

# On the biogeography of salt limitation: A study of ant communities

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Edited by James H. Brown, University of New Mexico, Albuquerque, NM, and approved September 25, 2008 (received for review May 9, 2008)

**Sodium is an essential nutrient whose deposition in rainfall decreases with distance inland. The herbivores and microbial decomposers that feed on sodium-poor vegetation should be particularly constrained along gradients of decreasing sodium. We studied the use of sucrose and NaCl baits in 17 New World ant communities located 4–2757 km inland. Sodium use was higher in genera and subfamilies characterized as omnivores/herbivores compared with those classified as carnivores and was lower in communities embedded in forest litter than in those embedded in abundant vegetation. Sodium use was increased in ant communities further inland, as was preference for the baits with the highest sodium concentration. Sucrose use, a measure of ant activity, peaked in communities 10–100 km inland. We suggest that the geography of ant activity is shaped by sodium toxicity near the shore and by sodium deficit farther inland. Given the importance of ants in terrestrial ecosystems, changing patterns of rainfall with global change may ramify through inland food webs.**

ants | biogeochemistry | geography | limitation | sodium

There are many reasons why the availability of sodium should impact population and ecosystem processes. Sodium is vital for maintaining osmotic balance, muscle activity, and nervous system function (1), and body sodium is tightly regulated (2). Terrestrial plants contain *ca.* 1.0 mg · kg<sup>-1</sup> sodium (3), whereas the herbivores and decomposer microbes that eat them maintain sodium levels 100- to 1000-fold higher (2, 4). Because sodium (unlike, e.g., nitrogen) is readily lost through excretion, basal consumers often must expend considerable energy to find and harvest it (5–7).

Sodium supply varies geographically. It is carried inland by oceanic aerosols, and the sodium content in rainfall can drop up to 1000-fold with increasing distance from the ocean (8, 9). Furthermore, inland ecosystems with high rainfall may experience significant sodium loss through leaching (10). As a consequence, continental populations of moose (6), gorillas (11), parrots (12), butterflies (13), and bees (14) often search for “salt licks” (15) or resort to cannibalism (16) to maintain sodium balance. However, aside from the salient work of Blair-West *et al.* (17) on introduced rabbits in Australia, we know of no detailed studies of how attraction to sodium, and potentially sodium limitation, varies geographically.

Ant communities (Hymenoptera: Formicidae) are a model system for exploring the biogeography of salt limitation. Ants are common players in terrestrial ecosystems (18, 19). Moreover, ants are trophically diverse, with herbivores that feed on sugary exudates common in “green” food webs with abundant vegetation (e.g., forest canopies and grasslands) and predators that feed on the consumers of decomposer microbes, common in litter (or “brown”) food webs (20–23).

We quantified recruitment to NaCl and sucrose in 17 ant communities to test 2 hypotheses concerning the biogeography of salt limitation. The first hypothesis posits that decreased sodium inputs via rainfall should increase the sodium limitation of consumers. We predicted that the harvesting of NaCl solution

by ants increases from the coastline to the interior of continents. The second hypothesis posits that herbivores are more sodium-limited than predators. We predicted that carnivorous taxa would use NaCl baits less frequently than herbivores and omnivores, and that communities embedded in green food webs would show more attraction to sodium than those from brown food webs.

## Results

We sampled ant communities from a variety of ecosystems [supporting information (SI) Table S1], ranging from coastal forest and scrub 4 km inland to an Amazonian rain forest 2757 km inland. These communities yielded between 4 (Panama litter) and 18 genera (Peru rain forest litter; median, 8) and were sampled at temperatures from 26 °C (Peru rainforest) to 34 °C (Florida scrub; median, 28 °C).

Ant recruitment to sucrose increased from concentrations of 0.5% to 20% ( $F_{1,66} = 83.14$ ;  $P < 0.0001$ ), with 20% sucrose (Fig. 1) uniformly trending the highest. Ant recruitment to sucrose also varied with distance inland ( $F_{3,66} = 4.9$ ;  $P < 0.004$ ). A significant distance–concentration interaction ( $F_{3,66} = 6.8$ ;  $P < 0.0005$ ) indicated that the exponent of the recruitment curve varied with distance to the ocean. When communities in each distance bin were analyzed separately, the percentage of vial hits scaled as sucrose concentration<sup>0.15</sup> at < 10 km inland ( $F_{1,8} =$ ;  $P = 0.18$ ;  $r^2 = 0.21$ ) up to 0.88 at 10–100 km inland ( $F_{1,17} = 5.7$ ;  $P = 0.0001$ ;  $r^2 = 0.65$ ), and down to 0.50 ( $F_{1,29} = 7.8$ ;  $P = 0.0001$ ;  $r^2 = 0.68$ ) and 0.48 ( $F_{1,12} = 9.71$ ;  $P = 0.0001$ ;  $r^2 = 0.89$ ), respectively, at 100–1000 and > 1000 km inland. The highest sucrose recruitment values (66% and 68%) were 10–100 km at concentrations of 10% and 20%, respectively. Further inland, these concentrations averaged 49% and 55%, and 54% and 61% hits (Fig. 1).

