

SHORT COMMUNICATION

Activity pattern of the neotropical marsupial *Didelphis aurita* in south-eastern Brazilian Atlantic Forest

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Abstract: Nocturnal activity is generally presumed, but rarely evaluated in studies of tropical small mammals. When evaluated, activity is frequently presented as a fixed pattern, but actually it is the influence of biotic and abiotic factors on individuals. We investigated effects of age, sex, minimum temperature, reproductive and climatic seasons on activity of *Didelphis aurita* (Didelphimorphia, Didelphidae) in an Atlantic Forest area in south-eastern Brazil from June 2009 to December 2010. We captured 37 individuals, 51 times (28 females and 23 males). Activity of *D. aurita* was mostly nocturnal, but some individuals were active during the day (12%, N = 6). Nocturnal activity was bimodal, more prevalent in adults than juveniles, and different from most neotropical marsupials. Contrary to expectations, individuals of *D. aurita* were more active at colder temperatures, which indicates that low temperatures do not limit foraging behaviour.

Key Words: behaviour, bimodal activity, diel activity, small mammal, temperature

The activity pattern of an organism is a key aspect of its biology, and understanding how biotic and abiotic factors determine daily cycles of activity may reveal adaptations and constraints on animal species (Halle 2001). Periods of high activity should coincide with the best compromise between potentially conflicting demands, for instance feeding vs. reduction of predation risk vs. physiological stress. Temperature affects activity patterns in small mammals (McManus 1969). Endothermic animals maintain body temperature during periods of high or low ambient temperatures, resulting in a high amount of energy lost in thermoregulation. To minimize energy expenditure, possible strategies are to reduce activity, cease activity, or change it to other periods (Fitch & Shirer 1970, McManus 1969). In neotropical marsupials, the only species that reaches temperate latitudes, *Didelphis virginiana*, decreases activity below 8°C (Ryser 1995), and is rarely active in the hottest hours of the day. Even in tropical and subtropical latitudes, temperature may limit activity of species, such as with *D. marsupialis* (Morrison 1946).

Activity of marsupials can vary between sexes, ages and seasons, among other factors (Atramentowicz 1982, Galliez *et al.* 2009, Ladine 1997). In didelphid marsupials, the opossum *Didelphis aurita* Wied-Neuwied, 1826 is one of the most abundant in the Atlantic Forest (Gentile *et al.* 2004). Reproductive and climatic seasons affect its movements; males use larger areas less intensively in the reproductive period, and females increase movements in the dry season in response to reduced food availability (Loretto & Vieira 2005). Differences between sexes and seasons in activity could occur if affected by the same environmental cues, such as food availability and search for mates.

This study describes the activity pattern of *D. aurita* and evaluates the influence of sex, age class, reproductive and climatic seasons, and minimum temperature. We expected temporal differences between sexes and seasons, as observed in movement patterns in the same population (Loretto & Vieira 2005). Additionally, we predicted low temperatures will reduce activity.

We conducted the study in the Serra dos Órgãos National Park (PARNASO; 22°28'S, 42°59'W), an Atlantic Forest area, state of Rio de Janeiro, south-eastern Brazil. The climate is mild humid-mesothermic

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(Nimer 1989), and during the study total monthly rainfall ranged between 2.8 mm and 708.8 mm, with mean minimum and maximum monthly temperatures of 4.5 °C (June 2009) and 34.8 °C (March 2010). We obtained meteorological data from a climatic station of the PARNASO (22°26'S, 42°59'W) at 980 m asl. Sampling points and the climatic station are on the same slope of the mountain range and within the same valley. Therefore, despite the altitudinal difference between climatic station and our study site, minimum temperatures should follow the same pattern of variation throughout the region, and up to the altitude of the climatic station used.

We captured individuals in three grids, in 10 sessions of five nights each from June 2009 to December 2010 (for details see Gentile *et al.* 2004). We equipped Tomahawk live traps with digital timing devices that indicated the time the animal was captured. We recorded capture time, sex, reproductive condition and age class. Individuals were classified in three age classes based on tooth eruption sequence: juveniles, subadults and adults (Macedo *et al.* 2006). The reproductive period of *D. aurita* occurs from July to February, and was defined as the period when every adult or subadult female showed reproduction signs.

