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Potential feeding event of *Priodontes maximus* (Cingulata: Dasypodidae) by *Desmodus rotundus* (Chiroptera: Desmodontinae) in the Cerrado, Western Brazil

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ABSTRACT

Hematophagous bats are fascinating mammals due to their social structure and complex anatomical and physiological characters. We present evidence of the potential use of the giant armadillo (*Priodontes maximus*) in the feeding spectrum of the common vampire bat *Desmodus rotundus*. Observations were made in the Parque Sesc Serra Azul, located in the municipality of Rosário Oeste, Mato Grosso, Brazil, using camera traps. Although hematophagous bats use native prey in nature, records of this feeding activity are less common. The use of camera traps has become an efficient tool to obtain ecological data on many mammals and allows us to assess interactions and associations between species in non-invasive ways.

Keywords: camera traps, common vampire bat, giant-armadillo, hematophagous, potential prey

RESUMO - Potencial evento de alimentação de *Priodontes maximus* (Cingulata: Dasypodidae) por *Desmodus rotundus* (Chiroptera: Desmodontinae) no cerrado, oeste do Brasil. Os morcegos hematófagos se caracterizam por uma complexa estrutura social e características anatômicas e fisiológicas particulares. São apresentadas evidências do uso potencial do tatu-canastra (*Priodontes maximus*) no espectro alimentar do morcego-vampiro-comum *Desmodus rotundus*. As observações foram realizadas no Parque Sesc Serra Azul, município de Rosário Oeste, Mato Grosso, Brasil, usando armadilhas fotográficas. Embora os morcegos hematófagos predem espécies nativas na natureza, os registros dessa atividade alimentar são raros. O uso de armadilhas fotográficas tem se tornado uma ferramenta eficiente para obter dados ecológicos sobre diversos mamíferos e permite avaliar as interações e associações entre as espécies de forma não invasiva.

Palavras chave: armadilhas fotográficas, hematófago, morcego-vampiro-comum, presa potencial, tatu-canastra

The subfamily Desmodontinae includes three species, *Desmodus rotundus* (É. Geoffroy, 1810), *Diaemus youngii* (Jentink, 1893), and *Diphylla ecaudata* Spix, 1823

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(Simmons & Cirranello 2020). The species of this subfamily live in sympatry, are endemic to the New World, and subsist exclusively on blood (Reis et al. 2017). They have specific adaptations to their diet, such as large, pointed, stylet-shaped incisors that are longer than the canines and the deeply furrowed lower lip (Greenhall & Schmidt 1988; Reis et al. 2017). Their prey includes birds, wild mammals, domestic animals, and humans (Gonçalves et al. 2002; Dantas-Torres et al. 2005; Galetti et al. 2016). Among these species, *D. rotundus*, or common vampire bat, has the largest distribution, being present in all countries of South America, Central America to Mexico (Greenhall et al. 1983; Simmons & Cirranello 2020).

The common vampire bat has several morphological adaptations (e.g., walking, running, and jumping) that allow the species to feed on the ground. These characteristics are associated with the exploitation of a variety of prey and its greater complexity in foraging strategies (Greenhall 1972; Sazima & Uieda 1980; Riskin & Hermanson 2005; Riskin et al. 2006; Reyes-Amaya & Jerez 2013; Reyes-Amaya et al. 2017). They have an extremely elongated cecum stomach, which allows a considerable amount of blood to be stored (Greenhall & Schmidt 1988), and the ability to inhibit hemostasis in the host, producing natural anticoagulants (Greenhall 1972; Greenhall & Schmidt 1988; Gohlke et al. 1996).

The activity pattern of *D. rotundus* is more intense in the early night hours, with the highest peak in the second hour after sunset (Marinho-Filho & Sazima 1989). They tend to fly low while moving during foraging and attack prey, usually between 50 and 150 cm in height (Greenhall et al. 1971; Ávila-Flores et al. 2019). All or part of the group may disperse or feed together (Carter & Wilkinson 2013; Stockmaier et al. 2018). They also have a visual acuity comparable to some nocturnal rodents, being very likely to use vision for long-distance orientation. Olfactory acuity is similar to that of other small mammals, but it is more sensitive than an insectivorous bat (Bahlman & Kelt 2007). Therefore, they could locate the prey resting areas by the smell of droppings (Bahlman & Kelt 2007). Additionally, vampire bats possess a thermal sensitive organ, the Jacobson organ, which aids identifying the warmest vascularized region of its prey (Greenhall & Schmidt 1988).

