

Day and night in the subterranean: measuring daily activity patterns of subterranean rodents (*Ctenomys aff. knighti*) using bio-logging

Milene G Jannetti¹, C Loren Buck³, Veronica S Valentinuzzi² and Gisele A Oda¹

¹Laboratorio Binacional Argentina-Brasil de Cronobiología, Departamento de Fisiología, Instituto de Biociências, Universidade de São Paulo, São Paulo, Brazil

²Laboratorio Binacional Argentina-Brasil de Cronobiología, Centro Regional de Investigaciones Científicas y Transferencia Tecnológica (CRILAR), Entre Ríos y Mendoza, s/n, Anillaco, La Rioja 5301, Argentina

³Department of Biological Sciences, Northern Arizona University, PO Box 4185, Flagstaff, Arizona 86011, USA

***Corresponding author:** Departamento de Fisiología, Instituto de Biociências, Universidade de São Paulo—Rua do Matão, 321, Travessa 14, sala 321, 05508-900 São Paulo, SP Brazil. Tel: +55 11 30917480. Email: gaoda@ib.usp.br

While most studies of the impacts of climate change have investigated shifts in the spatial distribution of organisms, temporal shifts in the time of activity is another important adjustment made by animals in a changing world. Due to the importance of light and temperature cycles in shaping activity patterns, studies of activity patterns of organisms that inhabit extreme environments with respect to the 24-hour cyclicity of Earth have the potential to provide important insights into the interrelationships among abiotic variables, behaviour and physiology. Our previous laboratory studies with Argentinean tuco-tucos from the Monte desert (*Ctenomys aff. knighti*) show that these subterranean rodents display circadian activity/rest rhythms that can be synchronized by artificial light/dark cycles. Direct observations indicate that tuco-tucos emerge mainly for foraging and for removal of soil from their burrows. Here we used bio-logging devices for individual, long-term recording of daily activity/rest (accelerometry) and time on surface (light-loggers) of six tuco-tucos maintained in outdoor semi-natural enclosures. Environmental variables were measured simultaneously. Activity bouts were detected both during day and night but 77% of the highest values happened during the daytime and 47% of them coincided with time on surface. Statistical analyses indicate time of day and temperature as the main environmental factors modulating time on surface. In this context, the total duration that these subterranean animals spent on surface was high during the winter, averaging 3 h per day and time on surface occurred when underground temperature was lowest. Finally, transport of these animals to the indoor laboratory and subsequent assessment of their activity rhythms under constant darkness revealed a switch in the timing of activity. Plasticity of activity timing is not uncommon among desert rodents and may be adaptive in changing environments, such as the desert where this species lives.

Key words: Activity patterns, subterranean rodents, *Ctenomys*, nocturnality/diurnality, bio-logging, chronobiology

Editor: Andrea Fuller

Received 28 December 2018; Revised 11 June 2019; Editorial Decision 14 June 2019; Accepted 17 June 2019

Cite as: Jannetti MG, Buck CL, Valentinuzzi VS, Oda GA (2019) Day and night in the subterranean: measuring daily activity patterns of subterranean rodents (*Ctenomys aff. knighti*) using bio-logging. *Conserv Physiol* 7(1): coz044; doi:10.1093/conphys/coz044.

Introduction

Massive changes in climate patterns can compromise organismal fitness and population persistence of many species. Biological systems are dynamic but exhibit stability across time while being responsive to changes in external and internal environments. Characteristically, animals are capable of change either through phenotypic plasticity or microevolutionary processes. While most studies of the impacts of climate change have investigated shifts in the spatial distribution of organisms (Root, 1988; Root *et al.*, 2003; Sinervo *et al.*, 2010), shifts in seasonal phenology (Chevin *et al.*, 2010; Lane *et al.*, 2012; Shen *et al.*, 2015; Williams *et al.*, 2017) or daily timing of activity are other important adjustments that can be made by animals in a changing world (Daan, 1981; Halle and Stenseth, 2000; Kronfeld-Schor and Dayan, 2003; Levy *et al.*, 2018).

The light/dark cycle of Earth imposes a fundamental 24-hour pace that structures the timing of biological activity. In mammals, this is achieved by light/dark entrainment of endogenous, hypothalamic circadian clocks, resulting in rhythmic, 24-hour activity/rest alternations (Rusak, 1982). This endogenously generated rhythm can be profoundly altered by environmental cues, a phenomenon known as 'masking' (Aschoff *et al.*, 1982). For example, ambient temperature shapes activity patterns of mammals via direct, acute stimulation or inhibition of behavioural activity (Kenagy, 1973; Melcher *et al.*, 1990; Kenagy *et al.*, 2002a; Rezende *et al.*, 2003; Long *et al.*, 2005). Due to the importance of light and ambient temperature cycles in shaping activity patterns, it is interesting to study their effects on organisms that inhabit extreme environments with respect to the 24-hour cyclicity of Earth. The subterranean environment offers an opportunity for these studies, due to the constant darkness and attenuated temperature cycles of the burrow environment, characteristics thought to diminish the selective pressure of keeping daily activity patterns (Beale *et al.*, 2016).

There are eight lineages of extant subterranean rodents comprised of approximately 146 species (Cook and Lessa, 1998). The most speciose family of subterranean rodents is Ctenomyidae (parvorder Caviomorpha), comprising almost 60 extant species (Cook and Lessa, 1998; Parada *et al.*, 2011) all from the same genus *Ctenomys* sp. (tuco-tucos). This is a relatively understudied group with respect to their activity patterns in the field. Ctenomyids are endemic to South America and are widely distributed from high altitudes to sea level, from moist grass lands to arid deserts (Vassallo, 1998; Lacey *et al.*, 2000). Most of ctenomyids' diet is composed of aerial parts of plants; therefore, their foraging occurs primarily on the surface (Lacey *et al.*, 2000). This may explain why tuco-tucos have maintained functional eyes of moderate size, in contrast to spalacids and bathyergids of Africa and Asia (Borghi *et al.*, 2002).

Direct observations of individual tuco-tucos from northern Monte desert Argentina (*Ctenomys aff. knighti*), indi-

cate that they emerge to the surface mainly for foraging and soil removal associated with tunnelling (Tomotani *et al.*, 2012). Automated recordings of light-exposure using light-loggers confirmed that tuco-tucos emerge to the surface on a daily basis (Flôres *et al.*, 2016). Standard chronobiological lab protocols showed that their circadian clocks are functional and entrained by light/dark cycles (Valentinuzzi *et al.*, 2009; Flôres *et al.*, 2013) and through computer simulations it is clear that the unique light exposure patterns of tuco-tucos observed in the field are sufficient to synchronize their biological clocks (Tomotani *et al.*, 2012; Flôres *et al.*, 2016).

Several studies in non-subterranean rodents have shown that air and operative temperatures play an important role in determining whether the animal stays on the surface or retreats to underground burrows for thermal shelter (Melcher *et al.*, 1990; Kenagy *et al.*, 2002a; Rezende *et al.*, 2003; Long *et al.*, 2005). Conversely, subterranean tuco-tucos spend most of their time belowground, only emerging to the surface for brief episodes. It is not clear, in this scenario, whether air temperature plays a similar role in promoting surface emergence or how underground temperature influences this decision.

