NATURE NOTES

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Diet composition of an escaped captive-born southern tamandua (Tamandua tetradactyla) in a nonnative habitat in Asia

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Abstract

Studies on the role of natural predatory instincts in captive-born mammalian myrmecophagy are rare. Consequently, researchers rely extensively on case reports to learn more about the contexts in which predatory behavior occurs among such animals. In this study, we recorded an uncommon case of a captive-born southern tamandua (Tamandua tetradactyla) that accidentally escaped from a zoo into a nonnative habitat in Asia. The southern tamandua was found alive 3 months later. Two fresh fecal samples were obtained, and the diet composition was examined. Three termite species (one family, three genera), and 14 ant species (four subfamilies, nine genera) were identified in the fecal samples. The studied southern tamandua preyed on terrestrial and arboreal ants and termites, as the wild populations of its species do. Ants of the subfamily Myrmicinae and termites of the subfamily Nasutitermitinae were the most abundant prey items in the samples, which is consistent with related reports on the wild populations. Soldier ants constituted <1% of the prey items in the fecal samples, suggesting that the southern tamandua likely avoided preying on ants of the soldier caste. Fungus-growing termites Odontotermes (Isoptera: Macrotermitinae), which are not native to neotropical regions, were also ingested by the southern tamandua. This study provides information on how a captive-born mammalian myrmecophagy applies its natural feeding instincts in nonnative natural settings.

KEYWORDS

anteater, exotic species, myrmecophagy, natural predatory instincts, tamandua, zoo

TAXONOMY CLASSIFICATION

Behavioural ecology; Conservation ecology; Zoology

| INTRODUCTION 1

Xenarthra is one of the four main clades of placental mammals, which includes armadillos, sloths, and anteaters (Gardner, 2008). Anteaters are the least diverse xenarthran group and include pygmy anteaters Cyclopes didactylus, giant anteaters Myrmecophaga tridactyla, and tamanduas Tamandua spp. (Gibb et al., 2016). Two species of tamandua have been described: T. mexicana, also called the northern tamandua, which is distributed from southern Mexico to the northwest Andes in South America (Navarrete & Ortega, 2011), and T. tetradactyla, also called the southern tamandua, which lives in northern and central South America east of the Andes (Hayssen, 2011; Figure 1a). Tamanduas occur in tropical and subtropical areas and inhabit diverse habitats, including

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evergreen forests, transitional forests, savanna, and areas degraded by agricultural use (Hayssen, 2011; Montgomery, 1985; Navarrete & Ortega, 2011; Rodrigues et al., 2001). Both species are capable of moving, feeding, and resting on the ground and in trees (Handley, 1976; Lubin & Montgomery, 1981; Montgomery, 1985). Tamanduas are highly specialized predators, feeding predominantly on ants and termites (Lubin & Montgomery, 1981; Montgomery, 1985; Vaz et al., 2012) and other small arthropods (Hayssen, 2011; Redford, 1985), along with occasional fruit (Meritt, 1976). The proportions of prey items consumed by tamanduas vary among species, individuals, and regions (Montgomery, 1985; Pages, 1970; Rodrigues et al., 2008). Seasonal variations in prey composition and nutritional content among wild tamanduas have also been reported (Lubin & Montgomery, 1981). Consequently, meeting the dietary and behavioral needs of captive and captive-born tamanduas, such as those bred in modern zoos, is challenging (Oyarzun et al., 1996). Captive individuals typically receive artificial or commercial feed (Catapani et al., 2018).

The abundance and ready availability of food supplies in captivity may affect animals' natural predatory instincts (Maple, 1980; Poole, 1987). The factors that affect the survival of captive-born or reintroduced animals in the wild include prerelease training, habitat characteristics, food supplies, and anthropogenic disturbances in the environment (Maran et al., 2009). Jiménez Pérez et al. (2015)

