded leopards, and leopards. For instance, the dental size of lions are significantly larger than the one of the fossil teeth. Moreover, the attribution of the fossil teeth to snow leopards is unlikely since snow leopards are only present in high mountains. However, dental size is not a indicator for the distinguishment between tigers,
200 201 202 203 204 205 206	In previous studies (Meachen-Samuels & Van Valkenburgh, 2009), dental size is the only character that was used for distinguishing different pantherines, such as lions (<i>Panthera leo</i>), snow leopards (<i>Panthera uncia</i>), tigers, clouded leopards, and leopards. For instance, the dental size of lions are significantly larger than the one of the fossil teeth. Moreover, the attribution of the fossil teeth to snow leopards is unlikely since snow leopards are only present in high mountains. However, dental size is not a indicator for the distinguishment between tigers, clouded leopards, and leopards. This study, however, shows that several pantherines, including



Clouded leopards have two distinct features, including an undeveloped paraconid of p3 and a well-developed metaconid of m1, while both of which are absent in the fossil teeth from Longshia-dong Cave (Fig. 2). Thus, the assignment of the fossil teeth from Longshia-dong Cave to clouded leopards is here excluded. On the other hand, tiger teeth are characterized by a highly developed p4 paraconid, which is not seen in the fossil teeth (Figs. 2, 6). Moreover, the p3 of tigers have a lower protoconid than the one of the fossil (Figs. 2, 6). Tigers also present a wider crown in all teeth, especially in m1, than the ones of the fossil (Table 2).

The fossil pantherin teeth, in addition to their size, show many distinct features that are similar to those of the extant leopards, such as the presence of p3 paraconid and the shape of occlusal surface. Besides, the presence of m1 and slightly worn enamel indicate that the fossil teeth were belonging to a very young adult (Stander, 1997), though its gender is uncertain because of the lacking of morphological differences between male and female leopard teeth (Pocock, 1930) and the extremely poor sample size (n=1).

Traditional linear measurement

While our aforementioned morphological comparison indicates the affinity of the fossil teeth to leopards (*Panthera pardus*), traditional linear measurement was performed for further lines of evidence. The result of the first round of PCA based on the 22 dental dimensions shows a significant disparity between tigers and the others (clouded leopards and leopards) (Fig. 5a) (factor loading of each component shows on Supplemental Table 3), although the disparity between clouded leopards and leopards is inapparent. In the second round of PCA based on various ratios from the dimensions of p3, p4, and m1 (see Material and methods), while no pattern is indicated by the result of the PCA on p3 (Fig. 5b), the assignment of the fossil teeth to



235

236

237

238

239

240

241

242

243

244

245

246

247

248

249

250

251

tigers or to clouded leopards is respectively excluded based on the PCAs on p4 (Fig. 5c) and onm1 (Fig. 5d).

Geomorphometric analysis

In addition to the traditional linear measurement (Fig. 5), geomorphometric analysis were also performed for further lines of evidence. In the RWA based on all third premolars (p3), the first three relative warp axes accounted for 74.81% of the total variation, though no morphological disparities were revealed (Fig. 6). The RWA based on all fourth premolars (p4), however, shows the disparity between clouded leopards and the others in the plots of RW1 to RW2 (Fig. 7a) and RW2 to RW3 (Fig. 7c). The RW2 explains 24.52% of the total shape variance and relates primarily to the prominence of the mesial side toward buccal or lingual side (Fig. 7f). On the other hand, the RWA based on all first molars indicates the isolation of tigers in the plots of RW1 to RW2 (Fig. 8a) and RW2 to RW3 (Fig. 8c). Striking samely, the RW2 in the RWA based on all first molars, which accounts for 22.67% of the total variation, is also the best indicator that excludes tigers. In summary, the fossil pantherin teeth uncovered from Kenting, based on various lines of evidence, are assignable to leopards. While the plot of p3, p4, and m1 sizes of the fossil and extant leopards indicates that only the fossil m1 is encompassed within the variations of the extant leopard teeth, the slope of the regression line based on the three fossil teeth is similar to the ones of those based on the extant specimens. Thus, the three fossil teeth were very likely from a common origin.



Discussion

An integrative, qualitative method for the identification of leopard teeth

Morphological comparisons have been utilized in many studies for the species identification of various mammalian teeth, such as elephants (Todd, 2010), Hedgehogs (Gould, 2001) and moles (Van Cleef-Roders & Van Den Hoek Ostende, 2001). Based on the morphology and size of the fossil teeth (F056584, F056585, and F056586), we are only able to assign them to the subfamily of Pantherinae. Such a qualitative method, however, is not sufficient for the species identification of the subfamily of Pantherinae (lions excluded as their teeth are apparently larger than leopard teeth) because pantherin teeth are very similar both in size and morphology.

On the other hand, canine teeth are much more complex than felid teeth and thus have more homologous features for landmarks. Therefore, Asahara et al. (2015) have applied morphological comparisons in combination with geomorphometric analyses to canine teeth, which is an integrative method that has never been performed on felid teeth. This study, for the first time, demonstrates that an integration of morphological comparisons, linear measurements, and geomorphometric morphometric studies allows species identification of the subfamily of Pantherinae based on teeth.