We hypothesized that an interplay between these two temperatures would have masking effects on the timing of tuco-tuco emergence to the surface. To test this main objective, we outfitted tuco-tucos with light-loggers and simultaneously measured air, operative and underground temperatures and wind velocity.

We additionally quantified phase, amplitude and duration of overall daily activity by deploying miniature accelerometers to four of the animals with light-loggers. This second objective was motivated by the fact that the diurnal surface emergence of tuco-tucos in the field contrasts with the nocturnal wheel-running activity under laboratory conditions (Valentinuzzi *et al.*, 2009; Tachinardi *et al.*, 2014). This apparent difference in phase of activity between the laboratory and field could have been due to different methods of assessing activity (i.e. wheel running in the laboratory and light-loggers in the field); thus, our current study is the first to test activity timing in both the laboratory and field employing a single method (accelerometry) to quantify activity.

Methods

Study population

Tuco-tucos *Ctenomys aff. knighti* occur in Anillaco and surroundings, in La Rioja, Argentina (28°48'S; 66°56'W; 1445 m). This arid to semiarid area is located in the Monte desert with average daytime temperatures ranging between 18°C in winter and 28°C in summer and average nighttime temperatures between 9°C in winter and 18°C in summer [2006–16 AIRX3STM v006 dataset, obtained from Giovanni

online data system, developed and maintained by the NASA GES DISC (Acker and Leptoukh, 2007)]. Average annual precipitation is variable, ranging between 100 and 450 mm. Precipitation events are generally limited to summer and droughts can last up to 7 months (Abraham *et al.*, 2009). The soil is sandy and the predominant vegetation in these regions is a shrubby steppe, with characteristic Zygophylaceae, Fabaceae and Cactaceae flora (Fracchia *et al.*, 2011; Aranda-Ricket, 2014).

Seventeen adult tuco-tucos *Ctenomys aff. knighti* were live trapped in Anillaco from 2016 to 2017 using custom made traps (Tomotani *et al.*, 2012). Captured animals were hand carried in traps to the adjacent laboratory inside the *Centro Regional de Investigaciones Científicas y Transferencia Tecnológica*, located in Anillaco. Tuco-tucos were weighed (CSseries, OHAUS, ± 1 g precision) and placed individually into acrylic cages with wire tops ($53 \times 29 \times 27$ cm) for an average of 1 month. Individuals were uniquely marked with a subcutaneously injected microchip (Allflex®, Brazil). Cages were kept in a room with constant temperature $24 \pm 2^\circ\text{C}$, natural lighting and minimal noise. Animals were fed once a day at arbitrary hours. Food consisted of carrots, sweet potatoes, oatmeal, sunflower seed and rodent pellets. Tuco-tucos do not drink free water (Buffenstein, 2000).

All procedures of this work followed the guidelines established by the *American Society of Mammalogists for animal care and handling* (Sikes, 2016), and were approved by the Comissão de Ética no Uso de Animais from *Instituto de Biociências—USP* (nº 273/2016), and were authorized by the *Dirección General de Ambiente y Desarrollo Sustentable—Secretaría de Ambiente del Ministerio de Producción u Desarrollo Local*, La Rioja, Argentina (nº 028/2010 and 062/2008).

Arenas and environmental measurements

Tuco-tucos were maintained in one of three semi-natural outdoor enclosures (arenas) containing native vegetation. The arenas are adjacent to one another and surrounded by wire fencing above ground (arena 1, $10 \times 6 \times 1.5$ m or arenas 2 and 3, $12 \times 6 \times 1.5$ m) and concrete block 1 m below ground to inhibit escape of the tuco-tucos. A mesh covered each arena to prevent aerial predation. Only one animal at a time was kept inside each arena. No food was provided during the experiments; thus, animals were reliant upon foraging on natural vegetation.

A meteorological station (HOBO®, Onset, USA), 1 m from the arenas, provided hourly records of air temperature (T_{air}) (at 1.5 m high) and wind velocity (at 2.1 m height). Subsurface temperatures were measured and recorded each hour (HOBO®, Onset, USA; accuracy $\pm 0.53^\circ\text{C}$ from 0°C to 50°C) at 20 (T_{und}), 40 and 60 cm underground using sensors buried in sand. Reported average burrow depths of *Ctenomys* sp. range from 13 cm to 45 cm (Altuna *et al.*, 1999; Antinuchi and Busch, 1992). Operative temperature (T_e) was measured

and recorded each hour (HOBO®, Onset, USA; accuracy $\pm 0.53^\circ\text{C}$ from 0°C to 50°C) by a sensor wrapped in copper and inserted inside a taxidermied animal ('mannequin') (Bennett *et al.*, 1984; Kenagy *et al.*, 2002a) filled with cotton. The cotton filled mount was validated against a dead animal (Supplementary Fig. 1). The mannequin was exposed to direct sunlight in the soil surface of a fenced area of an arena. Continuous exposure to the elements and concomitant degradation required that mannequins be exchanged every 6 months. In 2015, 2016 and 2017, we obtained T_e data for 17%, 66% and 81% of the year, respectively. Environmental variables from the meteorological station and underground sensors were recorded without interruption from 2015 to 2017. Photoperiod, as well as civil twilight times (day and night onsets), were obtained from data available online (Time and Date AS, 2018).

Measurement of activity/rest rhythm

Bio-loggers were used to assess light exposure (i.e. time on surface) and gross motor activity of tuco-tucos in the arenas. Both loggers were affixed to a collar made of a cable tie inserted through pliable silicon tubing (Williams *et al.*, 2014). Seventeen animals were affixed with bio-loggers, 11 of them with both accelerometer and light-logger and 6 with only a light-logger. We obtained records for six individuals (five females $141 \text{ g} \pm 23 \text{ g}$ and one male 191 g). Four of these (three females and one male) had both a light logger and an accelerometer and the remaining two only had light loggers. One of our original 17 animals died of unknown causes and the remaining 10 individuals either escaped or were lost to predation. The loss of animals to predation and escape was both surprising and unfortunate given the construction of the arenas that included a 1-m buried wall and fencing around and across the tops of the arenas. Clearly, a 1-m buried wall is not always sufficient to contain tuco-tucos and wire mesh fencing is not adequate to block the entrance of some predators (e.g. snakes). In this context, light exposure data of six individuals recorded for 7 days in July and August of 2015, five of them from Flóres *et al.* (2016), were added to our analyses of time on surface, to increase sample size. The same environmental measurements described above were taken during all of these additional recordings, except for T_e . Since we had T_e data for only 17% of the year of 2015, only four of the additional individuals from Flóres *et al.* (2016) were used when verifying association between time on surface and operative temperature.

Time on surface was deduced from light exposure, sampled every 1 minute and only the maximum sample value recorded within each 5 minutes by light loggers ($15 \times 6 \times 6$ mm; 0.65 g; W 65, Migrate Technology, UK). Measurement range of light loggers is 1–19 000 lux and resolution is of 249 discrete levels, with no sensitivity to detect moonlight. Gross motor activity was recorded every 1-second in three spatial axes (XYZ) via accelerometry ($23 \times 12 \times 10$ mm; 2 g; Axy-3, TechnoSmart, Italy) with measurement range of ± 4 G-forces.

