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4 Predation of a Brazilian porcupine (*Coendou prehensilis*) by an ocelot (*Leopardus pardalis*) at a
5 mineral lick in the Peruvian Amazon

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21 **Abstract**

22 Many herbivorous and frugivorous Amazonian species, including several arboreal
23 animals, feed on earth and water at mineral licks in the Amazon region to supplement their diet
24 with micronutrients and clays. These species are vulnerable to predation during this activity. We
25 recorded an adult Brazilian porcupine (*Coendou prehensilis*) being predated by an adult male
26 ocelot (*Leopardus pardalis*) while drinking water at a mineral lick in the Maijuna-Kichwa
27 Regional Conservation Area (MKRCA) in northeastern Loreto, Peru. This observation provides
28 direct evidence arboreal species like the porcupine, which move slowly on the ground, are
29 particularly vulnerable to terrestrial predators while visiting mineral licks. Mineral licks are
30 important in the diets and ecology of Amazonian mammals, but arboreal prey must balance the
31 trade-off between using the resource and being hunted. We suggest that mineral licks may be
32 hotspots of risk in Amazonian prey species' landscape of fear.

33

34 **Keywords**

35 Felids, predation, mineral lick, ecology, behavior, Amazonia

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40 *Conflicts of interests*

41 The authors declare that they have no conflict of interest.

42 *Ethics approval*

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45 *Consent to participate*

46 Not applicable.

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48 Not applicable.

49 *Availability of data and material*

50 Our paper is an observational study, and does not include data beyond that which is
51 presented in the paper.

52 *Code availability*

53 We did not use code in our study.

54 *Authors’ contributions*

55 All authors contributed to the study conception and design. Data collection and analysis were
56 performed by Brian Griffiths. The first draft of the manuscript was written by Brian Griffiths and
57 all authors commented on previous versions of the manuscript. All authors read and approved the
58 final manuscript.

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61 Mineral licks are naturally occurring sites where animals exhibit geophagical behavior,
62 consuming soil or mud to supplement their diet with micronutrients or clays (Dudley et al., 2012;
63 Kreulen, 1985). Mineral licks are visited frequently by many different Amazonian herbivores
64 and frugivores, such as the Brazilian tapir (*Tapirus terrestris*), collared (*Pecari tajacu*) and
65 white-lipped peccary (*Tayassu pecari*), red brocket deer (*Mazama americana*), paca (*Cuniculus*
66 *paca*), black agouti (*Dasyprocta fuliginosa*), and red howler monkey (*Alouatta seniculus*)
67 (Montenegro, 2004; Tobler, Carrillo-Percestequi, & Powell, 2009; Tobler, 2008). Results from
68 several studies have indicated that mineral licks in Amazonia are also important sites for
69 predators (e.g. Izawa, 1993; Link et al., 2011; Link and Fiore, 2013; Matsuda and Izawa, 2008;
70 Montenegro, 2004) such as wild felids, seeking prey that visit the lick to consume soil. The
71 prevalence of predators at lick sites suggests that mineral licks may act as hotspots of risk in the
72 landscape of fear (Laundré et al., 2010) for prey species.

73 Behavioral studies provide further evidence that mineral licks form a key component of
74 the landscape of fear, particularly for arboreal species visiting the lick that are otherwise
75 inaccessible to terrestrial predators. For example, Link and Fiore (2013) and Ospina (2011)
76 showed that large-bodied primates increased their group size before descending to a lick, likely
77 to increase the probability of detecting a predator. In a study by Link et al. (2011), primates
78 visited licks most frequently on bright sunny days when visibility was highest, and employed
79 vigilance behavior from the trees while members of the group fed at the lick. Similar antipredator
80 behavior has been observed in the grouping behavior of birds visiting licks, in the formation of
81 mixed-species flocks (Brightsmith and Villalobos, 2011) that also employ vigilance and induce a
82 dilution effect (Hamilton, 1971). One study by Montenegro (1998) described antipredator

83 behavior of tapir at licks, although no direct observations of predation on species other than
84 primates at mineral licks have been published.