reported the survival rate of reintroduced giant anteaters in Iberá Nature Reserve, Argentina was 76% (11 died out of 47 released animals). There have been few studies on the role of natural predatory instincts in captive-born mammalian myrmecophagy in the wild. Here, we report a rare case of a 3-year-old captive-born female southern tamandua that accidentally escaped from Taipei Zoo into a nonnative habitat. The southern tamandua had not previously lived in the wild nor fed on natural prey items. The studied southern tamandua was found alive after 3 months in the wild. After locating the southern tamandua, we collected two fresh fecal samples and investigated the southern tamandua's diet composition. Specialist species are more susceptible to nonnatives diet than are generalist species because of their narrow feeding habits. Therefore, how this captive-born southern tamandua applied its natural feeding instincts in the wild and how it selected nonnative prey are of interest. We conducted this study to analyze what prey items the captive-born southern tamandua consumed in the nonnative habitat and whether these prey items were similar to those consumed by tamanduas in native habitats. Given that this is the first report of a captive-born southern tamandua surviving for 3 months in a nonnative habitat, it provides valuable insights regarding the adaptation of exotic species and the detailed dietary composition of the southern tamandua, further elucidating the characteristics and behaviors of this highly specialized predator.



FIGURE 1 Geographical distribution of *tamandua* spp. and the study site, Taipei, Taiwan (a). Map modified from Villa and Cervantes (2003); Hall (1981); Wetzel (1985); and Hayssen (2011). Locations where the *T. tetradactyla* escaped and was captured (b). Tamandua photo credit: Yi-tang Chang.

2 | METHODS

2.1 | Study area

The study area is close to Taipei Zoo (24.994676, 121.586293) in northern Taiwan (Figure 1b). Taipei Zoo is bordered by hills to the south and east and has an average elevation of 100 m. The climate of northern Taiwan is generally subtropical. The average monthly temperature in 2020 was 22.6°C (range: 16.2°C in January to 28.3°C in July; ShengKeng station, Taiwan Central Weather Bureau, http:// www.cwb.gov.tw). The annual precipitation in 2020 was 2132.5 mm (Taiwan Central Weather Bureau). The vegetation cover of the study area consists of evergreen broad-leaved forests, bamboo forests, grasslands, and farmlands.

2.2 | Subject

The southern tamandua was captive-born and brought to Taipei Zoo at 12 months in August 2018. The captive environment at the zoo had natural structural features, including tree trunks, branches, and plants with dirt substrates. The southern tamandua received 300g of feed, composed of meal worms, bee pupae, fruits, chitin powder, and vegetables, daily. As of November 3, 2019, the southern tamandua weighed 6.2 kg. On November 28, 2019, the southern tamandua was paired. On May 10, 2020, the southern tamandua gave birth. The cub was nursed by the studied southern tamandua. At 2:30 a.m. on September 1, 2020, the studied southern tamandua escaped from Taipei Zoo with the cub and reached a nearby forest (Figure 1b). The cub was found alive the next day in the culvert of the zoo. After 97 days, on December 6, 2020, the studied southern tamandua was found in a tree den in a secondary forest 3 km away from the zoo (Figure 1b), and its identity was confirmed through microchip scanning. The southern tamandua was subsequently brought to Taipei Zoo for clinical examination. Its body weight was 5.5 kg at the time. On the same day, the southern tamandua received 450g of regular feed.

2.3 | Analysis of fecal composition

On December 7, two of the southern tamandua's fecal samples were collected from its enclosure. The feces from which the samples were collected were defecated 30 min apart. In captivity, tamanduas normally defecate every 2 to 6 days (Meritt, 1975), indicating the southern tamandua feces collected in this study presumably represented its diet composition in the wild from within the preceding week. Both of the fecal samples consisted of the cuticular remains of ants and termites. We sampled 1 g of dry mass from each defecation for prey composition analysis. We performed fecal content analysis and prey item identification to extract prey items and debris by following the fecal filtering procedures proposed by Sun et al. (2019). The filtered prey items were macrophotographed and digitized (see

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Sun et al., 2019 for details). The termites and ants were identified to species level, and the number of individuals was counted using the macrophotographs (Sun et al., 2019). The morphological characteristics of ant head capsules were used for species identification and individual quantification. In termites, the left mandible is more morphologically diverse than the right mandible, and the numbers of left and right mandibles in the feces of animals consuming termites do not differ significantly (Liang, 2017). We therefore used termite left mandibles for species identification and individual quantification. For conehead soldier termites of genus *Nasutitermes*, head capsules were used for species identification and individual quantification (as illustrated in Figure 2). Finally, the prey item information collected in this study (such as subfamily, genus, and lifestyle) was compared with that reported in previously published literature.