Time of capture was converted to minutes after sunset (min as). Subsequent captures of the same individual during the following nights were ignored because of potential stress-related changes in activity. For the purpose of classification of activity patterns, we pooled captures in classes of 2-h intervals. Individuals were classified as diurnal, nocturnal, crepuscular, non-circadian, or acyclic based on time of activity, and as unimodal, bimodal or multimodal according to the distribution of activity (Bartness & Alberts 2000). Activity distributions were modelled as finite mixtures, using the package mixtools (version 1.0.0). We formulated nine models to determine the influence of sex, age class, nightly minimum temperature, reproductive (breeding and non-breeding) and climatic seasons (humid and super-humid) on the time of capture of *D. aurita*. We used Akaike's Information Criterion corrected for small sample size (AICc) to compare models based on the maximum likelihood (Burnham & Anderson 1998). We included an intercept-only model to compare the explanatory power of predictor variables relative to other unaccounted sources of variation. Model selection was performed with the package MuMIn (version 1.9.11). All analyses were performed in R environment (version 2.13.0, R Development Core Team).

We captured 37 individuals of *D. aurita* 51 times (28 females and 23 males): 19 females and 17 males in the reproductive period, nine females and six males in the non-reproductive period. When grouped by climatic season, 14 females and 13 males were captured during the super-humid season, and 14 females and 10 males in the humid season. Regarding age, we captured nine

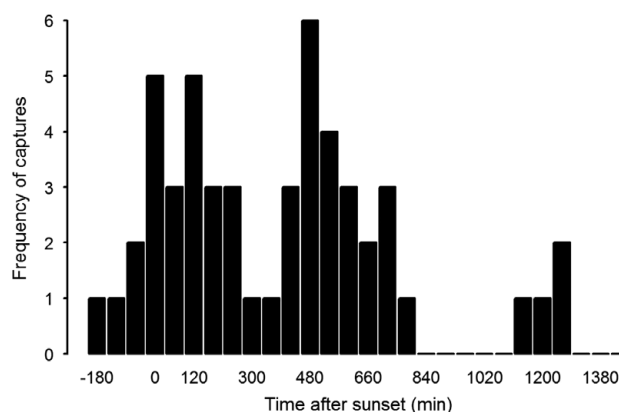


Figure 1. Frequency of captures of the marsupial *Didelphis aurita* in periods of 60 min with relation to local sunset time (zero mark) in an Atlantic Forest area, south-eastern Brazil. Negative values represent activity before sunset.

juveniles, 19 subadults and 23 adults. *Didelphis aurita* was mostly nocturnal, with a bimodal activity pattern in the night, but some individuals were also active during the day (Figure 1). The first activity peak started around sunset, and declined thereafter until achieving minimum activity between 240 and 300 min as. Latter, activity increased, and a second peak was recorded until 780 min as. Daytime activity encompassed 12% (N = 6) of the total activity, and was recorded mostly in adults, with only one juvenile male. Most daytime activity was in the afternoon, but the weak activity in the morning could be a result of interference by researchers who checked traps.

Due to the bimodal distribution of activity, we evaluated the effect of biotic and abiotic factors in the first (117 ± 90.0 min as) and second (560 ± 137 min as) peak of activity separately. Minimum temperature was the top-ranked variable for both activity periods (Appendix 1). In the first peak, the most plausible model included only minimum temperature. The second-ranked model, which also included age, was less plausible. In the second peak, minimum temperature and age formed the only plausible model, with juveniles and subadult less captured than adults.

The number of individuals active during the night reduced as minimum temperature increased. The negative relationship between number of individuals active and minimum temperature, observed in both activity peaks, reveal that low temperatures do not seem to limit activity of *D. aurita* (Table 1). Activity in a wide range of temperatures shows that *D. aurita* is able to persist at different temperatures and may be active at lower temperatures than its congener, *D. virginiana* (McManus 1971).