Experimental protocol

This study was conducted across 2 years (2016–17) during the months of April–August. The six animals from which we have data were released individually inside the arenas and were re-captured to recover devices. Deployment duration ranged from 8 to 68 days, because the duration was adjusted throughout the experiment, considering that increased deployment durations also increased chance of death, predation or escape. Individual animals were only tested in the arenas one time. Re-captured animals were taken to the laboratory where devices were removed and data were downloaded. Animal #221 took part on a parallel experiment after its recapture. Therefore, three of the four animals with accelerometers were kept with their sensors upon arrival and then transferred to cages placed inside light-tight ventilated boxes, under constant darkness (DD) and constant temperature ($24^{\circ}\text{C} \pm 2^{\circ}\text{C}$) conditions. Food was provided once every two nights at arbitrary times. Activity of these animals was recorded by accelerometers over at least 10 days.

Analysis

All mathematical operations and statistical analyses were performed using R software (R Core Team, 2018). Light-logger recordings with values ≥ 2 lux were considered episodes on surface. Acceleration data were collected from the three orthogonal axes and used to calculate the overall dynamic body acceleration (ODBA) as described in Williams *et al.* (2016). This method consists of subtracting the moving average from the data of each axis, using a 10-second time window. ODBA sums the resulting absolute values. Activity episodes were defined by episodes where ODBA values were higher than the arithmetic mean of all values from that animal, which was ground truthed against laboratory observations. All ODBA values below the arithmetic mean were considered episodes of ‘low body movement’, equivalent to rest. Additionally, high activity episodes for each animal were defined as episodes of body movement higher than half of the maximum value from that animal, which represents the majority of the activity bouts that can be visually observed in the actograms and supposedly corresponds to energy demanding activities.

Time on surface and gross activity data were visualized in actograms made with El Temps software (Díez-Noguera, 2019). Data from both the light and activity loggers were superposed to qualitatively evaluate the temporal association between the two variables.

For the four individuals from which we have information on both acceleration and light exposure, we calculated the percentage of time, relative to 24 hours, in which activity episodes occurred (‘activity’ vs. ‘rest’). From total daily activity time, we also calculated the percentage of ‘diurnal’ vs. ‘nocturnal’ activity, correcting for the day length

of each season, using the following formula modified from Halle and Stenseth (2000)

$$P = \frac{\frac{\sum c_L}{h_L}}{\frac{\sum c_L}{h_L} + \frac{\sum c_D}{h_D}} \times 100,$$

where P is the percentage of diurnal activity, $\sum c_L$ and $\sum c_D$ are the sum of activity records obtained during the day and night, respectively and h_L and h_D are the duration of day and night (civil twilight times). As noted above, the light logger only detects time on surface that occurs in daylight. Therefore, we also calculated the percentage of diurnal activity that occurred on the surface (‘surface’ vs. ‘subterranean’). High activity bouts (body movements higher than half of the maximum value of each animal) were also quantified. We obtained percentage of high activity bouts that happened during daylight hours and those that happened when animals were on the surface. For each of the three animals transferred to the lab, the average value of overall activity in the enclosures and in the lab were compared in order to quantitatively measure the difference between them. The resulting proportions for each animal were averaged.

Gathering data from light loggers of the six animals (2016–17) and the six additional animals recorded in 2015, we verified possible differences between the amount of time on surface of males and females with unequal variance t-test (Ruxton, 2006). We used a generalized linear mixed model (‘glmmTMB’ function) to evaluate the degree of masking on time on surface by environmental variables (Zuur *et al.*, 2009). Light logger data from 10 animals were used in this analysis, since Te data was not collected simultaneously for two animals recorded in 2015. Proportion of surface episodes within 3-hour intervals were used as a response variable. This was calculated as the percentage of episodes classified as aboveground (light-logger values equal to or higher than 2 lux) out of a total of 36 recordings (3-hour interval). Explanatory variables in our model were Te, T_{und}, wind speed (Wind) and time of day as a categorical variable, divided in 3-hour blocks (Hour), which included the following categories: ‘A’ for hours 09:00, 10:00 and 11:00; ‘B’ for 12:00, 13:00 and 14:00; ‘C’ for 15:00, 16:00 and 17:00; and ‘D’ for 18:00, 19:00 and 20:00, considering that environmental measurements were made each hour. Animal identification number (ID) was inserted as a random effect variable. Possible linear correlations between independent variables were verified in linear models, considering correlations significant when $|R| \geq 0.5$. Variance inflation factor (VIF) was also calculated to check for collinearity between independent variables included in each model. GVIF values presented for Hour were corrected for degrees of freedom according to Fox and Monette (1992) and squared for comparison with VIF values of other variables. Final model selection was based on absence of correlated independent variables and use of AIC value, selecting the model with lowest AIC.

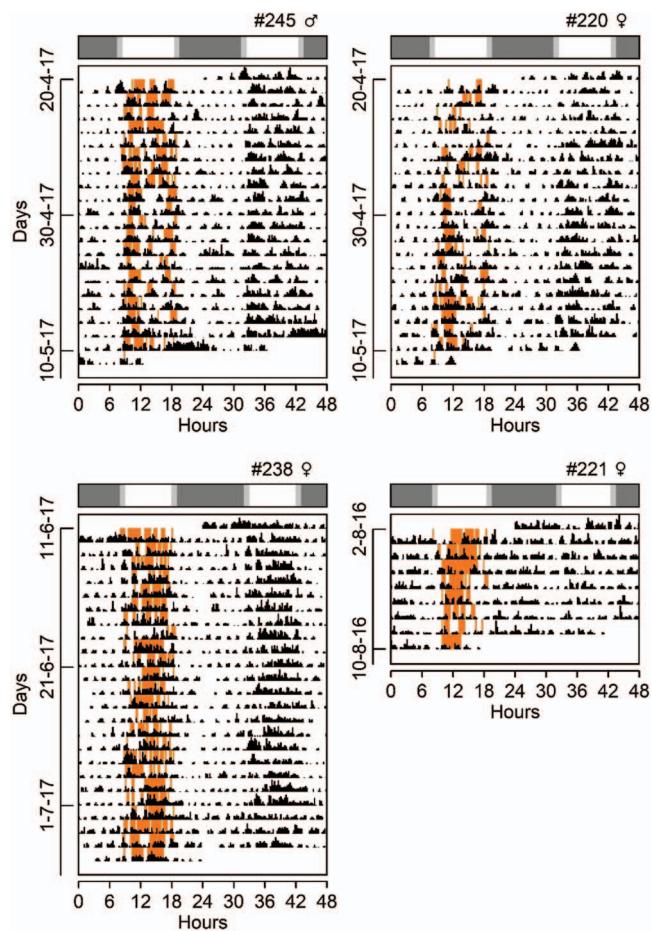


Figure 1: Double-plotted actogram of daily rhythms of gross motor activity and diurnal time on surface of tuco-tucos kept inside arenas in the field in Anillaco, La Rioja, Argentina, from April to August of 2016 and 2017. Orange marks: time on surface during daylight hours. Black marks: gross motor activity measured with accelerometers. Bar above actograms: natural photoperiod at which each recording started (white: day; grey: night). Top right of each actogram: animal identification number and sex.