Taphonomic implications for an adult origin of the fossil teeth

A previous study by Stander (1997) suggested that tooth wear in leopards is a reliable indicator for their ages. The eruption of permanent teeth in one to two years old leopards is completed and thus are whitish and wearless. The crown tips are wore in individuals older than three years old, and the tooth wear appears first on incisors, then on canines, on premolars, and finally on molars. The wear is apparent on the teeth of the individuals of five or six years old. Although the fossil teeth in this study only preserve p3, p4, and m1, their complete eruption

indicates an age older than one year. Moreover, the slight wear on their crown tips offers further 275 276 information that the fossil leopard should have been younger than five years old at death. 277 A behavioral or sedimentary origin? 278 Extant leopards tend to carry their kills to a safe, isolated location for storage (de Ruiter & Berger, 2000). Many studies, based on field observation, indicated that the leopards in South 279 Africa prefer carrying their kills into their caves, thus contributing to the large number of skeletal 280 remains in many caves in South Africa (Le Roux & Skinner, 1989). In addition to the fossil 281 282 leopard teeth, many mammalian fossils, such as deers (*Cervus*), macaques (*Macaca* sp.), and hystrixes (*Hystrix* sp.), were also uncovered from Longshia-dong Cave. These mammalian 283 fossils were once considered the kills brought back by leopards. Nevertheless, a previous study 284 285 pointed out that the deposit in Longshia-dong Cave is a result of multiple reworkings (Wang, 286 2015). The co-occurrence of the mammalian fossils in Longshia-dong Cave probably represents 287 a fossil accumulation over thousands of years. 288 Moreover, leopards left various bite marks on prey's bones (Shi & Wu, 2011; Binford, 2014), but Lin (2017) examined fossil bones from Longshia-dong Cave and failed to find any 289 290 common bite marks. Most deer fossils from the cave are mandibles or limb bones, all of which are unfavorable to leopards (Li, 2007). Thus, we suggest that the accumulation of mammalian 291 292 fossils in Longshia-dong Cave is very likely a result of multiple reworkings. 293 A comparison of the fossil teeth from Longshia-dong Cave to China leopard fossils Leopards are widely distributed throughout Asian and African continents, but only a few of 294 295 them are currently present on islands such as Java and Sri-Lanka (Pocock, 1932). No written 296 literature in Taiwan has reported the presence of leopards; however, this study reported the first



298 Pleistocene of Taiwan. 299 Many previous studies suggested a continental origin of Taiwanese mammalian fossils based on 300 various lines of evidence (Otsuka & Lin, 1984; Lai, 1989; Qi et al., 1999; Chen, 2000a; Fooden 301 & Wu, 2001). The Late Pleistocene of China has produced numerous leopard fossils from 302 various sites, including Beijing Mentougou Niuyan Cave (Deng et al., 1999), the first and 303 thirteenth locations of Beijing Zhoukoudian Site (Pei, 1934), Shaanxi Lantian Gongwangling Site (Hu, 1978), Anyang Yinxu Site (de Chardin & Young, 1936), and Guangxi Liucheng Cave 304 (Pei, 1987) (Fig. 1a). Among these reports, Deng et al. (1999) claimed that the two lower first 305 306 molars of the three fossil leopard teeth uncovered from Mentougou Niuyan Cave are the smallest (19.3×9.2 mm and 19.5×8.5 mm) in comparison with the other Chinese fossil leopard teeth. The 307 size of the fossil teeth in our study (16.77×7.95 mm; Table 3), however, are much smaller than 308 309 the teeth reported in Deng et al. (1999). The smaller dental size can be explained by two hypotheses: (1) ontogenetic variation and (2) insular dwarfism. The first hypothesis is here 310 311 precluded as both of our specimen and the Mentougou Niuyan Cave specimen are permanent 312 teeth. For the second hypothesis, Meiri et al. (2005) indicated that the m1 size in carnivores 313 randomly varies in different habitats (Meiri et al., 2005), albeit not regularly and predictably with 314 either area or isolation. They concluded three selective forces, including resource limitation, 315 predation and interspecific competition. Our study area, Kenting National Park, has produced 316 less various fossils than all aforementioned Chinese sites, suggesting a habitat with fewer resources for carnivores. Besides, the leopard described in this study is the only carnivores found 317 in Kenting National Park to date, indicating lower predation pressure by other carnivores and the 318 absence of interspecific competition. While the absence of interspecific competition would have 319

leopard fossils from Longshia-dong Cave and thus suggested the presence of leopards in the Late





resulted in gigantism, both of the fewer resources and lower predation pressure possibly contributed to the smaller size of Taiwanese leopards than Chinese leopards. Yet, more specimens are required for further studies.

Conclusions

Leopards are a group of carnivores widely distributed throughout Asian and African continents, yet they are no longer found in eastern Asia due to civilization. Fossils thus represent the only clue indicating prehistoric leopards. While their teeth are significantly smaller than the other pantherines and thus are easily identified, poorly preserved fossils hinders further investigations. This study, based on an integration of morphological and geomorphometric analyses, reveals the assignment of the fossil teeth excavated from Longshia-dong Cave to *Panthera pardus*, which is currently absent on Taiwan Island and suggests the presence of leopards in the Late Pleistocene of Taiwan. Such a record of prehistoric leopards in Taiwan thus adds up to the carnivore biodiversity of Taiwan. However, whether the panthera individual is aboriginal or migrated is still uncertain. Our study also indicates the smaller size of Taiwanese leopard fossil teeth than Chinese ones. Such a smaller dental size was possibly a result of insular dwarfism. To conclude, the discovery of the leopard fossil in Taiwan opens shed some light on the origin of the Kenting Fauna. Nevertheless, the insular dwarfism of, and the migration history of, the prehistoric leopards in Taiwan require more specimens and studies.