It is known that common vampire bats feed mainly on the blood of medium to large mammals (Greenhall et al. 1983; Gonçalves et al. 2020; Carter et al. 2021). Some direct observations of this interaction, with *D. rotundus* feeding on capybaras (*Hydrochoerus hydrochaeris* (Linnaeus, 1766)), were made in Apure in Venezuela (Carranza 1982; Greenhall & Schmidt 1988). In Brazil, 17 observations of these interactions with capybaras were recorded (Gonçalves et al. 2020). Vampire bats were observed feeding on the ear of sea lion (*Otaria flavescens* (Shaw, 1800); Mann 1951 *apud* Catenazzi & Donnelly 2008); this is a common record for the region, where there are also reports of the vampire bat sharing caves with sea lions on the islands off the coast of northern Chile (Mann 1951 *apud* Catenazzi & Donnelly 2008).

Camera traps have helped record wild animals chased or preyed upon by vampire bats. The most common records in the literature are interactions where vampire bats approach other wild animals, such as tapir (*Tapirus terrestris* (Linnaeus, 1758);



Castellanos & Banegas 2015; Galetti et al. 2016; Gnocchi & Srbek-Araujo 2017; Zortéa et al. 2018), and some less common records as with puma (*Puma concolor* (Linnaeus, 1771)), southern tamandua (*Tamandua tetradactyla* (Linnaeus, 1758); Kays 2016), peccary (*Pecari tajacu* (Linnaeus, 1758); Voigt & Kingston 2016; Zortéa et al. 2018), and preying on deers (*Odocoileus virginianus* (Zimmermann, 1780); Sánchez-Cordero et al. 2011; Tello-Mera & Mandujano 2016), and *Mazama americana* (Erxleben, 1777) (Galetti et al. 2016; Zortéa et al. 2018).

Birds are considered a secondary food source of the common vampire bat (Bobrowiec et al. 2015). The few records show interactions with sea birds, such as pelicans, cormorants (Mann 1951 *apud* Catenazzi & Donnelly 2008), and penguins (Luna-Jorquera & Culik 1995). However, with the introduction of domesticated animals in the Neotropics, along with the ongoing defaunation process, vampire bats have expanded their diet to include cattle, horses, goats, pigs and sheep, and less frequent, poultry and humans (Goodwin & Greenhall 1961; Delpietro et al. 1992; Galetti et al. 2016). The introduction of exotic species has also provided new and abundant food resources. *Desmodus rotundus* was photographed preying on a chital deer (*Axis axis* (Erxleben, 1777)) in Argentina, the sambar deer (*Cervus unicolor* Kerr, 1792) in Brazil, and feral pigs (*Sus scrofa* Linnaeus, 1758) in Argentina, Brazil, and Mexico (Costa & Esbérard 2011; Galetti et al. 2016; Pereira et al. 2016; Calfayan et al. 2019; Hernández-Pérez et al. 2019).

We used camera traps to monitor mammals in the protected area of the Parque Sesc Serra Azul, located on the left bank of the Cuiabazinho River, municipality of Rosário Oeste, Mato Grosso, western Brazil (Fig. 1). The study region is part of the Cerrado ecoregion and covers approximately 5,000 hectares (Louzada et al. 2015). The vegetation varies considerably in the area and is partially fragmented by anthropic activities but in an advanced state of recovery, presenting varied physiography as extensive pastures, cerrado, seasonally forests, and flooded forests (Pennington et al. 2006). The climate of the region is typical of the savannah, type “Aw”, according to the classification of Köppen-Geiger, with concentrated rainfall in the summer months (Alvares et al. 2013). Annual rainfall varies between 1,000 and 1,600 mm (Alvares et al. 2013).

Seven camera traps (CamPark, T70 Trail Cameram, and Bushnell) were installed at sampling points within the vegetation zones of the park for exploratory purposes. The cameras operated 24 hours a day, and for each event, a video of 15 or 30 seconds was recorded. One of the videos was recorded from a camera that operated locally for 36 days (December 1, 2019–January 5, 2020; Video S1) and another for 39 days (April 26–June 4, 2021; Video S2).