We calculated and summarized the average duration of time on surface for each hour of the day. To every animal's record, we obtained a curve, calculating the average duration of time on surface for each time of day. All animals' mean curves were then summarized as an average curve ($n=12$). Similar steps were done to obtain duration of time on surface for each interval of T_e ($n=10$) and each interval of T_{und} ($n=12$). All data collected at night were excluded from these calculations since light-loggers do not detect time on surface when it is dark. Pearson's linear correlation coefficient was calculated for the average points of surface episodes $\times T_e$ and surface episodes $\times T_{und}$ final curves. Finally, data of duration of surface episodes per hour and per temperature intervals (T_e or T_{und}) were visualized simultaneously in three axes figures.

Results

During the April to August (winter) study period, T_e daily range was 28.9°C on average. The maximum T_e was 50.2°C

at 13:00 in August 2015 and T_e minimum was -11.8°C at 08:00 in July 2017. Average daily range of T_{und} was 3.7°C . T_{und} maximum was 23.8°C at 00:00 in May 2015 and T_{und} minimum was 5.2°C at 15:00 in June 2016. The lowest values of T_{und} at 20 cm deep occurred regularly near noon, when T_e values were the highest. When T_{und} was compared to deeper measurements, the latter presented a more delayed and flatter variation profile (Supplementary Fig. 2). In La Rioja, maximum day length is 14.03 hours at summer solstice and minimum day length is 10.25 hours at winter solstice.

Accelerometer data revealed that, in addition to activity during the day, *Ctenomys aff. knighti* are also active during the night (Fig. 1). However, on average, 77% of high activity bouts occurred during the daylight hours. Additionally, 47% of these high activity bouts corresponded to when the animals were on the surface. Interestingly, two of the four individuals outfitted with both light and acceleration loggers displayed episodes of low body movement (ODBA below the arithmetic mean) while on the surface (Fig. 2).

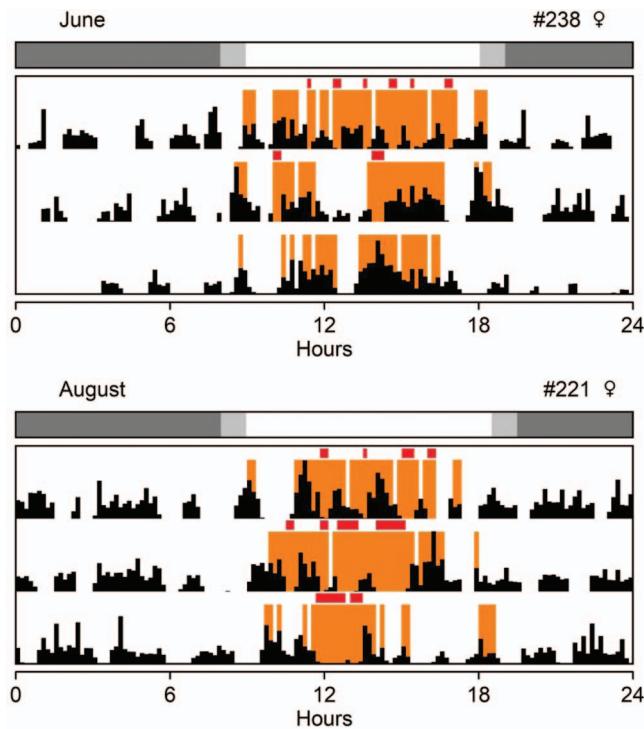


Figure 2: Three-day actogram of individuals #238 and #221 inside arenas in the field. Orange marks: time on surface during daylight hours. Black marks: gross motor activity measured with accelerometers. Red marks: aboveground episodes with low body movement. Bar above actograms: natural photoperiod at which each recording started (white: day; grey: night). Measurements were made on August 2016 (#221) and June 2017 (#238).

Considering all activity episodes (ODBA values above the arithmetic mean), animals were active on average 14.47 hours (60%) of the day, and three of the four individuals expressed >60% (8 hours) of their activity during daylight hours. There was an interindividual standard variation of 1 hour (8.6%) in the time that animals spent on the surface (Table 1). Considering all light logger recordings ($n = 12$), the amount of time on surface did not differ between males and females (unequal variance t-test $P = 0.9$) (Supplementary Table 1).

On average, time on surface was concentrated around noon (Fig. 3). Despite the high standard deviation of average values of time on surface, this variable increased with increasing values of T_e (Pearson correlation coefficient for average points: $R^2 = 0.88$, $P < 0.001$). Standard deviation of time on surface increased when $T_e > 25^\circ\text{C}$. The time tuco-tucos spent on surface increased when underground temperature at 20 cm deep decreased. There was one exception to this pattern where, in a particularly cold, snowy day, individual (#193) was on the surface for an average 5.6 minutes (10% of 1 hour) when T_{und} was 6.5°C and remained underground (0% time on surface) when T_{und} was 4.6°C (Pearson correlation coefficient for average points: $R^2 = 0.20$, $P = 0.10$ considering these occurrences and $R^2 = 0.92$, $P < 0.001$ without these occurrences). When both time of day and T_e or T_{und} effects are considered together, it is evident that while time on surface

between 10:00 and 15:00 occurs across a wide range of T_e , within this timespan, episodes on surface occur mostly when T_{und} values are limited to $7\text{--}12^\circ\text{C}$.

The final generalized linear mixed model included T_{und} , T_e , Hour and Wind, which explained almost 60% of the duration of time on surface (Table 2), accounting for the random effect of animal's ID (contribution of 7.7%). Models with both T_e and T_{air} were not considered since linear correlation between these variables was 67% ($P < 0.001$) and these models resulted in VIF of 5.6 or higher for T_e . The second highest linear correlation value was between T_e and Hour (45%, $P < 0.001$). Considered separately, T_e explained 31% of time on surface (12% attributed to random effect); however, due to its correlation with Hour being close to the threshold of 50% previously established, high R^2 values of $T_e + \text{Hour}$ models must be viewed with caution (in T_e and Hour model, GVIF for Hour: 1.3; VIF for T_e : 2.4). Meanwhile, when T_{und} is considered separately, this model's R^2 is 16% (5% attributed to random effect). Furthermore, the model with T_{und} and Hour (GVIF for Hour: 1.1; VIF for T_{und} : 1.2) received an AIC score 30 units lower than that including T_e and Hour, indicating a higher contribution of T_{und} than T_e to explain time on surface. This is observed in the projection of the model (Fig. 4), where time on surface increases around 12:00, when T_{und} is minimal and T_e is maximal.

Table 1: Quantification of daily activity duration of the four tuco-tucos with both activity and light loggers, when kept inside arenas in the field

Animal number	D	Activity (%)	Rest(%)	Diurnal (%)	Nocturnal (%)	Surface (%)	Underground (%)
245	20	61.8	38.2	64.7	35.3	30.9	69.1
220	20	54.1	45.9	64.1	35.9	18.2	81.8
238	24	60.5	39.5	66.7	33.3	36.2	63.8
221	8	64.9	35.1	44.9	55.1	37.7	62.3
Average	18 ± 7	60.3 ± 4.5	39.7 ± 4.5	60.1 ± 10.2	39.9 ± 10.2	30.8 ± 8.9	69.2 ± 8.9

Shown is the percentage of gross motor activity vs. rest (body movements below average value for each individual) related to the 24 hours of the day; percentage of diurnal vs. nocturnal activity related to total activity time (corrected for day length, using formula described in 'Methods'); and percentage of surface vs. underground activity related to total diurnal activity. D: number of recording days included in the calculations for each animal.

