Lust for Salt in the Western Amazon

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ABSTRACT

Although the use of mineral licks by diverse Amazonian birds and mammals is well-known, the ultimate motivation for such behavior remains unclear. As aerosol deposition of salts declines with distance from oceanic sources, lick visitation in the western Amazon can be explained by demand for sodium, given the low concentration of this micronutrient in the plant tissues consumed by these taxa. Sodium limitation also influences ant foraging behavior, and impinges on ecosystem rates of carbon cycling. The biogeographical context of sodium availability has been largely overlooked, but has substantial pantropical implications for herbivore and decomposer performance in inland rain forests.

Abstract in Spanish is available in the online version of this article.

Key words: ant; decomposition; geophagy; herbivory; licks; litter; sodium; vertebrate.

ECOLOGICAL RESEARCH IN THE TROPICS OFTEN INVOLVES STUDIES OF HERBIVORY. We suggest that an important feature of this plantanimal interaction, namely salt limitation, has been neglected by such studies, and yet has important implications for the physiological condition of herbivores, as well as for litter decomposition and carbon cycling more generally within inland tropical forests. Animals need dietary salt to maintain homeostasis, and exhibit corresponding behaviors to offset metabolic loss of sodium, chloride, and potassium ions (Denton 1982). This phenomenon is particularly pronounced in herbivores, for which dietary ingestion of intrinsically low-sodium plant tissue is often insufficient to meet nutritional requirements (National Research Council 2005). In consequence, taxa as diverse as African elephants (Holdø et al. 2002), moose (Risenhoover & Peterson 1986), and mountain gorillas (Rothman et al. 2006) seek out dietary supplementation of sodium. The use of mineral licks worldwide is well-documented for ungulates and other herbivorous mammals, although the role of compounds supplemental to sodium cannot necessarily be excluded (Jones & Hanson 1985, Klaus & Schmid 1998). Here, we propose that the diverse vertebrate visitors to the well-known mineral licks of the western Amazon have a common goal of sodium supplementation in a region deprived geographically of salt. Moreover, the outcome of sodium deprivation extends to the phytophagous arthropod community, including key taxa in the decomposition ecology of leaf litter.

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SALT-SEEKING IN THE AMAZON BASIN

Mineral licks of the western Amazon represent premier examples of geophagy; together with mineral-enhanced seepages and pools to which animals are attracted, these sites are known in Spanish as collpas, saladeros, and salados. Known sites on riverbanks to which parrots flock to ingest soil are prominent tourist destinations, and exceed fifty in South America (Brightsmith & Muñoz-Najar 2004, Lee et al. 2009). Less appreciated but equally relevant are comparable behaviors at clay and mineral licks reported for Amazonian primates (Ferrari et al. 2008, Blake et al. 2010), phyllostomid bats (Bravo et al. 2008), and various other mammals (see Izawa 1975, Brightsmith et al. 2008). Multiple hypotheses not mutually exclusive to sodium seeking have been proposed to explain these behaviors, including the use of clays to facilitate alkaloid detoxification, and medicinal functions of ingested soil (Table 1). Correlational analyses between salt content at licks and attraction behavior cannot rule out such alternative motivating factors. For the iconic geophagy exhibited by Amazonian parrots, however, recent studies have concluded that sodium acquisition is the primary motivating factor. Clay licks occur primarily in the western Amazon, with limited occurrence in the central and northern regions of the basin, but never near the Atlantic shoreline (Lee et al. 2009). Analysis of soil content at licks confirms elevated sodium levels (and in some cases magnesium levels) relative to control sites (Brightsmith et al. 2008, Powell et al. 2009). Geophagy by other sympatric avian taxa, such as columbids and cracids (Brightsmith & Muñoz-Najar 2004), is similarly consistent with sodium seeking associated with obligately phytophagous diets.

TABLE 1. Non-mutually exclusive hypotheses for mineral lick visitation and geophagy by vertebrates (see also reviews by Kreulen 1985 and Klaus & Schmid 1998).

