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1	Title: A leopard ate a chimpanzee: The first evidence from East Africa
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19 Introduction

20	Primates may have suffered predation pressures throughout their evolutionary history.
21	Hominoids have been sympatric with large carnivores since the early Miocene in Africa
22	(Werdelin and Peigné, 2010), and it is thought that predation pressure by large
23	carnivores has played a significant role in their evolution (Hart and Sussman, 2005). For
24	example, carnivore predation on <i>Proconsul</i> has been inferred from site R114
25	("Pot-hole") on Rusinga Island, Kenya, where the partial skeleton of <i>P. heseloni</i>
26	KNM-RU 2036 was recovered (Walker and Shipman, 2005).
27	In addition to fossil evidence, data on predation on living primates is also
28	important for reconstructing the predation pressure on our human ancestors. Among
29	potential nonhuman predators of living African apes, leopards (Panthera pardus) and
30	lions (Panthera leo) have been known to actually prey upon apes. There has been only
31	one report of lion predation on apes (Tsukahara, 1993). Lions are usually allopatric with
32	apes because they are absent from tropical rainforests (Nowell and Jackson, 1996)
33	where the majority of apes live. On the other hand, because leopards occur in most parts
34	of sub-Saharan Africa (<i>ibid</i> .), they may be more likely than lions to prey upon apes.
35	There has been limited information on leopard predation on apes, and all data have
36	come from West and Central Africa (Table 1). Moreover, despite long-term research on

37	chimpanzees (Pan troglodytes schweinfurthii) at several sites in East Africa, no
38	instances of leopard predation have been reported. This is probably because of the
39	recent extirpation of leopards from most of the research sites. According to personal
40	communications from experienced field researchers, there has been no evidence of the
41	presence of leopards for a decade or more at the research sites of Gombe (Wilson ML)
42	in Tanzania, and Kalinzu (Hashimoto C), Kibale (Struhsaker T, Mitani JC, and Mills
43	DR), and Budongo (Newton-Fisher NE) in Uganda. The only exception is Mahale in
44	Tanzania where leopards have lived sympatrically with chimpanzees, without evidence
45	of predation by the former on the latter (Nishida 2012). There have been several reports
46	of encounters between leopards and chimpanzees from Tanzania including Mahale
47	(reviewed in Pierce, 2009). Responses of chimpanzees to leopards varied from emitting
48	loud, fearful calls, vigilance, and acting in a threatening manner (e.g., Pierce, 2009);
49	stalking a leopard that had called in the distance (Mitani JC, personal communication);
50	to surrounding a den and killing a cub (Hiraiwa-Hasegawa et al., 1986).
51	Boesch (2009) asserted that all well-studied East African chimpanzee
52	populations face little or no predation pressure. Although he recognized the presence of
53	leopards at Mahale, he insisted that leopards were rare there, so that predation was
54	negligible. During a systematic survey of leopard scats, we found the first evidence of

the consumption of an eastern chimpanzee at Mahale. Here, we report the details of thisevidence.

57

58 Methods

We collected leopard scats in the Mahale Mountains National Park, Tanzania (Nishida, 592012) for 41 days in June–August 2012. Although other large carnivores (lions, hunting 60 dogs, and hyenas) are reported to inhabit the park, no direct or indirect evidence of the 61 62former 2 species has been observed in the study area for more than 2 decades (also, no 63 domestic dogs have been confirmed in the area). Hyena scats were observed in 2005 64 and 2008 but were distinguishable from those of leopards by its very whitish appearance and finer digestion of bones. Thus, it is unlikely that we misidentified leopard scats with 65 those of other species. We walked observation trails within the home range of the 66 67 habituated M group chimpanzees (387 min/day on average). When a scat was found, we 68 recorded its location with GPS and carried it back to our camp. After being dried and weighed, we inspected its contents for hair, bones, and skin. To confirm whether the 69 contents were of chimpanzees, we conducted morphological investigations and DNA 7071analysis. Taxonomic identification was based on previous work on comparative primate postcranial morphology (Ward et al., 1995; Nakatsukasa et al., 2003). DNA was 72

73	extracted from a small bone using a TBONE EX KIT (DNA Chip Research Inc., Japan)
74	and a 331-base pair segment of the mitochondrial hypervariable control region (Inoue et
75	al., 2011) was analyzed. We also examined the XY homologous gene amelogenin for
76	sex identification and 8 microsatellite loci for individual identification (Inoue et al.,
77	2008).