The three individuals outfitted with accelerometers and transferred to laboratory under DD conditions also displayed activity bouts during both day and night in the lab. However, while in the outdoor arenas these animals expressed 77% of their high activity bouts during daytime, this proportion decreased to just 15% upon transfer to the lab (i.e. 85% of their high activity bouts happened during external nighttime hours) (Fig. 5). Finally, comparing average field and lab levels for each individual, overall daily activity levels were on average 1.6 times higher in the field than in the lab.

Discussion

Studies of activity patterns in the field and laboratory allow us to understand ecological aspects as well as endogenous mechanisms of biological timing and its plasticity (Marques and Waterhouse, 2004). However, recording of activity in the field constitutes a logistical challenge (Halle and Stenseth, 2000; Dominoni *et al.*, 2017), particularly for subterranean rodents, with varying degrees of underground and aboveground activity. While most studies of activity patterns in free-living subterranean/fossorial mammals have been conducted with mole rats from Africa and Asia (Nevo *et al.*, 1982; Ben-Shlomo *et al.*, 1995; Lovegrove and Muir, 1996; Riccio and Goldman, 2000; Oosthuizen *et al.*, 2003; Skliba *et al.*, 2007; Vlasatá *et al.*, 2017), caviomorph subterranean/fossorial species from South America have also added important insights to this comparative study (Kenagy *et al.*, 2002a, 2002b; Rezende *et al.*, 2003; Tomotani *et al.*, 2012; Estevan *et al.*, 2016). In particular, the influence of ambient temperature on the surface activity of subterranean/fossorial rodents has been well studied in the South American degu (Kenagy *et al.*, 2002a, 2002b) and coruro (Rezende *et al.*, 2003).

Several studies with epigaeous species have analysed the correlation between burrow retreats and ambient temperatures, indicating that burrows serve as thermal refugia for these species that live mostly aboveground (Melcher *et al.*, 1990; Long *et al.*, 2005; Fick *et al.*, 2009). This is particularly important for desert rodents that can unload excess heat inside cool burrows, saving evaporative water in thermoregulation (Bartholomew 1964; Chappell and

Bartholomew, 1981; Bennett *et al.*, 1984). Conversely, underground temperature plays an important role for subterranean tuco-tucos, in the opposite movement of emergence to the surface, in winter. Across most of day in winter, underground temperature (14.3°C on average) is below the thermoneutral zone of tuco-tucos (23–33°C) (Tachinardi *et al.*, 2017) and a 24-hour variation is detectable at 20-cm depth with amplitude 3.7°C. Interestingly, temperature maxima occur around midnight and minima around noon, coinciding with the highest operative temperatures aboveground. Thus, emergence to the surface during the day is favoured for thermoregulatory reasons, which explains the high total time subterranean tuco-tucos spend on the surface (24% of day time, 3 hours per day on average) in this season. Finally, a higher contribution of T_{und} than T_e to explain time on surface is further observed in the projection of our generalized linear mixed model (Fig. 4). Although time on surface increases around 12:00 when T_{und} is minimal and T_e is maximal, it starts to decrease while T_e remains high, from 13:00 to 15:00, along with an increasing T_{und} .

Interestingly, combined light-logger and accelerometer data in winter revealed several incidences of low activity while animals were on the surface (Fig. 2). This may be indicative of basking, which would be in accordance with the above association between time on surface and thermoregulation. Basking has been traditionally associated with ectotherms (Bogert, 1949) but recent findings have established their role in small rodents during early mornings, rewarming from daily torpor (Geiser *et al.*, 2002; Warnecke *et al.*, 2008). Basking at random times has been seen less often, but has been identified in subterranean rodents such as the African ice rat (*Otomys sloggetti robertsi*) (Schwaibold and Pillay, 2006) and African mole rat (*Tachyoryctes microcephalus*) (Vlasatá *et al.*, 2017). We hypothesize that tuco-tucos are also basking during those times on the surface without movement. Our previous observational studies indicated that tuco-tucos do not wander on the surface for foraging purposes; but rather, they only leave the burrow for very few seconds to acquire vegetation and immediately return (Tomotani *et al.*, 2012). The only activity they perform with their whole body exposed for long durations (minutes to an hour) is soil removal, an activity that coincides with their highest locomotor activity aboveground