Hypothesis	References
Detoxification of plant secondary	Oates (1978), Gilardi
compounds, especially alkaloids	et al. (1999)
2. Mineral supplementation of diet, especially	Jones and Hanson (1985),
with sodium and calcium ions	Powell et al. (2009)
3. Acquisition of soil to supplement	Gionfrido and Best
mechanical grinding in the avian crop	(1996)
4. Zoopharmacognosy, with particular	Knezevich (1998), Vermeer
reference to internal parasites and	and Ferrell (1985)
alleviation of diarrhea	
5. Buffering of gastric pH	Oates (1978), Mahaney
	et al. (1999)

Extensive work on bat attraction to salt licks in Amazonian Peru has also implicated sodium deprivation as a motivating factor. Pools and depressions in soil, some of which are created by terrestrial geophagous taxa, are visited by no fewer than 26 species of phyllostomid bats (Bravo et al. 2008, 2010b). Water from the bat-visited collpas is enhanced in sodium, and (less so) in calcium and potassium; bats prefer to drink water from the lick sites when matched with controls (Bravo et al. 2010a). Using nitrogen isotope ratios, Voigt et al. (2008) showed that short-tailed fruit bats (Carollia perspicillata, Phyllostomidae) with plant-dominated diets were more likely to be captured at salt licks in Ecuador, and concluded that calcium seeking and buffering of plant secondary compounds were likely explanations for this behavior. However, given that plant tissues typically contain orders of magnitude less sodium compared with animals, this finding also supports a conclusion of sodium deprivation among more phytophagous individuals.

Visitors to mineral licks in the Amazon can also be found among some primates and arthropods. Geophagy in New World monkeys is well-documented, but its restricted occurrence in Amazonian as opposed to Central American forests has not been well-understood (Izawa 1993, Campbell et al. 2005, Ferrari et al. 2008). We suggest that this behavior in frugivorous primates is driven by a search for sodium, and is unlikely to occur in most of lowland Central America or near the South American littoral, given their proximity to oceanic sources. Moreover, ants in the western Amazon favor low-concentration sodium solution relative to either carbohydrate or protein baits, and the intensity of this response varies inversely with distance from oceanic sources at continental scales (Kaspari et al. 2008). Inland attraction to sodium is, however, substantially diminished for ant taxa that are predominantly carnivorous, as would be predicted given their animal prey (Kaspari et al. 2008). Insects from at least five other orders have also been observed engaging in geophagy at mineral licks in Peru (Brightsmith et al. 2008, Molleman 2010). These behaviors are not confined to the tropics; the attraction of some temperate-zone swallowtail butterflies to various sodium salts is well-known (Arms et al. 1974).

THE GEOGRAPHY OF SALT

In aggregate, the evidence supporting vertebrate attraction to salt in the Amazon basin is both compelling and geographically constrained. The vast majority of mineral licks utilized by parrots and other vertebrates are in the western part of the basin thousands of kilometers from the Atlantic Ocean. Directly to the west of these sites, the Andean range has precluded influx of marine aerosols from the Pacific since the late Miocene (Graham 2009). Sodium concentrations in Amazonian river waters are accordingly highest at the Atlantic, and decline monotonically with increasing distance upriver (Stallard & Edmond 1981, Tardy et al. 2005). This geographical gradient suggests that the western Amazon is salt-deprived, and that an increasing physiological demand for dietary salt supplementation is correspondingly manifested by herbivores at mineral licks. Carnivorous and omnivorous taxa, by contrast, are less likely to be sodiumstressed, and one commonality of all documented vertebrates that visit Amazonian salt licks and collpas is a predominantly phytophagous diet.

Altitudinal variation in salt availability may similarly be relevant to animal nutritional ecology, given greater leaching by precipitation at higher elevations. Differential responses of herbivores from lowland and alpine regions may correspondingly ensue (Blair-West et al. 1968). The eastern slopes of the Andes should be particularly susceptible to sodium deprivation, to which end an elevational transect of environmental availability would be informative. Water at salt licks consumed by the mountain tapir in Colombia is enhanced in both nitrogen and sodium relative to a control stream (Lizcano & Cavelier 2004), but no further information is available on such behaviors in Andean contexts.