We describe the potential predation between *D. rotundus* and the giant armadillo (*Priodontes maximus* (Kerr, 1792), a species in the Vulnerable category, VU; IUCN 2021), based on video records from camera traps. The giant armadillo is the largest species of the Cingulata Order, and an adult can reach 155.9 cm in body length and weigh up to 44.4 kg in nature (Silveira et al. 2009). The hairless upperparts are composed of bone armor, usually between 11 to 13 moving bands at the back and 3 to 4 at the back of the neck; these shields are highly flexible between each other (Emmons & Feer 1999).



The common vampire bat was recorded on the back of a giant armadillo on December 21, 2019, at 01:30 (Latitude -14.514306; Longitude -55.756278). The bat was in the median position on the armadillo's back, probably associated with the areas between mobile plates during animal displacement, which facilitate feeding through malleable tissues near the vascularized region (Fig. 2). On May 8, 2021, a second registration was made at 23:59 (Latitude -14.477583; Longitude -55.702083). In this video, the common vampire bat chases and possibly tries to feed on the armadillo's tail (see videos in Supplementary online material).

The two species, bat and armadillo, have nocturnal activity patterns. However, the low incidence of vampire bat records feeding on *P. maximus*, may be directly related to the circadian rhythm of both species and the rarity of the armadillo. Observations through camera traps of a common vampire bat preying on a giant armadillo were recorded in a previous study at 02:45 (Zortéa et al. 2018); however, both species have divergent activity peaks. The vampire bats, more active in the early evening, being able to reduce their activity during the periods of greater intensity of the moonlight (Flores-Crespo et al. 1972; Marinho-Filho & Sazima 1989), and the activity peak of the giant armadillo has been observed between 22:00 and 04:00 (Silveira et al. 2009; Aya-Cuero et al. 2017). Another species of armadillo, *Dasypus novemcinctus* Linnaeus, 1758, also was recorded with the highest activity peak between 18:00 and 24:00 hours (Ríos-Solís et al. 2021), reinforcing a mainly nocturnal pattern of activities for the armadillo's species. Moreover, giant armadillos are not easily found in degraded habitats, while populations of vampire bats tend to be higher in altered areas due to the presence of domestic animals (Delpietro et al. 1992; Gomes & Uieda 2004; Costa & Esbérard 2011; Mialhe 2013). Some other factors can also negatively influence the populations of these bats as decreased availability of food and daytime shelters (Voigt & Kelm 2006; Streicker & Allgeier 2016; Gonçalves et al. 2017). In captivity, the common vampire bat predation was observed on *Dasypus* sp. During the interaction, the bat bit between the bands of the armadillo's tail (Schmidt & Greenhall 1972; Greenhall & Schmidt 1988).

In the last two decades, camera traps have exponentially increased the survey of non-invasive information, which would otherwise be difficult to obtain (Kays 2016). It has become an efficient tool for getting ecological data from many mammals, including small mammals (flying or not), medium and large (Rovero et al. 2014; Kays 2016).

Fecal samples have been used extensively in mammalian diet studies based on the morphological classification of digested prey remains. However, conventional microscope-based fecal analyzes are not possible in hematophagous bats. Therefore, molecular analyses in feces or blood have been successfully used to infer the prey preference of the common vampire bat (Bobrowiec et al. 2015; Bohmann et al. 2018). Molecular analysis has corroborated that the diet of *D. rotundus* is based mainly on the blood of mammals (Bobrowiec et al. 2015; Bohmann et al. 2018), and secondarily on the blood of chickens (Bobrowiec et al. 2015). Although genetic approaches offer an attractive solution to rebuild vampire bat diets with high specificity, existing methods have not yet reached a technical level that would make them profitable and efficient for large-scale applications.



Studies using stable carbon isotopic has shown evidence of the predation of common vampire bats on cattle (Herrera et al. 1998; Voigt & Kelm 2006; Streicker & Allgeier 2016). Because livestock is temporally and spatially more stable, it may be a more viable resource than wild animals. Finally, serological analysis of digestive tract contents, using the precipitin test technique (Greenhall & Schmidt 1988), detect the blood of domestic and wild animals, including humans and squirrels (Greenhall & Schmidt 1988). A less common prey, the South American sea lion *O. flavescens* was recorded by recordings with night-vision cameras and stable isotopes in the diet of *D. rotundus* (Catenazzi & Donnelly 2008).