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79 Results
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80	We collected 142 leopard scats, among which one contained the right and left patellae,
81	the distal end of a manual proximal phalanx, and an intact manual intermediate phalanx
82	of a chimpanzee (Fig. 1). The patellae are morphologically similar and almost identical
83	in size. The completely fused proximal epiphysis of the intermediate phalanx,
84	morphology of the ligamentous insertion on the patella and modestly developed flexor
85	sheath ridges on the phalanges suggest these bones belonged to an adult (not old), and
86	probably a female based on general size and robusticity criteria. The distal part is
87	chewed off and trabeculae are exposed dorsally in both patellae (Fig. 1 a, b). The break
88	surface is mediolaterally long and gently concave. The dorsal (= posterior) break edge is
89	more proximal than in the ventral (= anterior) break edge, which approximates the
90	original distal border. The missing part includes the whole attachment area of the

91	patellar ligament. The cut surface suggests that the leopard filled its mouth with the
92	distal part of the quadriceps femoris muscles and the patella and tried to cut off the
93	mouth infill from the remaining carcass. Probably, the patella (and the ligament) was
94	not fully turned over, and the leopard's carnassials could not reach the patellar ligament
95	but only the ligament attachment area. The right patella also has a bite mark on the
96	lateral rim (Fig. 1 b). There is no gnaw mark on the intermediate phalanx (Fig. 1 c).
97	Along the proximal epiphysis of the intermediate phalanx, the cortex is damaged,
98	probably caused by the acid (or acidotic erosion and physical stress). The break on the
99	proximal phalanx is a common fracture pattern.
100	The scat was found on June 18, 2012, at the beginning of the dry season. It was
100 101	The scat was found on June 18, 2012, at the beginning of the dry season. It was not very fresh but estimated to be no older than 4 months from its appearance and the
101	not very fresh but estimated to be no older than 4 months from its appearance and the
101 102	not very fresh but estimated to be no older than 4 months from its appearance and the extent of preceding rainfalls (the main factor in scat decay). The location (Fig. 2) is
101 102 103	not very fresh but estimated to be no older than 4 months from its appearance and the extent of preceding rainfalls (the main factor in scat decay). The location (Fig. 2) is within the M group's home range and is used exclusively by the group (Nakamura et al.,
101 102 103 104	not very fresh but estimated to be no older than 4 months from its appearance and the extent of preceding rainfalls (the main factor in scat decay). The location (Fig. 2) is within the M group's home range and is used exclusively by the group (Nakamura et al., 2013). We had seen and heard evidence of leopards (foot prints, roars, etc.) on 10
101 102 103 104 105	not very fresh but estimated to be no older than 4 months from its appearance and the extent of preceding rainfalls (the main factor in scat decay). The location (Fig. 2) is within the M group's home range and is used exclusively by the group (Nakamura et al., 2013). We had seen and heard evidence of leopards (foot prints, roars, etc.) on 10 different days within the preceding month. On June 13, a field assistant had observed a

109	their mothers were available (Table 2). Analyses of DNA extracted from a phalanx of
110	the victim (hereafter "Bone2012") reconfirmed that Bone2012 was actually a female
111	chimpanzee. Her mitochondrial haplotype was B, a common haplotype in the M group
112	(Inoue et al., 2011), but among candidates, only a male AG had this haplotype.
113	Microsatellite analyses also showed that two candidates (CA and TZ) had different
114	alleles from Bone 2012 at 5 or more loci. Mothers of 3 other candidates (AG, ME, and
115	TZ09) did not share alleles with Bone2012 at 2 loci. Therefore, we conclude that
116	Bone2012 was not an individual from the M group.
117	
118	Discussion

- 119 We confirmed that a leopard ate an adult female chimpanzee at Mahale. This is the first
- evidence of leopard consumption of eastern chimpanzees (P. t. schweinfurthii), which 120
- 121adds another subspecies to the list of apes consumed by leopards.