341

342

343

344

345

346

347

348

Acknowledgements

The authors would like to thank Y.- J. Chen of NMNS, S.-W. Chang of the Endemic Species Research Institute, and T.-C. Chan of Taipei Zoo for their assistance in accessing the specimens. We are also grateful to P.-J. Chiang for his helpful comments and J. Chen for her help on illustrations. Appreciation also goes to all members of the Taiwan-Japan joint excursion in 2014. We also thank K.-C. Wang and T.-Y. Hsiao for their assistance in the excursion. We also thank J.-S. Chi for proofreading. This study was supported by the grants of the Ministry of Science and Technology (MOST), Taiwan (102-2116-M-178-004- to C.-H. Chang and 108-2116-M-178-003-MY2 to T.-R. Yang).

349

350

References

Asahara M, Chang C-H, Kimura J, Son NT, and Takai M. 2015. Re-examination of the 351 352 fossil raccoon dog (Nyctereutes procyonoides) from the Penghu channel, Taiwan, and an age estimation of the Penghu fauna. Anthropological Science 353 354 123:177-184. 10.1537/ase.150710 355 Binford LR. 2014. Bones: ancient men and modern myths. Cambridge, Massachusetts, 356 United States of America: Academic press. 357 Chen K-T. 2000a. On Taiwan mammalian faunas in different periods of time and related 358 problems: The background materials for Taiwan zooarchaeological studies, part 1. Bulletin of the Institute of History and Philology Academia Sinica 71:129-198. 359 Chen K-T. 2000b. On Taiwan mammalian faunas in different periods of time and related 360 361 problems: The background materials for Taiwan zooarchaeological studies, part 362 2. Bulletin of the Institute of History and Philology Academia Sinica 71:367-457.



363	Chen W-S, Lee W-C, Huang N-W, Yen I-C, Yang C-C, Yang H-C, Chen Y-C, and Sung
364	S-H. 2005. Characteristics of Accretionary Prism of Hengchun Peninsula,
365	Southern Taiwan: Holocene Activity of the Hengchun Fault Western Pacific Earth
366	Sciences.
367	Chi W-R. 1982. The calcareous nannofossils of the Lichi Melange and the Kenting
368	Melange and their significance in the interpretation of plate-tectonics of the
369	Taiwan region. Ti-Chih (Geology) 4:99-114.
370	Chiang P-J, Pei KJ-C, Vaughan MR, Li C-F, Chen M-T, Liu J-N, Lin C-Y, Lin L-K, and
371	Lai Y-C. 2015. Is the clouded leopard Neofelis nebulosa extinct in Taiwan, and
372	could it be reintroduced? An assessment of prey and habitat. Oryx 49:261-269.
373	Christiansen P. 2008. Species distinction and evolutionary differences in the clouded
374	leopard (Neofelis nebulosa) and Diard's clouded leopard (Neofelis diardi).
375	Journal of Mammalogy 89:1435-1446.
376	Christiansen P, and Kitchener AC. 2011. A neotype of the clouded leopard (Neofelis
377	nebulosa Griffith 1821). Mammalian Biology 76:325-331.
378	de Chardin PT, and Young C. 1936. On the Mammalian Remains from the
379	Archaeological Site of Anyang: Geological Survey of China.
380	de Ruiter DJ, and Berger LR. 2000. Leopards as taphonomic agents in dolomitic
381	caves—implications for bone accumulations in the hominid-bearing deposits of
382	South Africa. Journal of Archaeological Science 27:665-684.
383	Deng T, Huang W-B, and Hong-Jie W. 1999. The Late Pleistocene mammlian fossils
384	from Nyuyan Cave in Mentougou, Beijing, China. Vertebrata
385	Palasiatica:156~164.
386	Dunteman GH. 1994. Principal Component Analysis (PCA): New Delhi: Sage Toppan.



387	Fisher von Waldheim G. 1817. Adversaria zoological. <i>Mémoires de la Société Imperiale</i>
388	des Naturalistes de Moscou 5:368-428.
389	Fooden J, and Wu H-Y. 2001. Systematic review of the Taiwanese Macaque, <i>Macaca</i>
390	cyclopis Swinhoe, 1863. Zoology.
391	Fukawa M. 2000. Research methods for fossils Earth Sciencne education 53:282.
392	Gan Y. 2016. Study the Late Pleistocene Cat (Carnivora, Felidae) Fossils from a
393	Limestone Cave within Kenting Area, Southern Taiwan Master. National Cheng
394	Kung University
395	Gong S-Y. 1982. Study Stratigraphic and Paleoenvironmental of Hengchun Limestone.
396	PhD Thesis. Department of Geology, National Taiwan University
397	Gong S-Y, and Yui T-F. 1998. Meteoric diagenesis and stable isotopic compositions of
398	the Hengchun Limestone, southern Taiwan. Journal of Geological Society of
399	China 41:1-24.
400	Gould GC. 2001. The phylogenetic resolving power of discrete dental morphology
401	among extant hedgehogs and the implications for their fossil record. American
402	Museum Novitates 2001:1-52.
403	Gray JE. 1867. Notes on the skulls of the cats (Felidae). Proceedings of the Zoological
404	Society of London. p 258-277.
405	Gunz P, and Mitteroecker P. 2013. Semilandmarks: a method for quantifying curves and
406	surfaces. Hystrix 24:103-109.
407	Hillson S. 2005. Teeth: Cambridge university press.
408	Ho C-K, Qi G-Q, and Chang C-H. 1997. A preliminary study of Late Pleistocene
409	Carnivore fossils from the Penghu Channel, Taiwan. Annual of Taiwan Museum
410	40.