ECOLOGICAL IMPLICATIONS

As behavioral attraction to salt represents a temporary but intentional switch from macronutrient to micronutrient consumption, a number of ecological consequences may ensue. Metabolic costs of movement to mineral licks, together with an opportunity cost in time otherwise spent foraging, indicate net energetic losses. Exposure to predators may ensue at licks, particularly for arboreal taxa that otherwise rarely venture to the ground (Link et al. 2010), whereas mixed-species groups visiting salt licks may benefit from an overall enhanced group alertness to potential predators. Although the adverse physiological consequences of extreme sodium deprivation are clear (Denton 1982), chronic effects of suboptimal levels may cascade into other aspects of herbivore ecology, including reproductive and population dynamics (Freeland et al. 1985, Batzli 1986). This outcome would pertain to tropical as well as extra-tropical sites that, for reasons of physical geography, are sodium-restricted (e.g., Aumann & Emlen 1965).

Additional contexts can be identified within tropical forests that provide for enhanced sodium. The activity of fungal decomposers concentrates sodium within leaf litter, and also within decomposing logs (Cromack et al. 1977). Chimpanzees in central Africa consume decaying palm pith that is high in sodium (Reynolds et al. 2009), and this behavior may be more widespread among primates in tropical forests (Kaplin & Moermond 2000, Pinto & Setz 2004, Rothman et al. 2006). Non-trivial consumption of decaying wood has also been noted for spider monkeys at various Amazonian sites (Suarez 2006, Di Fiore et al. 2008) and, as with geophagy, is generally confined to the western parts of the basin.

Indigenous Amazonian peoples would also be predicted to seek out dietary supplements of sodium. Kayapó Indians in central Brazil have been reported to burn leaves of three palm species to obtain salt-enhanced ash (Posey 2002), but the geographical distribution of such activity has not been studied.

At the broader ecosystem scale, sodium availability may also influence rates of litter decomposition and carbon cycling in inland tropical forests. Low-level addition of 0.5 percent sodium chloride solution to standardized leaf litter bags in the Peruvian Amazon resulted in a marked increase in termite numbers and in overall mass loss relative to bags treated with stream water controls (Kaspari et al. 2009). These effects occurred over a timespan of only several weeks, and were attributed to effects of sodium enhancement alone given results with different salt combinations. Sodium limitations on decomposition in tropical rainforests may in fact be widespread given that approximately 80 percent of such humid regions lie more than 100 km inland. In sodium-limited temperate-zone ecosystems, anthropogenically supplemented salts may also have the potential to alter insect behavior and population growth rates (Braun & Flückiger 1984, Kaspari et al. 2010). The consequences of salt supplementation in both tropical and temperate zones represent an intriguing target for experimental work at behavioral and ecological scales.

CONCLUSIONS AND FUTURE DIRECTIONS

Evidence to date suggests that visitation by diverse mammalian and avian taxa to mineral licks in the western Amazon can be at least partially explained by the geographically imposed outcome of sodium deprivation. Alternative hypotheses, and, in particular, detoxification of alkaloids and other toxins, are not necessarily generalizable to the broad diversity of plant secondary compounds consumed by animal taxa, and also do not explain restriction of these behaviors to only the western parts of the Amazon basin. The recently observed geographical gradient in ant behavioral responses to salt baits extends the concept of sodium deprivation to a key invertebrate group, and indicates a more general phenomenon in herbivorous taxa. The Amazon represents the largest tract of inland rain forest on the planet. Identification of sodium as a potentially constraining physiological and ecological factor for herbivores has important implications for animal population dynamics as well as broader features of carbon cycling within this vast region.