85 We describe a predation event by an adult male ocelot (*Leopardus pardalis*) on an adult
86 Brazilian porcupine (*Coendou prehensilis*) at a mineral lick in the Maijuna-Kichwa Regional
87 Conservation Area (MKRCA) (-72.9311° W, -3.15442° S), a 391,000 hectare protected area
88 collaboratively managed by the indigenous Maijuna people in the northern part of the state of
89 Loreto in the northeastern Peruvian Amazon. The MKRCA is characterized by tropical, humid,
90 primary *terra firma* forest, floodplain forest, and a terraced habitat that is unique to the region
91 (Gilmore, 2005). The region has a mean annual rainfall of about 3100 mm and temperature of
92 26° C (Marengo, 1998).

93 Ocelots are a medium-bodied felid (6-18 kg) (de Oliveira et al., 2010) found throughout
94 South and Central America (de Oliveira et al., 2010; Fernandez, 2002). Terrestrial mammals
95 make up the bulk of the ocelot's diet, but insects, crabs, reptiles, birds, sloths, monkeys, and
96 other arboreal mammals are also common prey items. Ocelots are mainly recorded as hunting
97 from the ground, and so their mechanisms of hunting arboreal species are unknown (Moreno et
98 al., 2006), although it is assumed that many of these prey are hunted when they are scared to the
99 ground by the ocelot or are at the ground by some other circumstance (Bianchi and Mendes,
100 2007).

101 The Brazilian porcupine (*Coendou prehensilis*) is an arboreal (Roberts et al., 1985),
102 nocturnal hystricognath rodent of the Erethizontidae family that is widely distributed in moist,
103 humid forests throughout Amazonia (Eisenberg, 1978; Mares and Ojeda, 1982). *C. prehensilis* is
104 exclusively herbivorous, with a diet that consists of mainly buds, bark, fruits, and seeds (Charles-
105 Dominique et al., 1981; Eisenberg, 1978). The Brazilian porcupine is known to descend to the

106 forest floor to visit mineral lick sites across Amazonia (Gilmore and Young, 2010; Molina et al.,
107 2014).

108 Images of the predation event were recorded as part of a camera trapping study on
109 mineral licks in the Sucusari Basin, in the southern reaches of the MKRCA, on October 8, 2018
110 at 9:47pm. Camera traps (Bushnell Aggressor) were installed at 80 mineral licks in Sucusari
111 from August, 2018, to June, 2019 to measure mammal visitation at licks across the region. The
112 locations of the mineral licks were determined through participatory mapping exercises with
113 Majuna hunters in July, 2017. Camera traps were set to photo mode, taking a burst of three
114 images every time the motion sensor was activated. A delay of fifteen seconds between each set
115 of images was chosen to avoid expending the camera's batteries prematurely. Cameras were set
116 at a minimum of 50 centimeters from the ground, facing the active "face" and entrances to the
117 lick, following (Tobler et al., 2009). The locations of the face and entrances were determined
118 from signs of animal activity. In mineral licks that hunters noted flooded seasonally, cameras
119 were placed farther from the ground to avoid inundation, just above the high-water mark on
120 nearby vegetation. Camera traps were left at mineral licks for a minimum of 82 days and were
121 not disturbed after they were set.

122 Fifteen images of the ocelot stalking and attacking the porcupine were recorded (Fig. 1).
123 The ocelot stalked the porcupine for over one minute while the porcupine was drinking water in
124 the mineral lick and readjusted its position before pouncing on the porcupine and dragging it out
125 of the water. Over seven minutes pass between when the ocelot pounced and when it dragged the
126 porcupine from the water; presumably, it was killing the porcupine in the mineral lick during this
127 time. It is likely that the camera did not sense movement for seven minutes because the camera
128 was located behind and slightly above a fallen log and the mud in the lick was quite deep. While

129 the ocelot is stalking the porcupine, it does not have any mud on its fur. When the ocelot
130 pounced, dragging the porcupine into the mud behind the log, both animals were likely hidden
131 from view of the camera. When the ocelot pulls the porcupine onto the log, both animals are
132 covered in mud.

133 One previous study reports ocelots preying on porcupines, from a single scat sample
134 (Emmons, 1987), but it is thought that ocelots hunt arboreal species opportunistically from the
135 ground, including at mineral lick sites (Izawa, 1993; Link et al., 2011; Matsuda and Izawa,
136 2008). Arboreal species like the Brazilian porcupine, which move slowly on the ground may be
137 particularly vulnerable to terrestrial predators while visiting mineral licks, highlighting the
138 importance of mineral licks in the diets of Amazonian mammals.