Records of vampire bat predation on medium to large-mammals species are uncommon. They range from direct visualization and camera traps to more modern techniques, such as isotope and DNA analysis. Camera traps are an additional method as they allow assessing the interactions between species in a context closer to the functionality of natural communities. Compared to other methods, observation can be maintained for longer and is less expensive than laboratory analysis. Our contribution to the record, which is extremely rare and important to understand and predict predation by *D. rotundus*, would help in future studies on the behavior of the two species.

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SUPPLEMENTARY ONLINE MATERIAL

Supplement 1

Video S1. A common vampire bat (*Desmodus rotundus*) was recorded possibly feeding on a giant armadillo (*Priodontes maximus*) at Parque Sesc Serra Azul, Mato Grosso state, western Brazil.

Supplement 2

Video S2. The common vampire bat (*Desmodus rotundus*) chases and possibly tries to feed on the armadillo's tail (*Priodontes maximus*) at Parque Sesc Serra Azul, Mato Grosso state, western Brazil.



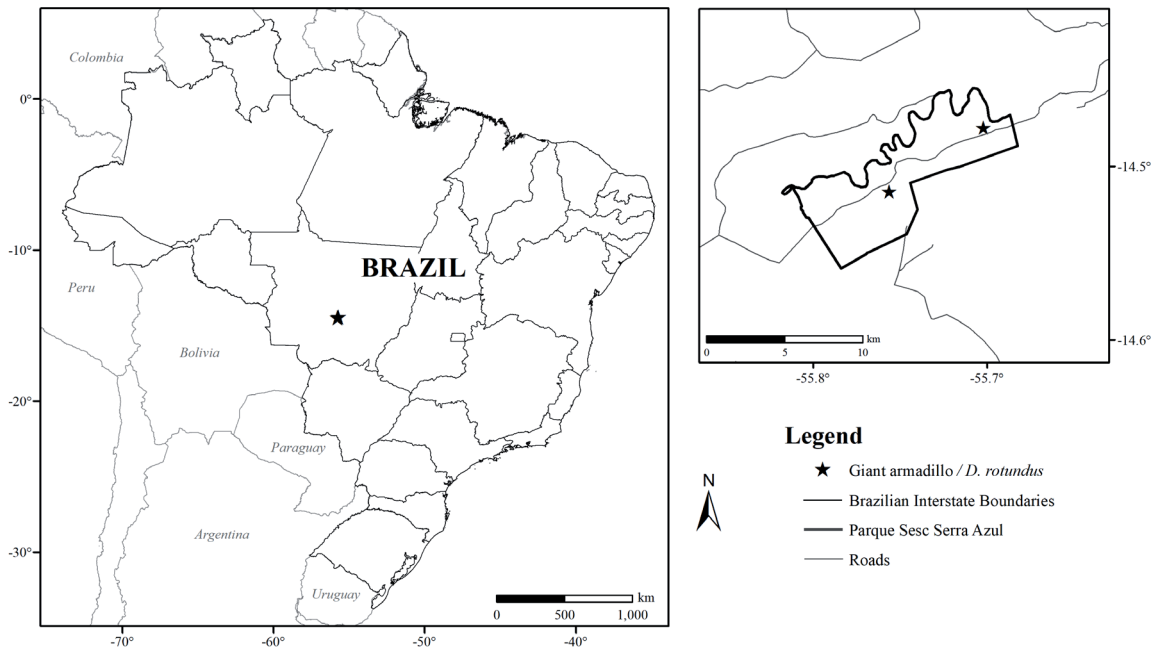


Figure 1. Map of Parque Sesc Serra Azul, Mato Grosso, western Brazil, location of the records made by the present study.



Figure 2. The moment when the common vampire bat (*Desmodus rotundus*) is recorded possibly feeding the giant armadillo (*Priodontes maximus*) in Parque Sesc Serra Azul, Mato Grosso state, western Brazil, recorded by a camera trap.

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