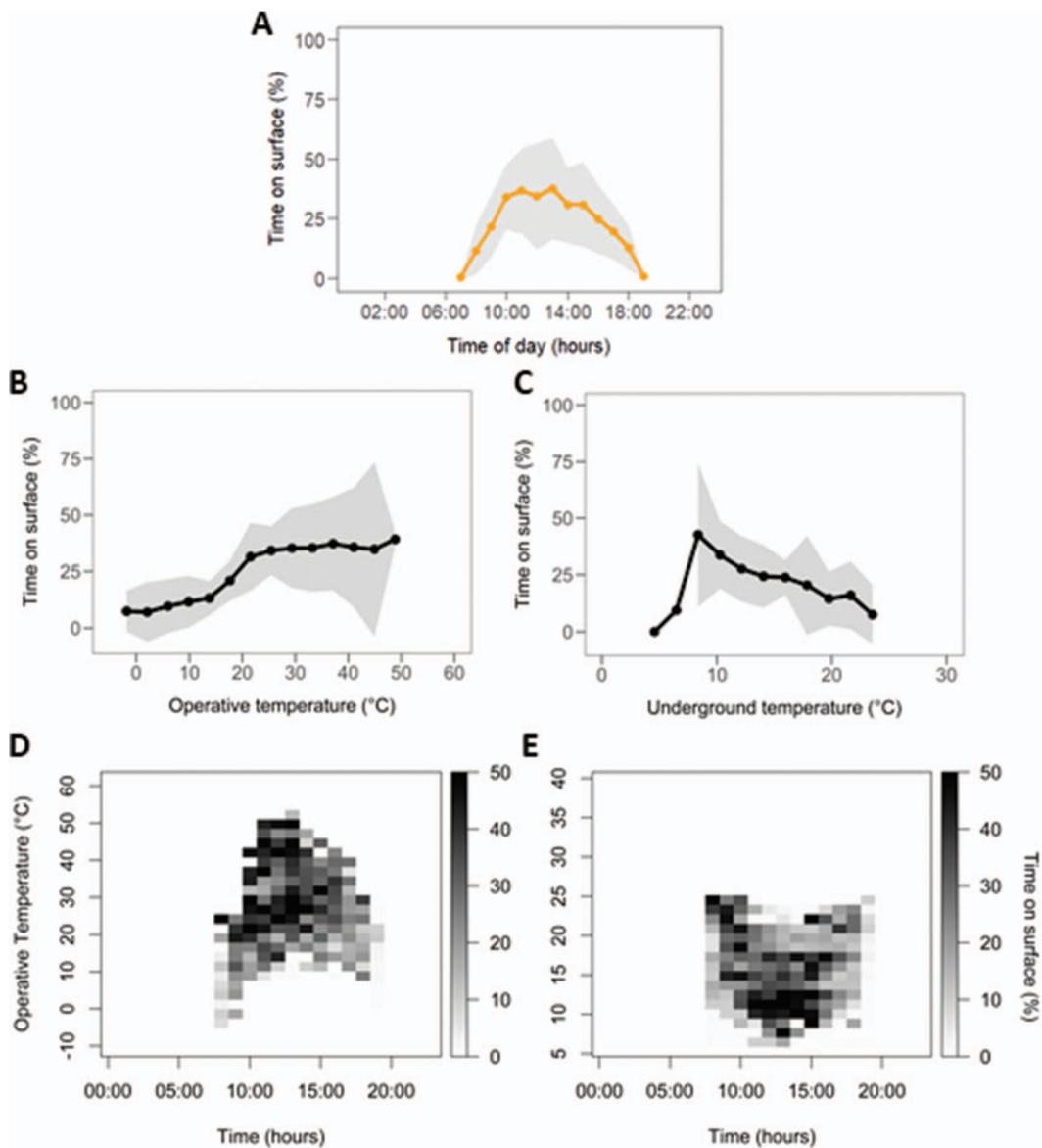


Figure 3: Average percentage of recordings of animals on surface when kept inside arenas in the field, from April to August of 2015–17, at each time of day (A), each operative temperature interval (B) or each underground temperature (at 20 cm) interval (C). Shaded area: standard deviation between individuals. (A) Sample size: 12 individuals; (B) Sample size for each point respectively: 1, 5, 10, 10, 10, 10, 10, 10, 10, 10, 10, 10, 9, 8, 8, 5, 2; (C) Sample size for each point respectively: 1, 1, 3, 7, 7, 8, 7, 9, 6, 5, 3. Simultaneous effect of time of day and environmental temperature is shown in (D) for operative temperature and (E) for underground temperature. Darker areas indicate intervals of temperature and time of day when percentage time on surface was higher.

(Tomotani *et al.*, 2012). However, they can stay up to 1 hour with only their heads out of the burrow entrance without doing any noticeable movement (M Jannetti, personal observation). While it also looks like a vigilant state, we hypothesize that this very common, immobile posture, with only the heads out of the burrow could be basking. If future studies using body temperature sensors confirm this hypothesis, we can conclude that thermoregulatory basking

is another important, winter behaviour of tuco-tucos on the surface, besides foraging and soil removal.

Based on the considerable amount of time tuco-tucos spend on the surface (Flôres *et al.*, 2016) and on the nature of their observed surface behaviours (Tomotani *et al.*, 2012), we had concluded previously that they are diurnal in the field. However, we had no knowledge of their activity timing and

Table 2: Comparison between generalized linear mixed models performed with environmental variables recorded, to explain variability of time on surface every 3 hours

Fixed effect variables	AIC	R ² (%)	χ^2
T_{und}, T_e, Hour, Wind	3600.3	59.9	512.6
T _{und} , T _e , Hour	3635.7	55.6	473.2
T _{und} , Tair, Hour	3683.7	57.5	425.2
T _{und} , Hour	3691.2	53.1	421.7
T _e , Hour	3721.4	59.4	391.5
Hour	3829.0	54.3	265.9
T _e	4006.0	31.1	84.9
T _{und}	4049.8	16.3	41.1
Wind	4067.5	11.3	23.4
Tair	4069.2	12.8	21.7

R² values consider the random effect variable, animal ID. Final model, in bold, is further discussed in the text. Number of observations: 633 from 10 animals. All models above were significant with P < 0.001.

behaviours underground. The combination of our light-logger and accelerometer data revealed that highest activity levels do in fact occur when tuco-tucos are on the surface. However, they also revealed that lower amplitude activity occurs underground during both the day and the night (Fig. 1). Because accelerometers can be used in both the field and lab, we then transferred three of the four individuals to the laboratory for continued monitoring of their activity timing under constant dark conditions. This was the first time we tracked a single activity parameter, the locomotor activity level from

ODBA from the same individuals under both captive and field conditions. Activity during both day and night was again detected in lab condition but, similar to our previous experiments using light-loggers in the field and wheel-running in the lab, accelerometry data confirmed the shift in highest activity time between field and lab conditions (Fig. 5). The night-time activity displayed by tuco-tucos under constant darkness in the lab represents the circadian clock controlled component (Hut *et al.*, 2011; Tomotani *et al.*, 2012) while the timing of field activity is defined downstream the clock (van der Vinne *et al.*, 2014). While the former depends mostly on the stable light/dark entrainment of the circadian clock (Flôres *et al.*, 2016), the latter is plastic and susceptible to prevailing environmental conditions, with higher activity levels during the day associated to economy in thermoregulatory costs (van der Vinne *et al.*, 2014, 2015). Finally, we verified that overall activity levels in the field is on average 1.6 times greater than in the lab, a discrepancy that has always been assumed to explain the temporal niche switches between field and lab conditions in rodents (Hut *et al.*, 2011; van der Vinne *et al.*, 2014).

Although higher levels of activity are concentrated during the day while in the field, accelerometry clearly revealed that tuco-tucos also display significant activity at night (40%) during the winter (Fig. 1). A similar pattern, but with a more evenly distributed activity bouts throughout day and night, has been observed in several primate species in the field (Erkert and Cramer, 2006). Erkert and Cramer (2006) have experimentally shown the circadian component behind this arrhythmic pattern and suggest that this may represent a transitory stage on the way from a nocturnal to a diurnal lifestyle. This is an interesting proposition, given the common

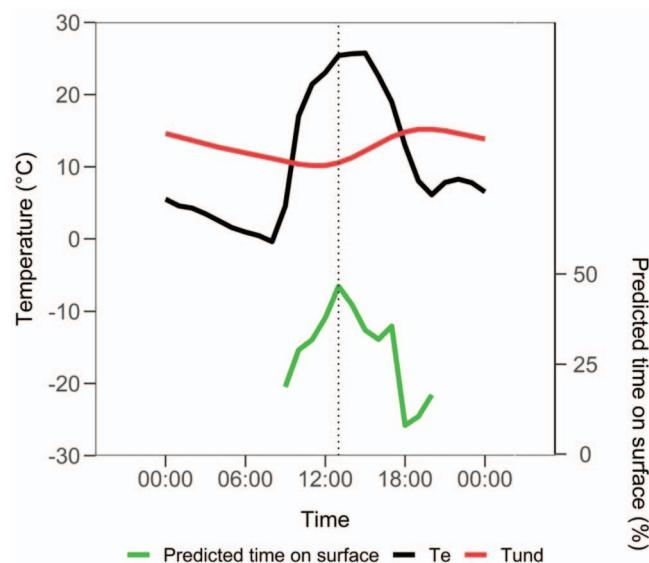


Figure 4: Predicted relationship of time on surface with T_e and T_{und} in tuco-tucos, based on the generalized linear model. This prediction was constructed with time of day, T_e, T_{und} and Wind values of an arbitrary day (15 July 2016), in Anillaco, La Rioja, Argentina. Green line: predicted time on surface; black line: T_e; red line: Tund. Wind speed variation was omitted for better clarity.