411	Hseu Z-Y, Wang H-H, Wu S-H, and Chang I-S. 2004. Pedogenesis and classification of
412	soils in the Kenting uplifted coral reef nature reserve, Southern Taiwan. Taiwan
413	Journal of Forest Science 19:153-164.
414	Hsu C-C. 2003. Geomorphometric study of Octopus and Cistopus (Cephalopoda:
415	Octopodidae) based on landmarks of beaks. Master Thesis. National Sun Yat-
416	sen University.
417	Hu C-K. 1978. Gongwangling Pleistocene mammalian fauna of Lantian, Shaanxi.
418	Palaeontologica Sinica, New Series C 21:1-45.
419	Huang C-Y, Cheng Y, and Jeh C. 1985. Genesis of the Kenting Formation in the
420	Hengchun Peninsula, southern Taiwan. Ti-Chih (Geology) 6:21-38.
421	Jolliffe IT. 2002. Springer series in statistics. Principal component analysis 29.
422	Kawamura A, Chang C-H, and Kawamura Y. 2016. Middle Pleistocene to Holocene
423	mammal faunas of the Ryukyu Islands and Taiwan: An updated review
424	incorporating results of recent research. Quaternary international 397:117-135.
425	10.1016/j.quaint.2015.06.044
426	Kawamura Y. 1992. Collecting Technique and Storage System for Micro-mammalian
427	Fossils. Honyurui Kagaku (Mammalian Science) 31:99-104.
428	King LM. 2012. Phylogeny of Panthera, Including P. atrox, Based on Cranialmandibular
429	Characters.
430	Lai K-Y. 1989. The fossil record of mammals in Taiwan. Monograph of the Symposium
431	on the Geographical Origin of the Fauna of Taiwan Taipei Zoo, Taipei. p 26-49.
432	Le Roux P, and Skinner J. 1989. A note on the ecology of the leopard {Panthera pardus
433	Linnaeus) in the Londolozi Game Reserve, South Africa. African Journal of
434	Ecology 27:167-171.



435	Li Q. 2007. Fauna analysis of Zhoukoudian Tianyuan Cave human fossil site Master.
436	Chinese Academy of Sciences Institute of Vertebrate Paleontology and
437	Paleoanthropology.
438	Lin C-Y. 2017. Study on Fossil Deer in a Limestone Cave from the Late Pleistocene of
439	Kenting Area, Southern Taiwan. Master. Tunghai University.
440	Linnaeus C. 1758. Systema naturae: Stockholm Laurentii Salvii.
441	Meachen-Samuels J, and Van Valkenburgh B. 2009. Craniodental indicators of prey
442	size preference in the Felidae. Biological Journal of the Linnean Society 96:784-
443	799.
444	Meiri S, Dayan T, and Simberloff D. 2005. Area, isolation and body size evolution in
445	insular carnivores. Ecology Letters 8:1211-1217.
446	Morrison D. 1976. The structure of multivariate observations: I. Principal components.
447	Multivariate Statistical Methods 2:266-301.
448	Otsuka H. 1984. Stratigraphic position of the Chochen vertebrate fauna of the
449	T'ouk'oushan Group in the environs of the Chochen District, Southwest Taiwan,
450	with special reference to its geologic age. Quarterly Journal of the Taiwan
451	Museum 37:37-55.
452	Otsuka H, and Lin C-C. 1984. Fossil rhinoceros from the T'ouk'oushan Group in
453	Taiwan. Journal of Taiwan Museum 37:1-35.
454	Pei W-C. 1934. On the Carnivora from locality 1 of Choukoutien. <i>Paleontologia Sinica</i> ,
455	Ser C 8:76-80.
456	Pei W-C. 1987. Carnivora, Proboscidea and Rodentia from Liucheng Gigantopithecus
457	cave and other caves in Guangxi. Memoirs of Institute of Vertebrate Paleontology
458	and Paleoanthropology, Academia Sinica 18:5-134.