To move beyond correlational interpretations of animal foraging behavior at salt licks, additional evidence should be obtained to test the hypothesis of sodium deprivation. For example, demonstration of an inverse relationship between behavioral attraction to sodium and its excretory loss (Holdø et al. 2002), as well as differential responses to multiple sodiumand chloride-containing compounds (Blair-West et al. 1968, Arms et al. 1974), can be used to hone in on particular ions as primary attractants. In the tropical regions characterized by a dry season and reduced deposition of salt via rainfall, behavioral responses to sodium may correspondingly become more pronounced. Inland salt licks at tropical sites in the Old World (Klaus et al. 1998, Matsubayashi et al. 2007) would also be appropriate for characterization of mineral content relative to herbivore behavior and distance from oceanic sources. In contrast to various alternative hypotheses (see Table 1), we predict sodium deprivation to be the one common factor underlying these otherwise taxonomically and geographically disparate phenomena.

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LITERATURE CITED

- Arms, K., P. Feeny, and R. C. Lederhouse. 1974. Sodium: Stimulus for puddling behavior by tiger swallowtail butterflies, *Papilio glaucus*. Science 185: 372–374
- AUMANN, G. D., AND J. T. EMLEN. 1965. Relation of population density to sodium availability and sodium selection by microtine rodents. Nature 208: 198–199.
- BATZLI, G. O. 1986. Nutritional ecology of the California vole: Effects of food quality on reproduction. Ecology 67: 406–412.
- BLAIR-WEST, J. R., J. P. COGHLAN, D. A. DENTON, J. F. NELSON, E. ORCHARD, B. A. SCOGGINS, R. D. WRIGHT, K. MYERS, AND C. L. JUNQUEIRA. 1968. Physiological, morphological and behavioural adaptation to a sodium deficient environment by wild native Australian and introduced species of animals. Nature 217: 922–928.
- BLAKE, J. G., J. GUERRA, D. MOSQUERA, R. TORRES, B. A. LOISELLE, AND D. ROMO. 2010. Use of mineral licks by white-bellied spider monkeys (Ateles belzebuth) and red howler monkeys (Alouatta seniculus) in eastern Ecuador. Int. J. Primatol. 31: 471–483.
- Braun, S., and W. Flückiger. 1984. Increased population of the aphid Aphis pomi at a motorway. Part 2: The effect of drought and deicing salt. Environ. Pollut. Series A. 36: 261–270.
- Bravo, A., K. E. Harms, and L. H. Emmons. 2010a. Preference for *Collpa* water by frugivorous bats (*Artibeus*): An experimental approach. Biotropica 42: 276–280.
- Bravo, A., K. E. Harms, and L. H. Emmons. 2010b. Puddles created by geophagous mammals are potential mineral sources for frugivorous bats (Stenodermatinae) in the Peruvian Amazon. J. Trop. Ecol. 26: 173–184.
- Bravo, A., K. E. Harms, R. D. Stevens, and L. H. Emmons. 2008. Collpas: Activity hotspots for frugivorous bats (Phyllostomidae) in the Peruvian Amazon. Biotropica 40: 203–210.
- Brightsmith, D. J., and R. A. Muñoz-Najar. 2004. Avian geophagy and soil characteristics in southeastern Peru. Biotropica 36: 534–543.
- BRIGHTSMITH, D. J., J. TAYLOR, AND T. D. PHILLIPS. 2008. The roles of soil characteristics and toxin adsorption in avian geophagy. Biotropica 40: 766–774.