The scat was found within the M group's home range but the victim was not 122

- 123from the group. Because the home range of a different chimpanzee group reaches to
- approximately 1 km north of the scat location, a small distance compared to known 124
- 125leopard range sizes (Jenny, 1996), the leopard may have eaten a chimpanzee of that
- group, then moved into the M group's home range and defecated. Alternatively, it may 126

127	have eaten an unknown female right after her emigration from her natal group to the M
128	group. If the leopard actually killed the victim, this means that a leopard can prey on a
129	full adult female chimpanzee. Female chimpanzees are smaller in body size, less
130	aggressive, and less gregarious than males, and thus could be more prone to predation.
131	The predation risk would be even higher when females transfer alone between
132	unit-groups. Thus, we might need to take such potential risk from female transfer into
133	account when discussing how a female-dispersal social structure, like that of
134	chimpanzees, could have evolved.
135	Contrary to the assumption of Boesch (2009) that no eastern chimpanzee
136	research sites have as many leopards as in the Taï forest, we found comparable, or even
137	greater, numbers of scats per day than at Taï or Lopé (Table 1). Nevertheless, we have to
138	be cautious about direct comparisons because research efforts might differ. Evidence of
139	leopards was quite frequent during the study period. On the basis of the 1-2 leopard
140	sightings per year at Taï (Boesch and Boesch-Achermann, 2000, Table A.2), we cannot
141	conclude that leopards are rarer in Mahale than in Taï.
142	Because leopards sometimes do scavenge (Bailey, 1993), we cannot determine
143	from scat evidence alone whether the leopard killed the chimpanzee or scavenged a
144	corpse. However, observations of 3 seriously injured M group chimpanzees may suggest

145	occasional leopard attacks on chimpanzees. In 2009, one male chimpanzee had a much
146	deeper wound than usually caused by fights among male conspecifics. Further, in 2011,
147	a mother-infant pair was found injured and the mother had 3 long, parallel scars from
148	the head to the back, seemingly caused by claws. Because researchers had thought that
149	leopards did not eat chimpanzees at Mahale (Nishida, 2012), they had assumed that all
150	serious injuries came from intraspecific fights. Now, because there has been confirmed
151	leopard consumption of a chimpanzee, we should also consider attacks by leopards as a
152	possible cause of injury or death for Mahale chimpanzees. Again, if the victim was
153	actually killed by a leopard, this means that predation pressure has been underestimated
154	for more than 40 years at Mahale or, alternatively, that leopards have recently shifted
155	their diet to include chimpanzees. If the former is the case, this might imply a wider
156	underestimation of predation pressures on primates in general. Because predation events
157	are rarely observable but can still be significant to primates' behavioral and social
158	evolution, we might need to acquire a better picture of carnivore-primate relationships
159	with long-term data. The ecology and behavior of Mahale leopards have yet to be
160	studied. Because information on predation pressure on living apes is still too scarce to
161	be used to infer the predation pressure on fossil hominins, we may need to investigate
162	further the ecology and behavior of leopards that are currently sympatric with apes.

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- 175

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Study site (Country)	Species/subspecies	Eviden	Source		
	-	Leopard scats ¹	Indirect evidence	Hoppe-Dominik, 1984	
Taï (Côte d'Ivoire)	Pan troglodytes verus	0/215 (15 months)	Three deaths possibly by		
		1/200 (1 year)	leopards	Boesch 1991; 2009,	
				Zuberbühler and Jenny	
				2002	
Dzanga-Sangha	Gorilla gorilla gorilla	Yes ²	N/A	Fay et al., 1995	
(Central African					
Republic)					
Lopé (Gabon)	G. g. gorilla	6/196 (8 years)	N/A	Henschel et al., 2005	
	P. t. troglodytes	4/196 (8 years)			
Petit Loango (Gabon)	P. t. troglodytes	N/A	Leopard scats and footprints	Furuichi, 2000	
			near chimpanzee corpse		
Lui Kotal	P. paniscus	Yes ²	N/A	D'Amour et al., 2006	
(Democratic Republic					
of Congo)					

Table 1 Reported leopard predation/consumption of African apes 231

¹ The figures indicate the number of scats containing ape bones/the total number of scats collected. The duration of fecal collection is shown in parentheses. ² No mention of the number of scats.