459	POCOCK R. 1930. The pantners and ounces of Asia. Volume II. J Bombay Nat Hist Soc
460	34.
461	Pocock R. 1932. The Leopards of Africa. Proceedings of the Zoological Society of
462	London: Wiley Online Library. p 543-591.
463	Qi G-Q, Ho C-K, and Chang C-H. 1999. The Pleistocene fossil suids from Chochen,
464	Tainan, southwestern Taiwan. Collection and Research:33-40.
465	R Core Team. 2013. A language and environment for statistical computing. R
466	Foundation for Statistical Computing. Vienna, Austria.
467	Rohlf FJ. 2005. tpsDig, digitize landmarks and outlines, version 2.05. Department of
468	Ecology and Evolution, State University of New York at Stony Brook.
469	Rohlf FJ. 2007. tpsRelw version 1.45. Department of Ecology and Evolution, State
470	University of New York, Stony Brook.
471	Shi M, and Wu X. 2011. The microscratch trace of animal bone surface and human
472	behavior study of Kua Hu Qiao Site in Xiaoshan District, Zhejiang Province.
473	Quaternary Sciences 31:723-729.
474	Shih T-T, Tsai W-T, Hsu M-Y, Mezaki S, and Koba M. 1989. The study of ages and
475	terraces of coral reef in the Kenting National Park Area. Construction and
476	Planning Agency Ministry of the Interior Kenting National Park Headquarters
477	Conservation Research Report.
478	Slice DE. 1996. Introduction to landmark methods. Advances in Morphometrics:
479	Springer, 113-115.
480	Stander P. 1997. Field age determination of leopards by tooth wear. African Journal of
481	Ecology 35:156-161.



482	Swinnoe R. 1862. On the Mammais of the Island of Formosa (China). Proceedings of
483	the Zoological Society of London: Wiley Online Library. p 347-368.
484	Swinhoe R. 1870. Catalogue of the mammals of China (south of the Yangtsze) and of
485	the Island of Formosa. Proceedings of Zoological Society of London. p 615-653
486	Teilhard C, and Pei W. 1941. The fossil mammals from Locality 13 of Choukoutien.
487	Palaeontologia Sinica. New Ser C:1-106.
488	Todd NE. 2010. New phylogenetic analysis of the family Elephantidae based on
489	cranial-dental morphology. The Anatomical Record: Advances in Integrative
490	Anatomy and Evolutionary Biology 293:74-90.
491	Tseng JZ, Wen H, and Chen S-Q. 2010. Geometric morphometrcs analysis of cranial
492	shape among Late Miocene Hyaenid Ecomorphologies in the Linxia Basin,
493	Gansu, China. Vertebrata Palasiatica 48:235-246.
494	Van Cleef-Roders JT, and Van Den Hoek Ostende LW. 2001. Dental morphology of
495	Talpa europaea and Talpa occidentalis (Mammalia: Insectivora) with a
496	discussion of fossil Talpa in the Pleistocene of Europe. Zoologische
497	Mededelingen 75:51-68.
498	Wang K-C. 2015. Rodent Fossils from Late Quaternary Limestone Cave in Kenting
499	Area, Southern Taiwan Master. National Cheng Kung University.
500	Wei K-Y. 2007. Quaternary mammalian fossils of Taiwan: an eclectic overview and
501	prospects for future study. Special Publication of the Central Geological
502	Survey:261-286.
503	Xu M-Y. 1989. A Geomorphological Study of Marine Terraces in Taiwan PhD. Chinese
504	Culture University.



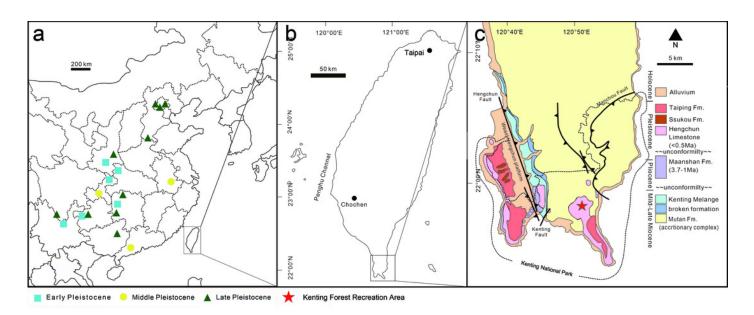


505	Yen T-P, and Wu C-Y. 1986. The Pliocene and younger formations in the Southern part
506	of the Hengchun Peninsula. Ti-Chih (Geology) 7:1-10.
507	Zelditch ML, Fink WL, and Swiderski DL. 1995. Morphometrics, homology, and
508	phylogenetics: quantified characters as synapomorphies. Systematic Biology
509	44:179-189.
510	Zelditch ML, Lundrigan BL, and Garland Jr T. 2004. Developmental regulation of skull
511	morphology. I. Ontogenetic dynamics of variance. Evolution & Development
512	6:194-206.
513	
514	



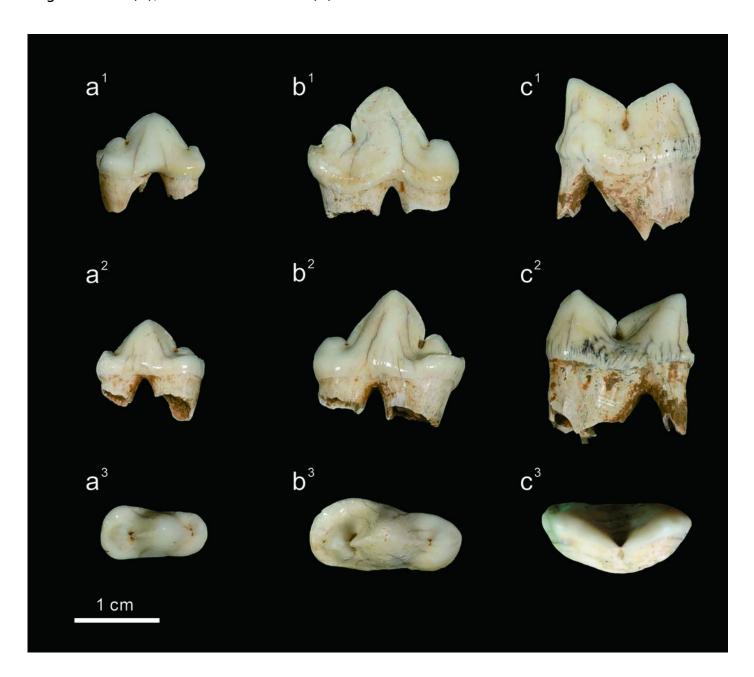
Distribution of the Pleistocene leopard fossils in East Asia