- CAMPBELL, C. J., F. AURELIA, C. A. CHAPMAN, G. RAMOS-FERNANDEZ, K. Matthews, S. E. Russo, S. Suarez, and L. Vick. 2005. Terrestrial behavior of Ateles spp. Int. J. Primatol. 26: 1039-1051.
- CROMACK, K., P. SOLLINS, R. L. TODD, D. A. CROSSLEY, W. M. FENDER, R. FOGEL, AND A. W. TODD. 1977. Soil microorganism-arthropod interactions: fungi as major calcium and sodium sources. In W. J. Mattson (Ed.). The role of arthropods in forest ecosystems, pp. 78-84. Springer-Verlag, New York.
- DENTON, D. A. 1982. The hunger for salt. Springer-Verlag, Berlin.
- DI FIORE, A., A. LINK, AND J. L. DEW. 2008. Diets of wild spider monkeys. In C. J. Campbell (Ed.). Spider monkeys: The biology, behavior and ecology of the genus Ateles, pp. 81-137. Cambridge University Press, Cambridge.
- FERRARI, S. F., L. M. VEIGA, AND B. URBANI. 2008. Geophagy in New World monkeys (Platyrrhini): Ecological and geographic patterns. Folia Primatol. 79: 402-415.
- FREELAND, W. J., P. H. CALCOTT, AND D. P. GEISS. 1985. Allelochemicals, minerals and herbivore population size. Biochem. Syst. Ecol. 13: 195-206.
- GILARDI, J. D., S. S. DUFFEY, C. A. MUNN, AND L. A. TELL. 1999. Biochemical functions of geophagy in parrots: Detoxification of dietary toxins and cytoprotective effects. J. Chem. Ecol. 25: 897-922.
- GIONFRIDO, J. P., AND L. B. BEST. 1996. Grit-use patterns in North American birds: The influence of diet, body size, and gender. Wilson Bull. 108: 686-696.
- Graham, A. 2009. The Andes: A geological overview from a biological perspective. Ann. Missouri Bot. Gard. 96: 371-385.
- HOLDØ, R. M., J. P. DUDLEY, AND L. R. McDowell. 2002. Geophagy in the African elephant in relation to availability of dietary sodium. J. Mammal. 83: 652-664.
- IZAWA, K. 1975. Foods and feeding behavior of monkeys in the Upper Amazon basin. Primates 16: 295-316.
- IZAWA, K. 1993. Soil-eating by Alouatta and Ateles. Int. J. Primatol. 14: 229-242.
- JONES, R. L., AND H. C. HANSON. 1985. Mineral licks, geophagy, and biogeochemistry of North American ungulates. Iowa State University Press, Ames, Iowa,
- KAPLIN, B. A., AND T. C. MOERMOND. 2000. Foraging ecology of the mountain monkey (Cercopithecus l'hoesti): Implications for its evolutionary history and use of disturbed forest. Am. J. Primatol. 50: 227-246.
- KASPARI, M., C. CHANG, AND J. WEAVER. 2010. Salted roads and sodium limitation in a northern forest community. Ecol. Ent. 35: 543-548.
- KASPARI, M., S. P. YANOVIAK, AND R. DUDLEY. 2008. On the biogeography of salt limitation: A study of ant communities. Proc. Natl Acad. Sci. U.S.A. 105: 17848-17851.
- Kaspari, M., S. P. Yanoviak, R. Dudley, M. Yuan, and N. A. Clay. 2009. Sodium shortage as a constraint on the carbon cycle in an inland tropical rainforest. Proc. Natl. Acad. Sci. U.S.A. 106: 19405-19409.
- KLAUS, G., C. KLAUS-HÜGI, AND B. SCHMID. 1998. Geophagy by large mammals at natural licks in the rain forest of the Dzanga National Park, Central African Republic. J. Trop. Ecol. 14: 829-839.
- KLAUS, G., AND B. SCHMID. 1998. Geophagy at natural licks and mammal ecology: A review. Mammalia 62: 481-497.
- KNEZEVICH, M. 1998. Geophagy as a therapeutic mediator of endoparasitism in a free-ranging group of rhesus macaques (Macaca mulatta). Am. J. Primatol. 44: 71-82.
- Kreulen, D. A. 1985. Lick use by large herbivores: A review of benefits and banes of soil consumption. Mammal. Rev. 15: 107-123.