3 | RESULTS

A total of 1019 termite individuals across three species (one family, three genera) and 583 ant individuals across 14 species (four subfamilies, nine genera) were identified from the fecal samples (Table 1). Two genera of termites, (Reticulitermes and Odontotermes) and six genera of ants (Technomyrmex, Polyrhachis, Aphaenogaste, Strumigenys, Tetramorium, and Brachyponera) were the first recorded prey items. The ant subfamily Myrmicinae appeared in the samples in the highest frequency (eight species, 57.1% of the total ant species consumed) and number (534 individuals, 92.1% of the total ant individuals consumed) than other ant subfamilies. The arboreal ant Crematogaster rogenhoferi comprised more than half of the ant individuals consumed (333 individuals, 63.4%). Moreover, soldier ants comprised an extremely low percentage of the ant individuals consumed (three individuals, <1%). Termites of the genus Nasutitermes comprised the majority of the termite individuals consumed (748 individuals, 73.4%). Among the Nasutitermes consumed, 629 and 119 were workers and soldiers, respectively. Odontotermes was the second most common genus of termites in the samples, comprising 24.9% of the termite individuals consumed. All of the Odontotermes workers consumed were major workers. Six elytra of a termitophilous beetle, Oreomicrus sp., were also present in the fecal samples (Figure 2).

4 | DISCUSSION

Although the findings of this study are based on limited fecal samples, we were able to collect detailed information regarding the prey composition of the southern tamandua. Our findings also provide insights into the feeding adaptation of a captive-born southern tamandua in a nonnative habitat. Overall, the prey composition of the studied southern tamandua exhibited some common patterns with those of wild populations of tamanduas in South America. For example, the studied southern tamandua consumed both terrestrial and arboreal ants and termites, which is consistent with previous records (Table 1).



FIGURE 2 Macrophotograph of cuticular remains of prey in southern tamandua (*Tamandua tetradactyla*) feces. The feces were collected from a captive-born southern tamandua that escaped from Taipei zoo and lived in the wild for 3 months in Taiwan. Scale bar = 2 mm.

The ant subfamilies Myrmicinae and Formicinae exhibit the highest species richness in the diets of wild tamanduas (Table 1). However, in this study, high species richness was only observed for the subfamily Myrmicinae. This was likely due to the limited number of fecal samples collected or the availability of Myrmicinae is generally abundant in suburban Taiwan. In addition, soldier ants comprised <1% of the ant individuals present in the fecal samples, which suggests that the southern tamandua likely avoided preying on colonies with large numbers of ants of the soldier caste. Social insects typically have well-developed chemical and physical defenses (Redford, 1985), and the defense behavior of ants and termites is effective in repelling ant-eaters (Montgomery & Lubin, 1977).

Montomery (Montgomery, 1985) and Lubin and Montgomery (1981) reported that Nasutitermes termites are an important part of the diets of both tamandua species. Nasutitermes parvonasutus are commonly found in suburban Taiwan (Liang & Li, 2016) and were the species of which the southern tamandua in this study consumed the most. Although N. parvonasutus do not inhabit neotropical regions, Nasutitermes spp. are found commonly in south America (Constantino, 2020). We speculated the studied southern tamandua preyed on a significant amount of N. parvonasutus likely due to the display of their natural predatory instincts. N. parvonasutus does not build arboreal nests but build shelter tubes on tree trunks; mature colonies of this species may also build nests underground (Liang & Li, 2016). In addition, Nasutitermes soldiers are present in greater numbers in nests and in lower numbers in shelter tubes and decomposing wood; and the defense of soldiers is effective in repelling large vertebrate predators, such as tamanduas (Lubin & Montgomery, 1981). Remarkably, a termitophilous beetle, Oreomicrus sp. (Hydrophilidae: Sphaeridiinae: Omicrini), was identified in the southern tamandua's fecal samples. Oreomicrus sp. was

observed in *N. parvonasutus* tunnels in decomposing wood by Liang and Li (2016) in Taiwan. The southern tamandua likely ingested the termitophilous beetles together with *N. parvonasutus* in shelter tubes and decomposing wood during foraging. Montgomery (1985) also reported that tamanduas ingest certain commensal arthropods of social insects.