139 While the importance of mineral licks to the diets of herbivorous mammals has been well
140 studied, the ecological role of Amazonian mineral licks to predators is largely unknown. Studies
141 targeting mineral licks in other regions have described predation risk at licks for other prey
142 species. For example, Couturier and Barrette (1988) observed moose in Quebec, Canada
143 exhibiting vigilance behavior at mineral licks. A seminal work by Cowan and Brink (1949)
144 described heavy predation on wild goats by bears and mountain lion at mineral licks in Canada.
145 Large felids were recorded visiting mineral licks in Nepal by Moe (1993), likely looking for
146 prey. Behavioral studies at Amazonian mineral licks have largely been limited to large-bodied
147 primates although they make up only a small portion of the species that frequent the licks. One
148 exception is a study done by Montenegro (1998), which described antipredator behavior by the
149 tapir at Amazonian licks.

150 Given the observations of predators and direct predation events at mineral licks,
151 including this one, we suggest that further research is needed to understand predator-prey

152 interactions at mineral licks for species beyond primates. Further behavioral studies in particular
153 will provide context as to what extent mineral licks present hotspots of risk in prey species’
154 landscape of fear in Amazonia.

155

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161 **Literature Cited**

- 162 Bianchi, R.D.C., Mendes, S.L., 2007. Ocelot (*Leopardus pardalis*) predation on primates in
163 Caratinga Biological Station, southeast Brazil. *Am. J. Primatol. Off. J. Am. Soc.*
164 *Primatol.* 69, 1173–1178.
- 165 Brightsmith, D.J., Villalobos, E.M., 2011. Parrot Behavior at a Peruvian clay lick. *Wilson J.*
166 *Ornithol.* 123, 595–602.
- 167 Charles-Dominique, P., Atramentowicz, M., Charles-Dominique, M., Gerard, H., Hladik, A.,
168 Hladik, C.M., Prévost, M.-F., 1981. Les mammifères frugivores arboricoles nocturnes
169 d’une forêt guyanaise: inter-relations plantes-animaux. *Rev. Décologie.*
- 170 Couturier, S., Barrette, C., 1988. The behavior of moose at natural mineral springs in Quebec.
171 *Can. J. Zool.* 66, 522–528.
- 172 Cowan, M., Brink, V.C., 1949. Natural game licks in the rocky mountain national parks of
173 Canada. *J. Mammal.* 30, 379–387.

174 de Oliveira, T.G., Tortato, M.A., Silveira, L., Kasper, C.B., Mazim, F.D., Lucherini, M., Jácomo,
175 A.T., Soares, J.B.G., Marques, R.V., Sunquist, M., 2010. Ocelot ecology and its effect on
176 the small-felid guild in the lowland neotropics. *Biol. Conserv. Wild Felids* 559–580.

177 Dudley, R., Kaspari, M., Yanoviak, S.P., 2012. Lust for salt in the western Amazon. *Biotropica*
178 44, 6–9. <https://doi.org/10.1111/j.1744-7429.2011.00818.x>

179 Eisenberg, J.F., 1978. The evolution of arboreal herbivores in the class Mammalia. *Ecol.*
180 *Arboreal Folivores* 135–152.

181 Emmons, L., 1987. Comparative feeding ecology of felids in a neotropical rainforest. *Behav.*
182 *Ecol. Sociobiol.* 20, 271–283. <https://doi.org/10.1007/BF00292180>

183 Fernandez, E.C., 2002. Ocelot (*Leopardus pardalis*) ecology in the Chamela-Cuixmala Biosphere
184 Reserve, Jalisco, Mexico (M.S.). University of Wyoming, United States -- Wyoming.

185 Gilmore, M., 2005. An ethnoecological and ethnobotanical study of the Maijuna Indians of the
186 Peruvian Amazon (PhD Thesis). Miami University.

187 Gilmore, M.P., Young, J.C., 2010. The Maijuna participatory mapping project: mapping the past
188 and the present for the future. *Perú Maijuna Rapid Biol. Soc. Invent. Rep.* 22, 233.