Like sucrose, ant recruitment to NaCl baits showed significant interactions ( $F_{3,71} = 4.4$ ;  $P = 0.007$ ), along with the effects of concentration ( $F_{1,71} = 7.06$ ;  $P = 0.0097$ ) and distance inland ( $F_{3,71} = 10.2$ ;  $P = 0.0001$ ; Fig. 1). Within 10 km of the ocean, vial hits marginally decreased as NaCl concentration<sup>-0.27</sup> ( $F_{1,8} = -2.02$ ;  $P = 0.08$ ;  $r^2 = 0.34$ ), failed to vary with concentration at 10–100 km ( $F_{1,18} = 1.4$ ;  $P = 0.17$ ;  $r^2 = 0.10$ ), increased with NaCl concentration<sup>0.33</sup> at 100–1000 km ( $F_{1,33} = 3.7$ ;  $P = 0.007$ ;  $r^2 = 0.30$ ), and increased most strongly with NaCl concentration<sup>0.44</sup> at > 1000 km inland ( $F_{1,12} = 3.3$ ;  $P = 0.0024$ ;  $r^2 = 0.55$ ). Unlike sucrose, the highest values for recruitment to NaCl were at the sites furthest inland.

Author contributions: M.K., S.P.Y., and R.D. designed research; M.K., S.P.Y., and R.D. performed research; M.K. analyzed data; and M.K. wrote the paper.

The authors declare no conflict of interest.

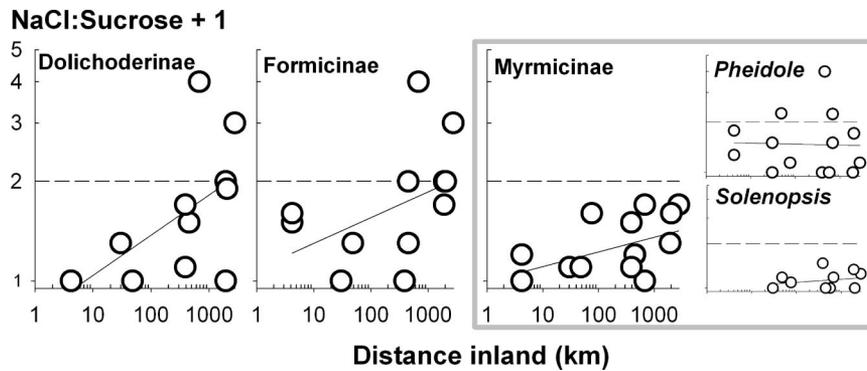
This article is a PNAS Direct Submission.

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This article contains supporting information online at [www.pnas.org/cgi/content/full/0804528105/DCSupplemental](http://www.pnas.org/cgi/content/full/0804528105/DCSupplemental).

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**Fig. 3.** Taxonomic decomposition of salt use across 17 ant communities. The ratio of hits on NaCl baits and sucrose baits is compared among three common subfamilies and two common genera of the subfamily Myrmicinae. The dashed line represents equal use; the straight curve is the best fit by linear regression.

and ability to harvest NaCl. If this were so, then we would expect to see self-similarity in patterns of ant recruitment to sodium as we decompose patterns for the family Formicidae into its constituent subfamilies, genera, and species. Within the limits of reasonable statistical power (10 sites with at least five hits per site), we show that three common subfamilies consistently increase NaCl use relative to sucrose further inland, but that two genera do not. Both genera, *Pheidole* and *Solenopsis*, may be buffered from NaCl deprivation by being carnivorous (16). Consistent with our second prediction, carnivorous ant genera and subfamilies are less likely to recruit to salt than those with lower  $\delta^{15}\text{N}$  signatures (24). Sodium stress may also be tempered by reducing body sodium (17), increasing the efficiency of osmoregulation (30, 31), switching sodium with phosphate or potassium (2), or feeding at higher trophic levels (16). Ultimately, the increasingly triangular scatterplots at finer taxonomic resolution may represent individual taxa, populations, or individuals implementing a variety of tactics to ameliorate sodium stress (32, 33). The degree to which our community-wide pattern is shaped by taxonomic replacement by more or less sodium-hungry taxa, by local adaptation, and/or by phenotypic plasticity awaits further study.