- LEE, A. T. K., S. KUMAR, D. J. BRIGHTSMITH, AND S. J. MARSDEN. 2009. Parrot claylick distribution in South America: Do patterns of "where" help answer the question "why"? Ecography 33: 503-513.
- Link, A., N. Galvis, E. Fleming, and A. Di Fiore. 2010. Patterns of mineral lick visitation by spider monkeys and howler monkeys in Amazonia: Are licks perceived as risky areas? Am. J. Primatol. 71: 1-11.
- LIZCANO, D. J., AND J. CAVELIER. 2004. Chemical characteristics of salt licks and feeding habits of mountain tapir (Tapirus pinchaque) in the central Andes of Colombia. Mastozool. Neotrop. 11: 193-201.
- MAHANEY, W. C., M. W. MILLNER, K. SANMUGADAS, R. G. V. HANCOCK, S. Aufreiter, S. Campbell, M. A. Huffman, M. Wink, D. Malloch, AND V. KALM. 1999. Chemistry, mineralogy and microbiology of termite mound soil eaten by the chimpanzees of the Mahale Mountains, western Tanzania. J. Trop. Ecol. 15: 565-588.
- Matsubayashi, H., P. Lagan, N. Majalap, J. Tangah, J. R. A. Sukor, and K. KITAYAMA. 2007. Importance of natural licks for the mammals in Bornean inland tropical rainforests. Ecol. Res. 22: 742-748.
- MOLLEMAN, F. 2010. Puddling: From natural history to understanding how it affects fitness. Entomol. Exp. Appl. 134: 107-113.
- National Research Council. 2005. Mineral tolerance of animals. National Academies Press, Washington, DC.
- OATES, J. F. 1978. Water-plant and soil consumption by guereza monkeys Colobus guereza: A relationship with minerals and toxins in the diet? Biotropica 10: 241-253.
- PINTO, L. P., AND E. Z. F. SETZ. 2004. Diet of Alouatta belzebul discolor in an Amazonian Rain Forest of northern Mato Grosso State, Brazil. Int. J. Primatol. 25: 1197-1211.
- Posey, D. A. 2002. Kayapó ethnoecology and culture. Routledge, London.
- POWELL, L. L., T. U. POWELL, G. V. N. POWELL, AND D. J. BRIGHTSMITH. 2009. Parrots take it with a grain of salt: Available sodium content may drive collpa (clay lick) selection in southeastern Peru. Biotropica 41: 279-
- REYNOLDS, V., A. W. LLOYD, F. BABWETEERA, AND C. J. ENGLISH. 2009. Decaying Raphia farinifera palm trees provide a source of sodium for wild chimpanzees in the Budongo Forest, Uganda. PLoS ONE 4:
- RISENHOOVER, K. L., AND R. O. PETERSON. 1986. Mineral licks as a sodium source for Isle Royale moose. Oecologia 71: 121-126.
- ROTHMAN, J. M., P. J. VAN SOEST, AND A. N. PELL. 2006. Decaying wood is a sodium source for mountain gorillas. Biol. Lett. 2: 321-324.
- STALLARD, R. F., AND J. M. EDMOND. 1981. Geochemistry of the Amazon. 1. Precipitation chemistry and the marine contribution to the dissolved load at the time of peak discharge. J. Geophys. Res. 86: 9844-
- SUAREZ, S. A. 2006. Diet and travel costs for spider monkeys in a nonseasonal, hyperdiverse environment. Int. J. Primatol. 27: 411-431.
- Tardy, Y., V. Bustillo, C. Roquin, J. Mortatti, and R. Victoria. 2005. The Amazon. Bio- geochemistry applied to river basin management. Part I. Hydro-climatology, hydrograph separation, mass transfer balances, stable isotopes, and modelling. Appl. Geochem. 20: 1746-
- VERMEER, D. E., AND R. R. J. FERRELL. 1985. Nigerian geophagical clay: A traditional antidiarrheal pharmaceutical. Science 227: 634-636.
- VOIGT, C. C., K. A. CAPPS, D. K. N. DECHMANN, R. H. MICHENER, AND T. H. Kunz. 2008. Nutrition or detoxification: Why bats visit mineral licks of the Amazonian rainforest. PLoS ONE 3: e2011.