(a), map of Taiwan showing the location of the Kenting National Park (b), and a geological map of southernmost Taiwan (c). The location of the excavation site, Longshia-dong Cave, is indicated by a red star in (c).



The three felid lower cheek teeth from Longshia-dong Cave

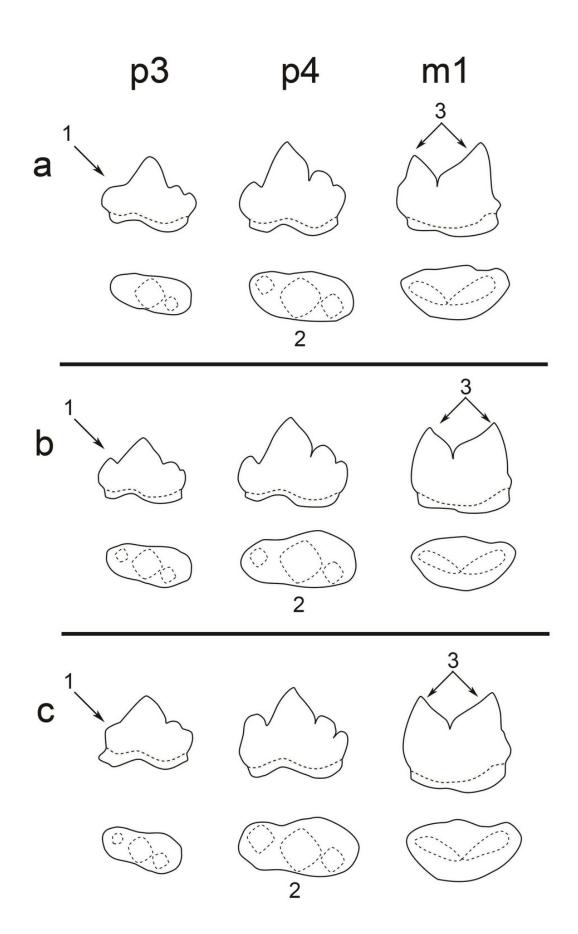
including p3 (a, F056584), p4 (b, F056585), and m1 (c, F056585), and their buccal views (1), lingual views (2), and occlusal views (3).





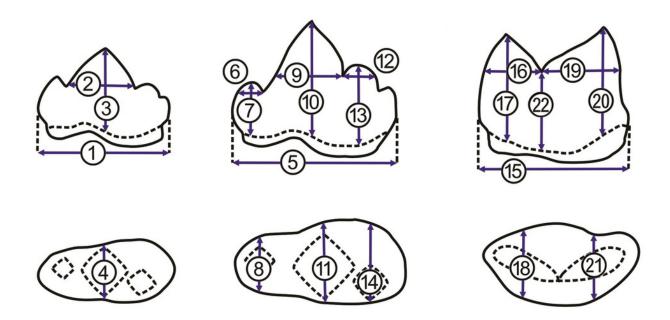
A comparison of the teeth (p3, p4, and m1) of three pantherines

including (a) clouded leopard, (b) leopard, and (c) tiger. Three significant characters are indicated by (1) the developmental level of p3 paraconid, (2) the shape of p4 occlusal surface (the difference between paraconid and protoconid widths), and (3) the size difference between m1 paraconid and m1 protoconid.



Measurements of p3, p4, and m1 from the buccal side (a, from left to right) and from the occlusal side (b, from left to right).

For the numbered dimensions, see Table 1.