Sodium availability also varies locally with soil type, topography, and irrigation by groundwater (34). Some variability in sodium and sucrose recruitment across sites is accounted for by

the local changes in habitat; for omnivorous taxa, access to sodium (and various other micronutrients) is likely higher in brown food webs, where fungi and bacteria concentrate sodium from sodium-poor leaf litter (4, 35). Our results support the hypothesis that ants in habitats surrounded by green vegetation are more herbivorous and sodium-stressed; however, the use of sodium baits even in litter communities appears to increase rapidly beyond *ca.* 300 km inland (Fig. 2).

In conclusion, the availability of NaCl appears to generate a geographic signature on the activity of a dominant group of terrestrial consumers, the ants. Sodium has a clear importance as a limiting nutrient in animal husbandry (2, 7). It also is toxic to plants; one-third of irrigated land is less productive due to NaCl contamination (2). The biogeography of salt limitation also may play out at the ecosystem level (29); for example, if low sodium input has little effect on plants but hampers the activity of microbial decomposers (4), then ecosystems of the continental interior (e.g., much of the Amazon basin, the grasslands of North America) may store more plant production as undecomposed carbon compared with ecosystems closer (but not necessarily adjacent) to the ocean. Given the importance of oceanic aerosols in sodium deposition and the uncertainty in predicting rainfall in a changing climate (36), this may be another unexpected forcing factor on global carbon cycles.

**Table 1.** Seventeen ant communities sampled for sucrose and NaCl use

Region	Name	Genera	Biome	Distance inland, km	Number	Latitude	Longitude	Food Web	Temperature, °C
Peru	Cicra	18	Rain forest	2757	6	12.56 S	70.10 W	Brown	26.7
Peru	Iquitos	15	Rain forest	2028	5	3.95 S	73.41 W	Brown	25.5
Peru <sup>1</sup>	ACTS litter	12	Rain forest	1920	5	3.25 S	72.91 W	Brown	26.7
Peru <sup>1</sup>	ACTS canopy	4	Rain forest	1920	–	3.25 S	72.91 W	Green	–
Oklahoma <sup>2</sup>	Ranch	8	Grassland	676	4	35.23 N	97.19 W	Green	25.6
Oklahoma <sup>2</sup>	Lake	6	Oak forest	676	4	35.20 N	97.13 W	Brown	27.3
Arkansas <sup>3</sup>	Hayfield	8	Old field	448	5	33.81 N	94.20 W	Green	30.3
Arkansas <sup>3</sup>	Cut	8	Mixed forest	452	5	33.81 N	91.28 W	Brown	27.0
Arkansas	Oak	5	Oak forest	567	5	34.82 N	92.49 W	Brown	27.5
Arizona <sup>4</sup>	Stateline	9	Warm desert	389	4	31.95 N	109.16 W	Green	28.5
Arizona <sup>4</sup>	SWRS	6	Pine forest	389	4	31.88 N	109.22 W	Brown	29.2
Costa Rica	La Selva	9	Rain forest	76	5	10.43 N	84.01 W	Brown	27.5
Florida	Everglades	10	Grassland	48	5	25.38 N	80.61 W	Green	28.2
Panama <sup>5</sup>	BCI	4	Rain forest	30	5	9.15 N	79.86 W	Brown	25.7
Panama <sup>5</sup>	BCI	9	Clearing	30	1	9.15 N	79.86 W	Green	32.5
Florida <sup>6</sup>	Scrub	11	Pine scrub	4	4	27.59 N	80.37 W	Green	34.1
Florida <sup>6</sup>	Hammock	5	Oak forest	4	4	27.59 N	80.37 W	Brown	32

Number refers to the number of sampling transects run. Temperature refers to the mean temperature at start of trials. Sites sharing the same superscript are green and brown food webs paired by region. Temperature was not recorded in canopy studies, where 15 trees were sampled during mid-morning over 2 days.

## Methods

We sampled the NaCl and sucrose use of 17 ant communities (Table S1). For each site, we measured the distance from the oceanic source of precipitation using Google Earth. Where this was ambiguous (i.e., the Everglades are in the center of the Florida peninsula), or where two oceans generated rainfall (i.e., the Chihuahuan desert of Arizona), multiple measures were taken and averaged. In six localities (Arizona, Arkansas, Florida, Oklahoma, Panama, Peru; Table 1), we sampled pairs of communities at similar ( $\pm 10$  km) distances inland. Each pair was characterized *a priori* as a brown food web (i.e., closed canopy forests with a thick litter layer and little vegetation in the understory) or a green food web (i.e., minimal canopy, if any, and abundant living vegetation accessible to the ground ant community). All brown communities were in forest litter. Green communities included grasslands, hayfields, lawns, and, in one case, the canopy of a tropical forest (see below).