The bimodal distribution agrees with observations on another population of *D. aurita* (Fernandez 1989), and on a population of the marsupial *Gracilinanus*

Table 1. Standardized parameters of the best models predicting activity pattern of *Didelphis aurita*.

Variables		Estimate	SE
First peak of activity			
Intercept		318	93.6
Tmin		− 10.1	5.24
Second peak of activity			
Intercept		1150	139
Tmin		− 37.23	8.37
Age	Juvenile	− 156	89.8
	Subadult	− 248	59.3

microtarsus in the study area (Matheus Dalloz, pers. comm.). This activity may be a result of intrinsic circadian rhythms, which generally are bimodal (Aschoff 1966). In the field, bimodal activity has also been recorded in bats (Erkert 2001), rodents (Bacigalupe *et al.* 2003) and carnivores (Julien-Laferrière 1993). Conversely, unimodal distribution of nocturnal activity was recorded in other marsupials (Galliez *et al.* 2009, McManus 1971, Ryser 1995). These contrasting results point to flexibility in the activity pattern within didelphid marsupials. However, generalizations are premature considering the few studies conducted with the group.

It is common for young to be active in distinct periods of adults in mammals (Ladine 1997). In this study, we detected differences in activity in the second peak, with juveniles and subadults less active compared with adults. Young opossums tend to move within small areas, and stay close to their dens (Fitch & Shirer 1970). These inexperienced individuals also require less food intake compared with adults, which could lead to a shorter period of activity, concentrated in the first hours of the night.

Different from expected based on movement patterns (Loretto & Vieira 2005), reproductive and climatic seasons did not influence the activity pattern of *D. aurita*. The use of larger areas by males in the reproductive season and by females in the driest season was not reflected in the time range used in the same population, despite the knowledge that, in marsupials, movements as well as activity patterns can be affected by both seasons (Atramentowicz 1982, Loretto & Vieira 2005, McManus 1971). Differences in activity during the year may be apparent when time allocated to distinct activities such as sleeping and eating is taken into account. In winter, when food availability is low, *D. virginiana* was active in the same period as in summer (between 17h00 and 07h00), but devoted almost twice the time to feeding and nest construction (McManus 1971). Nonetheless, time budget studies are difficult to conduct with cryptic animals in the field and studies in semi-natural environments are exceptions (McManus 1971).

The present study demonstrates the dynamic aspect of activity patterns. Individuals of *Didelphis aurita* are

generally nocturnal, as usually assumed, but a great number of individuals were active during the day. Besides, bimodal activity may not be simply a circadian, intrinsic rhythm, but also may have a behavioural and more plastic component; juveniles and subadults have a more unimodal activity. Finally, minimum temperatures commonly encountered in tropical regions do not seem to limit the foraging behaviour of *D. aurita*. Opossums were more active on colder nights, indicating a behavioural adjustment in this environment.

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Appendix 1. Comparison of the top models predicting activity patterns of *Didelphis aurita*. Explanatory variables were age (juvenile, subadult and adult), sex (female vs male), climatic season (humid and super-humid), reproductive period (breeding and non-breeding) and minimum temperature (Tmin). K = number of parameters of the model, AICc = Akaike Information Criteria corrected for small ratio sample size/number of parameters, $\Delta AICc = AICc_i - \text{minimum } AICc$, $w_i = \text{Akaike weight}$. Evaluated models: Tmin; Tmin + Age; Sex; Sex + Age; Sex + Climatic Season; Sex + Reproductive period; Climatic Season; Age; Reproductive period; Intercept-only.

Rank	Model	K	Log-likelihood	AICc	$\Delta AICc$	w_i
First peak of activity						
1	Tmin	3	–127.341	262.1	0.0	0.92
2	Tmin, Age	5	–126.484	267.0	4.9	0.08
3	Climatic season	3	–136.902	281.1	18.9	0
5	Intercept-only	2	–140.723	286.0	23.9	0
Second peak of activity						
1	Tmin, Age	5	–188.706	389.9	0.0	0.99
2	Tmin	3	–196.505	399.9	10.0	0.01
3	Reproductive period	3	–209.675	426.2	36.3	0
8	Intercept-only	2	–213.114	430.6	40.7	0