189 Hamilton, W.D., 1971. Geometry for the selfish herd. *J. Theor. Biol.* 31, 295–311.

190 Izawa, K., 1993. Soil-eating by *Alouatta* and *Ateles*. *Int. J. Primatol.* 14, 229–242.

191 Kreulen, D.A., 1985. Lick use by large herbivores: a review of benefits and banes of soil
192 consumption. *Mammal Rev.* 15, 107–123. [https://doi.org/10.1111/j.1365-](https://doi.org/10.1111/j.1365-2907.1985.tb00391.x)
193 [2907.1985.tb00391.x](https://doi.org/10.1111/j.1365-2907.1985.tb00391.x)

194 Laundré, J.W., Hernández, L., Ripple, W.J., 2010. The landscape of fear: ecological implications
195 of being afraid. *Open Ecol. J.* 3, 1–7.

196 Link, A., De Luna, A.G., Arango, R., Diaz, M.C., 2011. Geophagy in brown spider monkeys
197 (Ateles hybridus) in a lowland tropical rainforest in Colombia. *Folia Primatol. Basel* 82,
198 25–32. <http://dx.doi.org/10.1159/000326056>

199 Link, A., Fiore, A.D., 2013. Effects of predation risk on the grouping patterns of white-bellied
200 spider monkeys (*Ateles belzebuth belzebuth*) in Western Amazonia. *Am. J. Phys.*
201 *Anthropol.* 150, 579–590. <https://doi.org/10.1002/ajpa.22230>

202 Link, A., Galvis, N., Fleming, E., Di Fiore, A., 2011. Patterns of mineral lick visitation by spider
203 monkeys and howler monkeys in Amazonia: are licks perceived as risky areas? *Am. J. Primatol.*
204 73, 386–396.

205 Mares, M.A., Ojeda, R.A., 1982. Patterns of diversity and adaptation in South American
206 hystricognath rodents. *Mamm. Biol. S. Am.* 6, 393–432.

207 Matsuda, I., Izawa, K., 2008. Predation of wild spider monkeys at La Macarena, Colombia.
208 *Primates* 49, 65–68.

209 Moe, S.R., 1993. Mineral content and wildlife use of soil licks in southwestern Nepal. *Canadian*
210 *Journal of Zoology* 71, 933–936.

211 Molina, E., León, T., Armenteras, D., 2014. Characteristics of natural salt licks located in the
212 Colombian Amazon foothills. *Environ. Geochem. Health* 36, 117–129.
213 <https://doi.org/10.1007/s10653-013-9523-1>

214 Montenegro, O.L., 1998. The behavior of Lowland Tapir (*Tapirus terrestris*) at a natural mineral
215 lick in the Peruvian Amazon (Master's Thesis). University of Florida.

216 Montenegro, O.L., 2004. Natural licks as keystone resources for wildlife and people in
217 Amazonia (Ph.D.). University of Florida, United States -- Florida.

218 Moreno, R.S., Kays, R.W., Samudio, R., 2006. Competitive release in diets of ocelot (*Leopardus*
219 *pardalis*) and puma (*Puma concolor*) after jaguar (*Panthera onca*) decline. *J. Mammal.* 87,
220 808–816.

221 Ospina, A.L., 2011. Social and ecological determinants of “fission-fusion” sociality and
222 grouping strategies in the white bellied spider monkey (*Ateles belzebuth belzebuth*) in a
223 lowland rainforest in Western Amazonia (Ph.D.). New York University, United States --
224 New York.

225 Roberts, M., Brand, S., Maliniak, E., 1985. The biology of captive prehensile-tailed porcupines,
226 *Coendou prehensilis*. *J. Mammal.* 66, 476–482. <https://doi.org/10.2307/1380922>

227 Tobler, M.W., 2008. The ecology of the lowland tapir in Madre de Dios, Peru: using new
228 technologies to study large rainforest mammals. Texas A&M University.

229 Tobler, M.W., Carrillo-Percastegui, S.E., Powell, G., 2009. Habitat use, activity patterns and use
230 of mineral licks by five species of ungulate in south-eastern Peru. *J. Trop. Ecol.* 25, 261–
231 270.