Ant communities were sampled with transects of labeled Eppendorf vials. Each vial was half-stuffed with cotton saturated with sucrose or NaCl solution. Sucrose was used as a standard of overall ant activity, because sugars are widely used by ants and there was no *a priori* reason to expect geographic variation in sucrose use. Vials were baited with a series of four or five concentrations (sucrose: 0.1%, 0.5%, 1%, 5%, 10%, and 20% by mass; NaCl: 0.01%, 0.05%, 0.1%, 0.5%, and 1% by mass) plus a distilled water treatment. Fifteen vials of each concentration were snapped shut and thoroughly mixed in plastic bags. At each site, four or five linear transects were laid out (with one exception—"Panama clearing" had only one transect), and a vial was randomly selected, opened, and placed every 1 m. After 1 h, we collected the vials, snapping the cap shut and capturing the ants using the baits. These ants were identified to genus and counted in the laboratory.

In addition, at one site (ACTS in Peru), we used a canopy walkway to sample the ant communities of 15 tree crowns and the leaf litter communities beneath. Because vials could not be reliably placed and recovered from tree branches, we placed cotton balls soaked in sucrose, NaCl, and distilled water solutions in a row (with baits separated by 10 cm) on a large branch in the canopy or on the leaf litter. Solutions started at 0.1% NaCl and 1% sucrose and were observed for 15 min. Over successive 5-min intervals, concentrations were increased (NaCl: 0.05%, 0.1%, and 0.5%, sucrose: 5%, 10%, and 20%). We recorded the number of ants that drank from the solutions (assayed as 5 sec of continuous contact with the mandibles) over the period in which each concentration was presented. A total of 15 trees and 19 litter patches (10 m apart) were sampled in this way.

For each site analyzed with the vial method, sucrose and NaCl vials were scored, at each concentration, by the percentage (out of 15 vials) with ants minus the activity at distilled water. The cotton ball experiments were scored

similarly as the percentage of trials (15 tree, 19 litter) in which sucrose and/or NaCl was collected by ants. The ACTS litter sampled with the Eppendorf vials and cotton balls yielded similar values; only vial data are presented here.

To analyze patterns of recruitment with concentration and distance inland, we first calculated the average percentage of hits by ants for each concentration at a given site across each site's replicate transects. We used one-tailed tests to evaluate predictions of (1) higher sodium use further inland and (2) lower sodium use with increasing carnivory.

We tested prediction 1 in two ways. First, we used analysis of covariance to test the hypothesis that  $\log_{10}$  (ant recruitment) increased with  $\log_{10}$  (concentration for sucrose and NaCl), but that recruitment to NaCl, but not to sucrose, increased with distance inland. We did so by binning sites by distance inland ( $< 10$  km [ $n = 2$ ], 10–100 km [ $n = 4$ ], 100–1000 km [ $n = 7$ ], and  $> 1000$  km [ $n = 4$ ]). We also evaluated maximum sucrose and NaCl recruitment across sites across the range of concentrations (typically at 1% NaCl and 10%–20% sucrose) with distance inland. Because sodium use may be confounded by overall activity (as gauged by the activity at sucrose baits), we also evaluated changes in the ratio of sodium use to sucrose use per site. We evaluated taxonomic self-similarity for NaCl and sucrose use for subfamilies and genera that met the minimum criteria of the presence at 10 or more sites and at least five bait hits per site. In each case, we used analysis of covariance, with taxonomic membership as the class variable and distance inland as the continuous variable.

We also tested prediction 2 (lower sodium use with increasing carnivory) in two ways. First, we characterized genera by the Na:CHO ratio based on the total number of hits on NaCl versus those on sucrose. (Two genera, *Gigantiops* and *Pseudomyrmex* were assigned arbitrary values of 2, given absence of any workers at sucrose baits.) We used Davidson's (22, 24) studies of Amazon ants to assign 17 of 32 genera as "herbivore-omnivore" (6 genera) or "carnivore" (11 genera), based on Davidson's threshold of 5.5% delta 15N. We compared Na:CHO ratios for these two trophic groups using a Kruskal-Wallis test. As before, we then took subfamily means and repeated the test at a coarser taxonomic resolution. Second, we used a signed-rank test to compare NaCl and sucrose use in ants of for six pairs of matched green food web and brown food web communities.

**ACKNOWLEDGMENTS.** We thank F. Azorsa, N. Clay, M. Koinig, S. Price, C. Schaller, and N. Zegarra for field and laboratory assistance and O. Acevedo, P. Bucur, P. Jensen, S. Madigosky, N. Pitman, and R. S. Sikes for logistical support. The Panamanian Autoridad Nacional del Ambiente (ANAM), the Costa Rican Ministerio del Ambiente y Energía (MINAE), and the Peruvian Instituto Nacional de Recursos Naturales (INRENA) provided permits. The Amazon Conservatory for Tropical Studies, Amazon Explorama Lodges, the Smithsonian Tropical Research Institute, and the Organization for Tropical Studies facilitated access to field sites. This research was supported in part by the National Geographic Society and the Amazon Conservation Association.

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