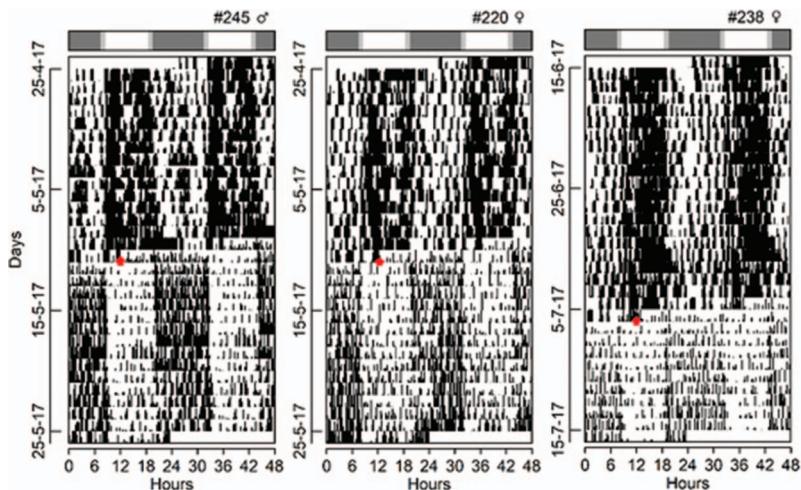


Figure 5: Double-plotted actogram of daily rhythms of gross motor activity of tuco-tucos kept inside arenas in the field and then released into constant darkness in the lab. Black marks: gross motor activity measured with accelerometers. Bar above actograms: natural photoperiod at which each recording started in the field (white: day; grey: night). Red arrows indicate day and time of transference. Top right of each actogram: animal identification number and sex. Because daily activity levels were lower in the lab than in the field, the upper limit of activity used in this actogram was lower than that used in Fig. 1, to allow visualization of activity/rest throughout the entire field/lab transition in a single actogram.

nocturnal ancestry of mammals (Gerkema *et al.*, 2013) and that the capacity of an organism to be active in either day or night, depending on prevailing environmental conditions, is viewed as adaptive in a changing environment.

The climate of South American deserts is changing, particularly with respect to increase in temperature and decrease in precipitation (Boulanger *et al.*, 2006; Labraga and Villalba, 2009). Desertification has become an enormous ecological problem, caused mainly by vegetation destruction, soil erosion and lack of water. In the desert of La Rioja province, Argentina, tuco-tucos play an important role in their environments, due to their burrowing habits, constantly altering the soils, dispersing diverse organic materials (Malizia *et al.*, 2000; Lara *et al.*, 2007), mycorrhizal fungi and endophytes, which form symbiotic relationships with the indigenous plant species (Fracchia *et al.*, 2011). However, their biology is poorly known and they live under constant human pressure since the local community encourages their extermination.

Climate projections suggest that vegetation cover and water availability will decrease under future climate scenarios (Therburg *et al.*, 2019); thus, limiting the ability of animals to remain active during the hotter day hours. The 24-hour time axis is an ecological resource that some animals can exploit to buffer impacts of climate change (Levy *et al.*, 2018). Plasticity of activity timing is seen in several other desert rodents (Levy *et al.*, 2007) and may be adaptive in changing environments. It is imperative to understand the capacity of animals to switch temporal niches, much as it is important to understand their capacity to migrate, when making predictions of how organisms will cope with these new climate-related challenges. Our data on biological timing

in tuco-tucos indicate a high degree of phenotypic plasticity in activity timing as evidenced by their capacity for temporal niche switching and both day and night activity.

Acknowledgements

The authors acknowledge Patricia Tachinardi for valuable discussions, Johanna Barros for excellent technical assistance, Tamiris Yassumoto, Giovane Improta and Danilo Flôres for all help with the experiments, Carlo Catoni from Technosmart for his kind assistance with his accelerometers, Natalia Yela for help with the meteorological station data, Cory Williams with kind help with accelerometry data analysis, Carlos Candia-Gallardo for his generous help with statistical analysis and the two anonymous referees for their constructive criticisms, suggestions and attention. MGJ, VSV performed experiments; MGJ analysed data; GAO, VSV, CLB conceived the experiments. MGJ, GAO, VSV, CLB wrote the paper.

Funding

V.S.V. was supported by FONCyT—Agencia Nacional de Promoción Científica y Tecnológica [grant PICT 2013/2753], CONICET—Consejo Nacional de Investigaciones Científicas y Técnicas [grant PIP-11220120100415CO]; CRILAR Institution was supported by CONICET—Consejo Nacional de Investigaciones Científicas y Técnicas [grant PUE-201622920160100125]; G.A.O. was supported by FAPESP—Fundação de Amparo à Pesquisa do Estado de São Paulo [grants 2014/20671-0, 2017/19680-2]; M.G.J. was supported

by scholarships [2014/09324-6, 2016/25058-0], Coordenação de Aperfeiçoamento de Pessoal de Nível Superior—Brazil (CAPES) [scholarship grant number 001].

Supplementary material

Supplementary material is available at *Conservation Physiology* online.

References

Abraham E, del Valle HF, Roig F, Torres L, Ares JO, Coronato F, Godagnone R (2009) Overview of the geography of the Monte Desert biome (Argentina). *J Arid Environ* 73: 144–153.

Acker JG, Leptoukh G (2007) Online analysis enhances use of NASA earth science data. *Eos* 88: 14–17.

Altuna CA, Francescoli G, Tassino B, Izquierdo G (1999) Ecoetología y Conservación de mamíferos subterráneos de distribución restringida: el Caso de *Ctenomys pearsoni* (Rodentia, Octodontidae) en el Uruguay. *Etología* 7: 47–54.

Antinuchi CD, Busch C (1992) Burrow structure in the subterranean rodent *Ctenomys talarum*. *Z Säugertierkd* 57: 163–168.

Aranda-Rickert A (2014) *Flora Del Parque Geológico Sanagasta*, Ed1st. Editorial Brujas, Córdoba, Argentina

Aschoff J, Daan S, Honma KI (1982) Zeitgebers, entrainment and masking: some unsettled questions. In J Aschoff, S Daan, G Groos, eds, *Vertebrate Circadian Systems*. Springer-Verlag, Berlin-Heidelberg

Bartholomew GA (1964) The roles of physiology and behaviour in the maintenance of homeostasis in the desert environment. *Symp Soc Exp Biol* 18: 7–29.

Beale AD, Whitmore D, Moran D (2016) Life in a dark biosphere: a review of circadian physiology in 'arrhythmic' environments. *J Comp Physiol B-Biochemical Syst Environ Physiol* 186: 947–968.

Ben-Shlomo R, Ritte U, Nevo E (1995) Activity pattern and rhythm in the subterranean mole-rat superspecies *Spalax ehrenbergi*. *Behav Genet* 25: 239–245.

Bennett AF, Huey RB, John-Alder H, Nagy KA (1984) The parasol tail and thermoregulatory behavior of the cape ground squirrel *Xerus inauris*. *Physiol Zool* 57: 57–62.

Bogert CM (1949) Thermoregulation in reptiles, a factor in evolution. *Evolution* 3: 195–211.

Borghesi CE, Giannoni SM, Roig VG (2002) Eye reduction in subterranean mammals and eye protective behavior in *Ctenomys*. *Mastozoología Neotrop* 9: 123–134.

Boulanger J-P, Martinez F, Segura EC (2006) Projection of future climate change conditions using IPCC simulations, neural networks and Bayesian statistics. Part 2: precipitation mean state and seasonal cycle in South America. *Clim Dyn* 28: 255–271.

Buffenstein R (2000) Ecophysiological responses of subterranean rodents to underground habitats. In EA Lacey, G Cameron, JL Patton, eds, *Life Underground: The Biology of Subterranean Rodents*. University of Chicago Press, Chicago, pp. 183–226.