The fungus-growing termite O. formosanus does not occur in neotropical regions, whereas in Taiwan, it is a widespread species (Chiu et al., 2010). O. formosanus constructs subterranean nests and gallery systems and builds foraging shelter tubes on the surfaces of tree trunks, dead grass, tree branches, leaf litter, and other natural materials (Chiu et al., 2018). The studied southern tamandua ingested a considerable amount of O. formosanus, which was probably because of prey accessibility and availability in the environment (Gallo et al., 2017; Reiss, 2000). In addition, only major worker of O. formosanus was present in the fecal samples; no minor workers, alates, or nymphs were present. Generally, major worker termites are responsible for foraging and building shelter tubes on the ground, whereas minor workers are more commonly involved in feeding and nursing within underground nest chambers (de Oliveira et al., 2021; Wang et al., 2009). Hence, we speculated that the southern tamandua fed on O. formosanus foraging aboveground instead of excavating underground nest chambers.

Reticulitermes are also not native to South America, and they were recently introduced to Chile, Argentina, and Uruguay (Austin et al., 2005; Constantino, 2002). The distribution of the introduced *Reticulitermes* does not overlap with that of south tamandua. *R. flaviceps* is an endemic species and only found in Taiwan, and they are commonly found in lowland forest and also build foraging tubes on the ground (Wu et al., 2019). Hence, we believe the studied tamandua preyed on *R. flaviceps* likely due to the availability on the ground.

	No. of prey species				
Prey items	T. mexicana	T. tetradactyla	Present study (worker, soldier)		Lifestyles
Ant					
Dolichoderinae					
Azteca	? ^h	1 ^{c,d}			Arboreal
Monacis		1 ^d			Arboreal
Technomyrmex			1	(8, NA)	Terrestrial/arboreal
Dorylinae					
Army ant (Unidentified)		1 ^e			Terrestrial
Neivamyrmex	1 ^h				Terrestrial
Ectatomminae					
Ectatomma	2 ^h				Terrestrial
Gnamptogenys		1 ^a			Terrestrial/arboreal
Ecitoninae					
Eciton	1 ^h				Terrestrial
Formicinae					
Brachymyrmex	2 ^h	1 ^a			Terrestrial
Camponotus	5 ^h	6 ^{a,d}	1	(3, 0)	Terrestrial/arboreal
Nylanderia		1 ^a			Terrestrial
Polyrhachis			3	(6, NA)	Terrestrial/arboreal
Myrmicinae					
Acromyrmex	1 ^h	1 ^a			Terrestrial/arboreal
Aphaenogaste			2	(68, NA)	Terrestrial/arboreal
Cephalotes		2 ^h			Arboreal
Crematogaster	2 ^h	2 ^a	1	(333, NA)	Arboreal
Monomorium	1 ^h				Terrestrial/arboreal
Pheidole	2 ^h	4 ^a	3	(123, 3)	Terrestrial/arboreal
Solenopsis	5 ^h	2 ^a			Terrestrial
Strumigenys			1	(1, NA)	Terrestrial
Tetramorium			1	(9, NA)	Terrestrial/arboreal
Trachymyrmex		1 ^a			Terrestrial
Ponerinae					
Brachyponera			1	(29, NA)	Terrestrial
Heteroponera		1 ^a			Terrestrial/arboreal
Odontomachus	1 ^h				Terrestrial
Pachycondyla		1 ^a			Terrestrial
Termite					
Kalotermitidae					
Calcaritermes	? ^f				Terrestrial
Macrotermitinae					
Odontotermes			1	(232*, 22)	Subterranean
Nasutitermitinae					
Nasutitermes	? ^f	4 ^{c,d,g}	1	(629, 119)	Terrestrial/arboreal

TABLE 1 Review of *Tamandua* spp. diet composition compared with that of the southern tamandua from the present study. The first two columns of number of prey species consumed are from previous studies.