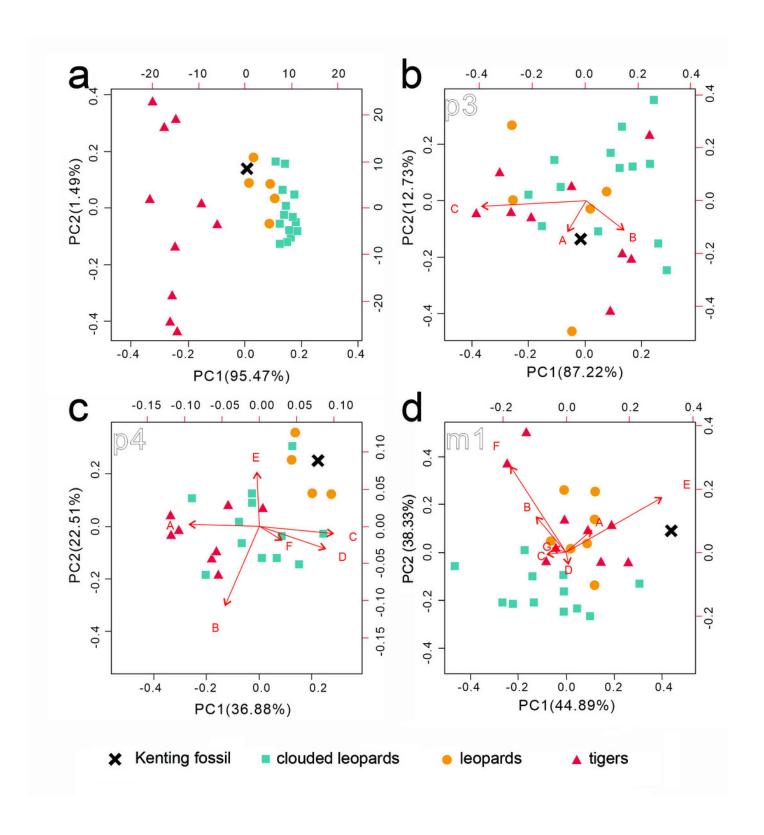




PCA analysis based on all datasets, Red triangles, *Panthera tigris*; yellow circles, *Panthera pardus*; cyan square, *Neofelis* sp.

black cross, the studied fossil felid teeth (a), three ratios from four p3 numbered dimensions (A: ②/①, B: ③/①, C: ②/③ in b), six ratios from seven p4 numbered dimensions (A: ⑥/⑤, B: ⑦/⑤, C:⑨/⑤, D: ⑩/⑥, E: □/⑥, F: □/⑥ in c), and seven ratios from eight m1 numbered dimensions (A: □/□, B: □/□, C: □/□, D: □/□, E: □/□, F: □/□, G: □/□ in d) (see the details of the numbered dimensions in Fig. 3 and Tab. 1). Red arrow represents a simplified ratio that shows the trend relating to the two principal components. Factor loadings of each principal component is indicated by the arrow length.

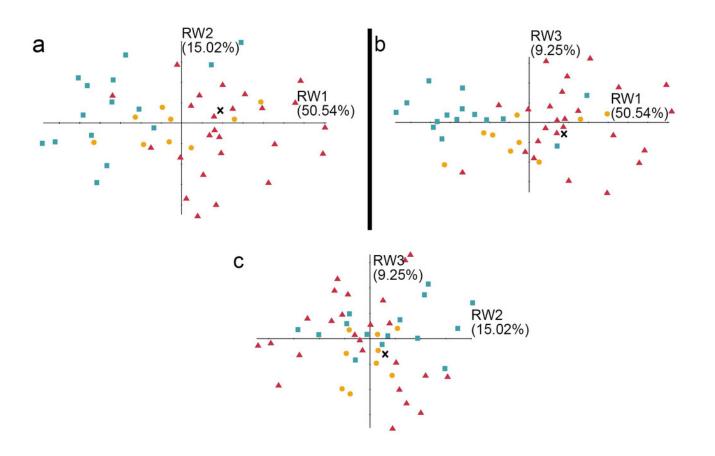






The shape variation of the occlusal surface of p3 in three pantherines, as revealed by a principal components analysis of three warp scores.

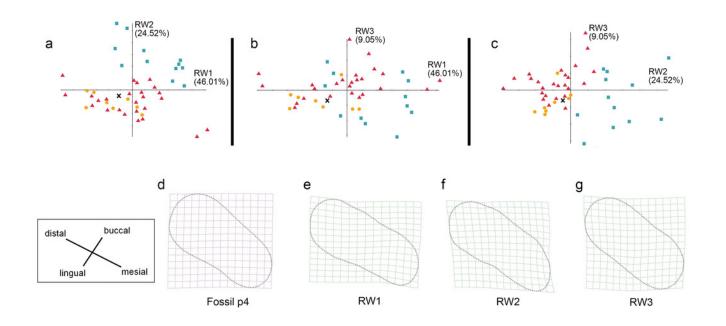
(a) RW1 versus RW2; (b) RW1 versus RW3; (c) RW2 versus RW3. RW1 in a positive direction explains 50.54% variance, RW2 in a positive direction explains 15.02% variance, and RW3 in a positive direction explains 9.25% variance. Red triangles, *Panthera tigris*; yellow circles, *Panthera pardus*; cyan square, *Neofelis* sp.; black cross, the studied fossil felid teeth.





The shape variation of occlusal surface of p4 in three pantherines, as revealed by a principal components analysis of warp scores.

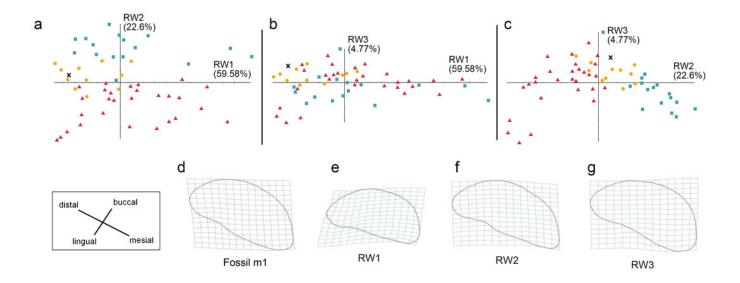
Plot of (a)RW1 versus RW2; (b) RW1 versus RW3; (c) RW2 versus RW3. Red triangles, Panthera tigris; yellow circles, Panthera pardus; cyan square, Neofelis sp.; black cross, the studied fossil felid teeth. (d) the shape of fossil p4, (e) thin-plate spline deformation grid depicting shape variation along RW1 in a positive direction explains 46.01% variance, (f) thin-plate spline deformation grid depicting shape variation along RW2 in a positive direction explains 24.52% variance, (g) thin-plate spline deformation grid depicting shape variation along RW3 in a positive direction explains 9.05% variance.