 $\begin{array}{c} 232\\ 233 \end{array}$

- Table 2 DNA profile of the chimpanzee bone found in a leopard scat in comparison with those of M group chimpanzees that 234
- disappeared in 2012 235

Names of	Age	Sex	Mitochondrial	Microsatellite loci ¹							
chimpanzees	(years)		haplotype ¹	D9s910	D11s2002	D2s1329	D12s66	D2s1326	D5s1470	D7s2204	D7s817
Bone2012	adult?	F	В	104/104	148/148	178/202		182/218	190/190	245/249	116/116
Candidates											
CA	52 ³	F	<u>C</u>	104/110	<u>148/156</u>	<u>182/198</u>	158/182	<u>182/186</u>		249/253	<u>112/124</u>
ΤZ	30 ³	F	<u>A</u>	104/110	148/148	<u>178/198</u>	154/154	<u>202/206</u>	<u>186/190</u>	245/249	144/148
AG^2	7	<u>M</u>	В	104/104	148/148	178/182	150/182	150/182	<u>194/194</u>	245/253	<u>148/152</u>
ME^2	10	F	<u>C</u>	104/113	148/148	<u>186/198</u>	182/182	182/182	186/190	249/249	<u>148/152</u>
TZ09 ²	2	F	A	104/110	148/148	178/198	154/154	<u>202/206</u>	186/190	245/249	<u>144/148</u>

236

¹Mitochondorial haplotype and microsatellite genotypes of the candidates are described in Inoue et al. (2011) and Inoue et al. (2008), respectively. ²DNA data from their mothers are shown because their own DNA is not available. A mother shares a mitochondrial haplotype and at least one allele at all 237loci with offspring. 238

³Estimated age 239

Underlined data show mismatches with Bone 2012. 240

242 Figure 1

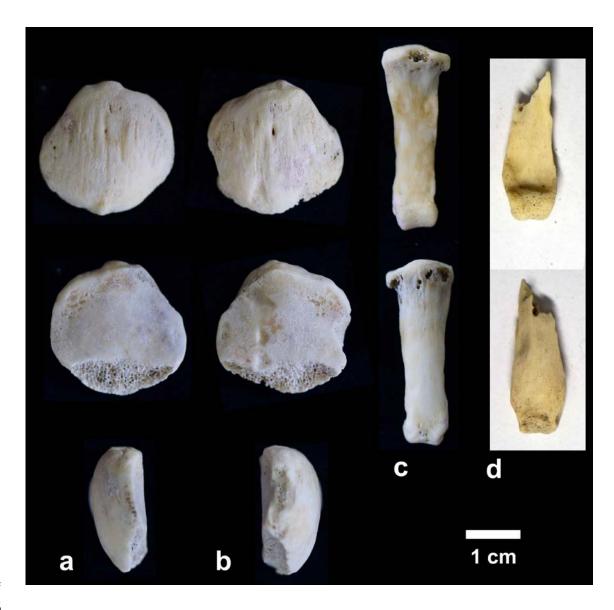
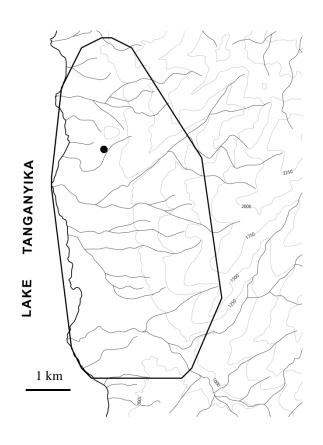


Figure 2





249 **Figure Legends**

- **Figure 1.** Recovered chimpanzee bones. a, b: left and right patellae, respectively (from
- top to bottom: anterior, posterior, and lateral views). c: middle phalanx (top: palmar
- view). d: distal fragment of a proximal phalanx (top: palmar view).

- **Figure 2.** The location where the leopard scat with chimpanzee bones was found (black
- dot) in relation to the home range of the M group chimpanzees (a polygon).