TABLE 1 (Continued)

	No. of prey spec	ies consumed			
Prey items	T. mexicana	T. tetradactyla	Present stu	dy (worker, soldier)	Lifestyles
Rhinotermitidae					
Coptotermes	? ^f				Terrestrial
Reticulitermes	? ^f		1	(15, 2)	Terrestrial
Termitinae					
Armitermes	? ^f				Arboreal
Microcerotermes	? ^f	1 ^c			Terrestrial/arboreal
Beetle					
Hydrophilidae					
Oreomicrus			1 ^b		Termitophilous

Note: * = major workers.

References used: [a] Gallo et al., 2017; [b] Liang & Li, 2016; [c] Lubin et al., 1977; [d] Lubin & Montgomery, 1981; [e] Montgomery & Lubin, 1977; [f] Navarrete & Ortega, 2011; [g] Oyarzun et al., 1996; [h] Sandoval-Gómez et al., 2012.

In Brazil, Rodrigues et al. (2001) reported that translocated southern tamandua usually remain close to their release locations, with the largest observed distance between the release site and the recapture site being 2.17 km. This indicates that southern tamandua may adapt to new environments rapidly. In this study, the distance from Taipei Zoo to where the studied southern tamandua was located was long—approximately 3 km. This was likely related to anthropogenic disturbances in the environment, such as free-roaming dogs (i.e., dogs that have owners but are not restricted to a prescribed indoor or outdoor space). The density of free-roaming dogs in northern Taiwan is high and negatively affects the wildlife these dogs encounter (Yen et al., 2019).

In summary, this study reveals the adaptations of a captive-born animal in a nonnative habitat and provides insights into how captiveborn animals apply their natural feeding instincts in natural settings. To expand upon the current results and further elucidate the dietary characteristics of tamanduas in natural and captive settings, the gut microbiomes of wild and captive southern tamandua were analyzed; the results will be provided in a future article.

AUTHOR CONTRIBUTIONS

Nick Ching-Min Sun: Conceptualization (lead); data curation (equal); formal analysis (equal); investigation (lead); methodology (equal); validation (equal); visualization (equal); writing – original draft (lead); writing – review and editing (supporting). Chung-Chi Lin: Conceptualization (supporting); data curation (equal); formal analysis (equal); investigation (supporting); methodology (supporting); supervision (supporting); writing – original draft (supporting). Chun-Chieh Liang: Data curation (supporting); formal analysis (supporting); methodology (equal); supervision (supporting); formal analysis (supporting); methodology (equal); supervision (supporting); formal analysis (supporting); methodology (equal); supervision (supporting); writing – original draft (supporting). Hou-Feng Li: Conceptualization (equal); data curation (equal); formal analysis (equal); funding acquisition (lead); investigation (supporting); methodology (equal); project administration (equal); supervision (lead); writing – original draft (supporting); writing – original draft (supporting); methodology (equal); project administration (equal); supervision (lead); writing – original draft (supporting); writing – original draft (supporting); writing – original draft (supporting); writing – review and editing (equal).

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CONFLICT OF INTEREST

None declared.

DATA AVAILABILITY STATEMENT

All the data are presented in the article (Table 1). No additional data will be uploaded at other place.

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REFERENCES

- Austin, J. W., Szalanski, A. L., Scheffrahn, R. H., Messenger, M. T., Dronnet, S., & Bagneres, A. G. (2005). Genetic evidence for the synonymy of two Reticulitermes species: Reticulitermes flavipes and Reticulitermes santonensis. Annals of the Entomological Society of America, 98, 395–401.
- Catapani, M. L., Pires, J. S. R., & Vasconcellos, A. S. (2018). Single- or pairhoused: Which is better for captive southern tamanduas? *Journal of Applied Animal Welfare Science*, 22, 289–297.
- Chiu, C. I., Li, H. F., & Yang, M. M. (2010). The geographical distribution and the dispersal flight season of Odontotermes formosanus (Isoptera: Termitidae) in Taiwan. Formosan Entomologist, 30, 193-202.
- Chiu, C. I., Yeh, H. T., Li, P. L., Kuo, C. Y., Tsai, M. J., & Li, H. F. (2018). Foraging phenology of the fungus-growing termite *Odontotermes*