The shape variation of occlusal surface of m1 in three pantherins, as revealed by a principal components analysis of warp scores.

(a) RW1 versus RW2; (b) RW1 versus RW3; (c) RW2 versus RW3.Red triangles, *Panthera tigris*; yellow circles, *Panthera pardus*; cyan square, *Neofelis* sp.; black cross, the studied fossil felid teeth. (d) the shape of fossil p4; (e) thin-plate spline deformation grid depicting shape variation along RW1 in a positive direction which explains 59.58% variance, (f) thin-plate spline deformation grid depicting shape variation along RW2 in a positive direction explaines 22.6% variance; (g) thin-plate spline deformation grid depicting shape variation along RW3 in a positive direction explains 4.77% variance.





Bivariate plots of selected dimensions (①, ④ , ⑤, \Box , \Box , \Box from Table 1) of p3, p4, m1 of the fossil teeth and extant leopards (in mm).

Dotted lines are the regression lines for the dimensions of the five extant specimens. The regression line for the dimensions of the fossil specimen is marked in red. Triangles, p3; squares, p4; circles, m1.

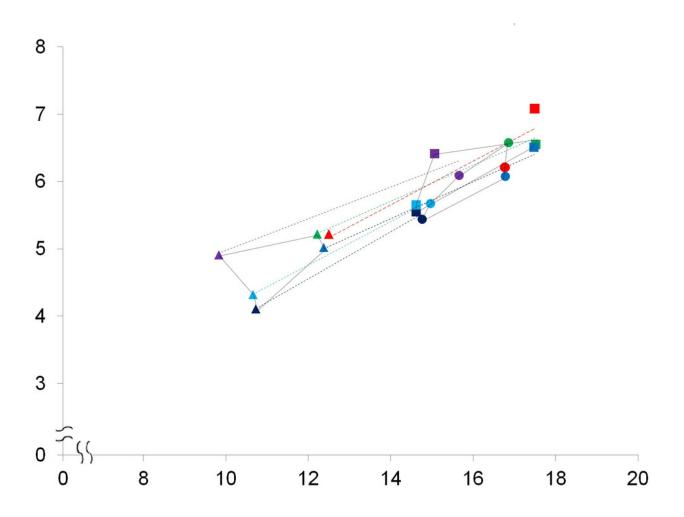




Table 1(on next page)

Selected dimensions of teeth



Table 1 Selected dimensions of teeth

dimensio	
n	
1	p3 crown length
2	p3 protoconid length
3	p3 protoconid height
4	p3 protoconid crown width
5	p4 crown length
6	p4 paraconid length
7	p4 paraconid crown height
8	p4 paraconid crown width
9	p4 potoconid length
10	p4 protoconid crown height
11	p4 protoconid width
12	p4 hypoconid length
13	p4 hypoconid height
14	p4 hypoconid crown width
15	m1 crown length
16	m1 paraconid length
17	m1 paraconid height
18	m1 paraconid width
19	m1 protoconid length
20	m1 protoconid height



PeerJ

21	m1 protoconid width
22	m1 carnassial notch height

2



Table 2(on next page)

Morphological comparisons of three extant species and the fossil. \bigcirc , present; -, absent; \triangle , uncertain.



- 1 Table 2 Morphological comparisons of three extant species and the fossil. o, present; -, absent;
- 2 Δ, uncertain.

	p3 paraconid	m1 metaconid	size comparison between the paraconid, and the protoconid, of m1	chubby talonid on p3 and p4	occlusal surface	
Longshia-dong Cave	0	-	similar	0	intermediate	
fossils						
Modern clouded leopards	-	Δ	bigger	-	narrow	
(Neofelis sp.)			paraconid			
Modern tigers	0	-	similar	0	intermediate	
(Panthera tigris)						
Modern leopards	Δ	-	similar	0	chubby	
(Panthera pardus)						



Table 3(on next page)

Comparisons of dental size measurements (mm) of Longshia-dong Cave fossil to Chinese leopard fossils (expressed by length*width).



- 1 Table 3 Comparisons of dental size measurements (mm) of Longshia-dong Cave fossil to
- 2 Chinese leopard fossils (expressed by length*width).

	Longshia-dong	Niuyan Cave			Zhoukoudian		Gongwangling	recent	
	Cave								
	(Kenting, Taiwan)	(Beijing, China)			(Beijing	g, China)	(Shaanxi, China)		
	-	V11799	V11800	V11801	location	location	V2980		
					1st	13th			
p3	12.5*6.39	13.4*6.5	-	-	16.3*9	16*7.5	14.4*8	14.8*7.9	12*6.3
p4	17.49*8.51	18.2*9.7	18.5*9.5	-	23.2*12	21*10	21*11.2	22.3*11	18*9.4
m1	16.77*7.95	19.3*9.2	19.8*8.5	19.5*8.5	24*12.2	21*10	22.2*12	22.7*12.3	18.3*8.6