232

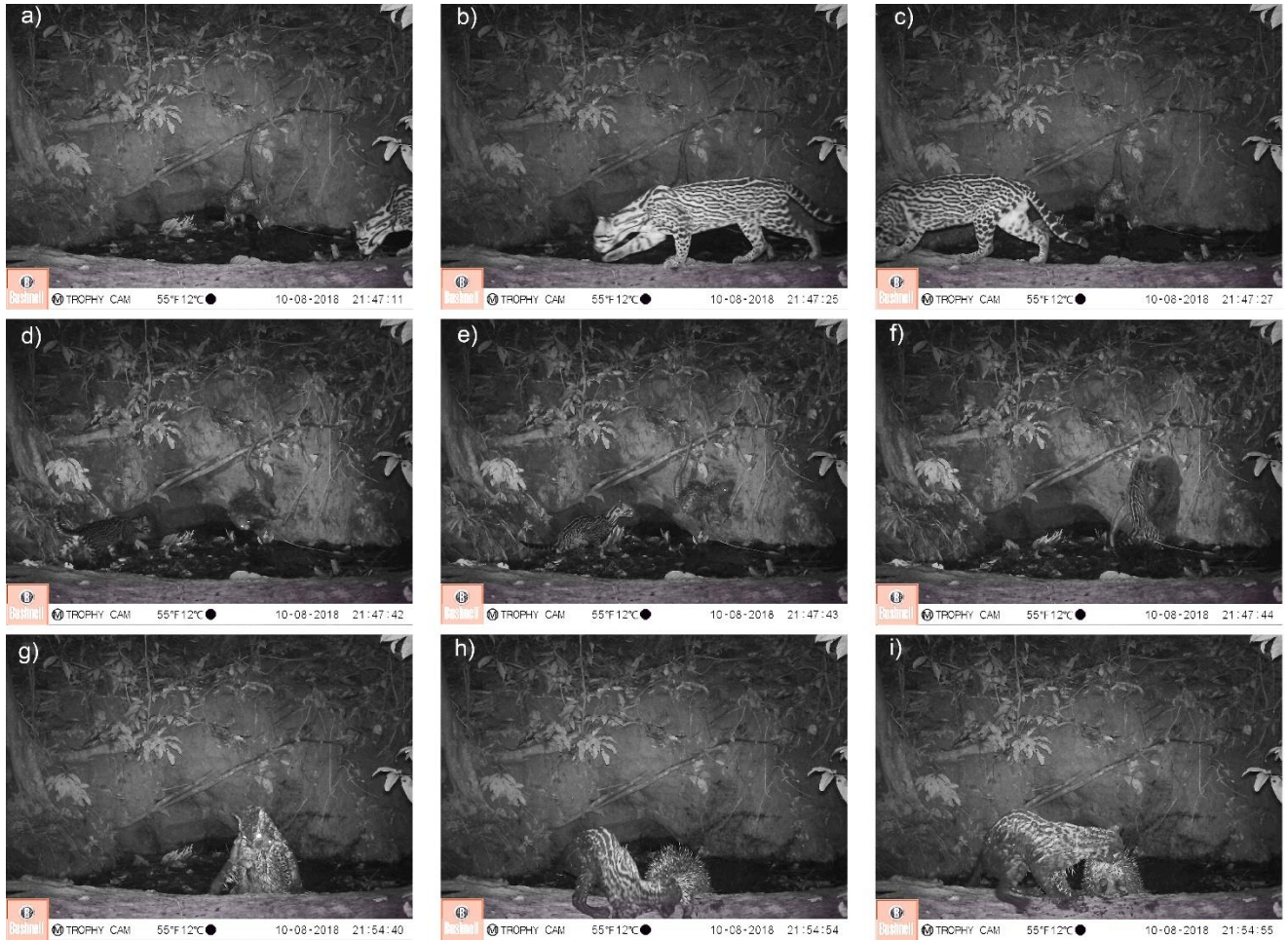
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239 **Fig. 1 a-b)** Adult porcupine (*Coendou prehensilis*) can be seen drinking water at the lick while a
 240 male ocelot (*Leopardus pardalis*) looks on; **c-d)** Ocelot walks along a log to approach the porcupine
 241 from the left; **e-f)** Ocelot pounces on the back of the porcupine, biting behind the head; **g-i)** Ocelot
 242 drags porcupine onto log and twists its neck

243