- Constantino, R. (2002). The pest termites of South America: Taxonomy, distribution and status. *Journal of Applied Entomology*, 126, 355–365.
- Constantino, R. (2020). Termite database. University of Brasília. http:// termitologia.net
- de Oliveira, M. H., Viana-Junior, A. B., Nascimento, C. C., & Bezerra-Gusmão, M. A. (2021). Worker dimorphism in nasute termites reflects different tasks during food collection. *Journal of Insect Behavior*, 34, 96–105.
- Gallo, J. A., Abba, A. M., Elizalde, L., Di Nucci, D., Ríos, T. A., & Ezquiaga, M. C. (2017). First study on food habits of anteaters, *Myrmecophaga tridactyla* and *Tamandua tetradactyla*, at the southern limit of their distribution. *Mammalia*, 81, 601–604.
- Gardner, A. L. (2008). Magnorder Xenartha. In A. L. Gardner (Ed.), Mammals of South America: Volume 1 marsupials, xenarthrans, shrews, and bats (pp. 127-128). University of Chicago Press.
- Gibb, G. C., Condamine, F. L., Kuch, M., Enk, J., Moraes-Barros, N., Superina, M., & Delsuc, F. (2016). Shotgun mitogenomics provides a reference phylogenetic framework and timescale for living xenarthrans. *Molecular Biology and Evolution*, 33, 621–642.
- Hall, E. R. (1981). The mammals of North America (2nd ed.). John Wiley & Sons, Inc.
- Handley, C. O. (1976). Mammals of the Smithsonian Venezuelan project. Brigham Young University Science Bulletin, Biological Series, 20, 1–89.
- Hayssen, V. (2011). Tamandua tetradactyla (Pilosa: Myrmecophagidae). Mammalian Species, 43(875), 64–74.
- Jiménez Pérez, I., Delgado, A., Di Blanco, Y. E., Abuin, R., Antúnez, B., Galetto, E., Masat, M., Peña, J., Pernigotti, R., Pontón, F., Solís, G., Spørring, K. L., & Heinonen, S. (2015). Reintroducción del hormiguero gigante (*Myrmecophaga tridactyla*) en la Reserva Natural Iberá (Argentina): ¿Misión cumplida? *Edentata*, 16, 11-20.
- Liang, C. C. (2017). Termite species composition in soil and feces of Formosan pangolin (Manis pentadactyla pentadactyla) at LuanShan, Taitung. [M.S. thesis], National Pingtung University of Science and Technology, Pingtung, Taiwan.
- Liang, W. R., & Li, H. F. (2016). Redescriptions of three nasutitermes species (Isoptera: Termitidae: Nasutitermitinae) occurring in Taiwan. Annals of the Entomological Society of America, 109, 779–795.
- Lubin, Y. D., & Montgomery, G. G. (1981). Defenses of Nasutitermes termites (Isoptera, Termitidae) against tamandua anteaters (Edentata, Myrmecophagidae). Biotropica, 13, 66–76.
- Lubin, Y. D., Montgomery, G. G., & Young, O. P. (1977). Food resources of anteaters (Edentata: Myrmecophagidae) I. a year's census of arboreal ants and termites on Barro Colorado Island, Panama Canal Zone. *Biotropica*, 9, 26–34.

Maple, T. L. (1980). Orangutan behavior. Van Nostrand Reinhold.

- Maran, T., Põdra, M., Põlma, M., & Macdonald, D. W. (2009). The survival of captive-born animals in restoration programmes Case study of the endangered European mink *Mustela lutreola*. *Biological Conservation*, 142, 1685–1692.
- Meritt, D. A. (1975). The lesser anteater, Tamandua tetradactyla, in captivity. International Zoo Yearbook, 15, 41–45.
- Meritt, D. A. (1976). The nutrition of edentates. *International Zoo* Yearbook, 16, 38-46.
- Montgomery, G. G. (1985). Movements, foraging and food habits of the four extant species of neotropical Vermilinguas (Mammalia: Myrmecophagidae). In G. G. Montgomery (Ed.), *The evolution* and ecology of armadillos, sloths, and vermilinguas (pp. 365–377). Washington, DC.
- Montgomery, G. G., & Lubin, Y. D. (1977). Prey influences on movements of Neotropical anteaters. In R. L. Phillips & C. Jonkel (Eds.), *Proceedings of the 1975 predator symposium* (pp. 103–131). Montana Forest and conservation Experiment Station, University of Montana.

- Navarrete, D., & Ortega, J. (2011). *Tamandua mexicana* (Pilosa: Myrmecophagidae). *Mammalian Species*, 43(874), 56–63.
- Oyarzun, S. E., Crawshaw, G. J., & Valdes, E. V. (1996). Nutrition of the tamandua: I. nutrient composition of termites (*Nasutitermes* spp.) and stomach content from wild tamanduas (*Tamandua tetradactyla*). *Zoo Biology*, 15, 509–524.
- Pages, E. (1970). Sur Tecologie et les adaptations de Torycterope et des pangolins sympatriques du Gabon. *Biologia Gabonica*, 6(1), 27–92.
- Poole, T. B. (1987). Social behavior of a group of orangutans (*Pongo pyg-maeus*) on an artificial Island in Singapore zoological gardens. *Zoo Biology*, 6, 315–330.
- Redford, K. H. (1985). Feeding and food preference in captive and wild Giant anteaters (*Myrmecophaga tridactyla*). *Journal of Zoology*, 205, 559–572.
- Reiss, K. Z. (2000). Feeding in myrmecophagous mammals. In K. Schwenk (Ed.), *Feeding* (pp. 459–485). Academic Press.
- Rodrigues, F. H. G., Marinho-Filho, J. S., & Santos, H. G. (2001). Home ranges of translocated lesser anteaters (*Tamandua tetradactyla*) in the Cerrado of Brazil. *Oryx*, 35, 166–169.
- Rodrigues, F. H. G., Medri, I. M., Miranda, G. H. B., Camilo-Alves, C., & Mourão, G. (2008). Anteater behavior and ecology. In S. F. Vizcaíno & W. J. Loughry (Eds.), *The biology of the Xenarthra* (pp. 257–268). University Press of Florida.
- Sandoval-Gómez, V. E., Ramírez-Chaves, H. E., & Marín, D. (2012). Registros de hormigas y termitas presentes en la dieta de osos hormigueros (Mammalia: Myrmecophagidae) en tres localidades de Colombia. *Edentata*, 13, 1-9.
- Sun, N. C. M., Liang, C. C., Chen, B. Y., Lin, C. C., Pei, K. J. C., & Li, H. F. (2019). Comparison of two faecal analysis techniques to assess Formosan pangolin *Manis pentadactyla pentadactyla* diet. *Mammalia*, 84, 41–49.
- Vaz, V. C., Santori, R. T., Jansen, A. M., Delciellos, A. C., & D'Andrea, P. C. (2012). Notes on food habits of armadillos (Cingulata, Dasypodidae) and anteaters (Pilosa, Myrmecophagidae) at Serra da Capivara National Park (Piauí state, Brazil). *Edentata*, 13, 84–89.
- Villa, R. B., & Cervantes, F. A. (2003). Los mamíferos de México. Instituto de Biología, Universidad Nacional Autónoma de México y Grupo Editorial Iberoamérica.
- Wang, Z. Y., Mo, J. C., & Lu, Y. J. (2009). Biology and ecology of Macrotermes barneyi (Isoptera: Termitidae). Sociobiology, 54, 777–786.
- Wetzel, R. M. (1985). The identification and distribution of recent Xenarthra (= Edentata). In G. G. Montgomery (Ed.), *The evolution and ecology of armadillos, sloths, and vermilinguas* (pp. 5–21). Smithsonian Institution Press.
- Wu, C. C., Tsai, C. L., Liang, W. R., Takematsu, Y., & Li, H. F. (2019). Identification of subterranean termite genus, *Reticulitermes* (Blattodea: Rhinotermitidae) in Taiwan. *Journal of Economic Entomology*, 112, 2872–2881.
- Yen, S. C., Ju, Y. T., Lee Shaner, P. J., & Chen, H. L. (2019). Spatial and temporal relationship between native mammals and free-roaming dogs in a protected area surrounded by a metropolis. *Scientific Reports*, 9, 8161.

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