Chappell MA, Bartholomew GA (1981) Standard operative temperatures and thermal energetics of the antelope ground squirrel *Ammospermophilus leucurus*. *Physiol Zool* 54: 81–93.

Chevin LM, Lande R, Mace GM (2010) Adaptation, plasticity, and extinction in a changing environment: towards a predictive theory. *PLoS Biol* 8: e1000357 10.1371/journal.pbio.1000357.

Cook JA, Lessa EP (1998) Are rates of diversification in subterranean south American tuco-tucos (genus *Ctenomys*, Rodentia: Octodontidae) unusually high? *Evolution (NY)* 52: 1521–1527.

Daan S (1981) Adaptive Daily Strategies in Behavior. In Aschoff J, ed., *Handbook of Behavioral Neurobiology*, Vol. 4. Biological Rhythms. Plenum Press, New York, pp 275–298.

Díez-Noguera A (2019) *El temps. Versión 1.292*. Universitat de Barcelona, Spain. URL <http://www.el-temps.com/>.

Dominoni DM, Åkesson S, Klaassen R, Spoelstra K, Bulla M (2017) Methods in field chronobiology. *Philos Trans R Soc B Biol Sci* 372.1734: 20160247. doi:10.1098/rstb.2016.0247

Erkert HG, Cramer B (2006) Chronobiological background to catemerality: circadian rhythms in *Eulemur fulvus albifrons* (Prosimii) and *Aotus azarae boliviensis* (Anthropoidea). *Folia Primatol* 77: 87–103.

Estevan I, Lacey EA, Tassino B (2016) Daily patterns of activity in free-living Rio Negro tuco-tucos (*Ctenomys rionegrensis*). *Mastozoología Neotrop* 23: 71–80.

Fick LG, Kucio TA, Fuller A, Matthee A, Mitchell D (2009) The relative roles of the parasol-like tail and burrow shuttling in thermoregulation of free-ranging cape ground squirrels, *Xerus inauris*. *Comp Biochem Physiol—A Mol Integr Physiol* 152: 334–340.

Flóres DEFL, Jannetti MG, Valentiniuzzi VS, Oda GA (2016) Entrainment of circadian rhythms to irregular light/dark cycles: a subterranean perspective. *Sci Rep* 6: 34264 10.1038/srep34264.

Flóres DEFL, Tomotani BM, Tachinardi P, Oda GA, Valentiniuzzi VS (2013) Modeling natural photic entrainment in a subterranean rodent (*Ctenomys* aff. *knighti*), the tuco-tuco. *PLoS One* 8: e68243. 10.1371/journal.pone.0068243.

Fox J, Monette G (1992) Generalized collinearity diagnostics. *J Am Stat Assoc* 87: 178–183.

Fracchia S, Krapovickas L, Aranda-Rickert A, Valentiniuzzi VS (2011) Dispersal of arbuscular mycorrhizal fungi and dark septate endophytes by *Ctenomys* cf. *knighti* (Rodentia) in the northern Monte Desert of Argentina. *J Arid Environ* 75: 1016–1023.

Geiser F, Goodship N, Pavey CR (2002) Was basking important in the evolution of mammalian endothermy? *Naturwissenschaften* 89: 412–414.

Gerkema MP, Davies WIL, Foster RG, Menaker M, Hut RA (2013) The nocturnal bottleneck and the evolution of activity patterns in mammals. *Proc R Soc B Biol Sci* 280: 20130508 10.1098/rspb.2013.0508.

Halle S, Stenseth NC (2000) *Activity Patterns in Small Mammals: An Ecological Approach*. Ecological Studies. Springer-Verlag, Berlin Heidelberg, New York.

Hut RA, Pilorz V, Boerema AS, Strijkstra AM, Daan S (2011) Working for food shifts nocturnal mouse activity into the day. *PLoS One* 6: e17527: 10.1371/journal.pone.0017527.

Kenagy GJ (1973) Daily and seasonal patterns of activity and energetics in a heteromyid rodent community. *Ecology* 54: 1201–1219.

Kenagy GJ, Nespolo RF, Vásquez RA, Bozinovic F (2002a) Daily and seasonal limits of time and temperature to activity of degus. *Rev Chil Hist Nat* 75: 567–581.

Kenagy GJ, Vásquez RA, Nespolo RF, Bozinovic F (2002b) A time-energy analysis of daytime surface activity in degus, *Octodon degus*. *Rev Chil Hist Nat* 75: 149–156.

Kronfeld-Schor N, Dayan T (2003) Partitioning of time as an ecological resource. *Annu Rev Ecol Evol Syst* 34: 153–181 10.1146/annurev.ecolsys.34.011802.132435.

Labraga JC, Villalba R (2009) Climate in the Monte Desert: past trends, present conditions, and future projections. *J Arid Environ* 73: 154–163.

Lacey EA, Patton JL, Cameron GN, eds (2000) *Life Underground: The Biology of Subterranean Rodents*. University of Chicago Press, Chicago.

Lane JE, Kruuk LEB, Charmantier A, Murie JO, Dobson FS (2012) Delayed phenology and reduced fitness associated with climate change in a wild hibernator. *Nature* 489: 554–557 10.1038/nature11335.

Lara N, Sassi P, Borghi C (2007) Effect of herbivory and disturbances by tuco-tucos (*Ctenomys mendocinus*) on a plant community in the southern Puna Desert. *Arct Antarct Alp Res*. 39: 110–116.

Levy O, Dayan T, Kronfeld-Schor N (2007) The relationship between the golden spiny mouse circadian system and its diurnal activity: an experimental field enclosures and laboratory study. *Chronobiol Int* 24: 599–613.

Levy O, Dayan T, Porter WP, Kronfeld-Schor N (2018) Time and ecological resilience: can diurnal animals compensate for climate change by shifting to nocturnal activity? *Ecol Monogr* 0: 1–21.

Long RA, Martin TJ, Barnes BM (2005) Body temperature and activity patterns in free-living arctic ground squirrels. *J Mammal* 86: 314–322.

Lovegrove BG, Muir A (1996) Circadian body temperature rhythms of the solitary cape mole rat *Georychus capensis* (Bathyergidae). *Physiol Behav* 60: 991–998.

Malizia AI, Kittlein MJ, Busch C (2000) Influence of the subterranean herbivorous rodent *Ctenomys talarum* on vegetation and soil. *Z Säugetierkd* 65: 172–182.

Marques MD, Waterhouse J (2004) Rhythms and ecology—do chronobiologists still remember nature? *Biol Rhythms Res* 35: 1–2.

Melcher JC, Armitage KB, Porter WP (1990) Thermal influences on the activity and energetics of yellow-bellied marmots (*Marmota flaviventris*). *Physiol Zool* 63: 803–820.

Nevo E, Guttman R, Haber M, Erez E (1982) Activity patterns of evolving mole rats. *J Mammal* 63: 453–463.

Oosthuizen MK, Cooper HM, Bennett NC (2003) Circadian rhythms of locomotor activity in solitary and social species of African mole-rats (Family: Bathyergidae). *J Biol Rhythms* 18: 481–490.

Parada A, D'Elía G, Bidau CJ, Lessa EP (2011) Species groups and the evolutionary diversification of tuco-tucos, genus *Ctenomys* (Rodentia: Ctenomyidae). *J Mammal* 92: 671–682.

R Core Team (2018) Version 3.5.1. R Foundation for Statistical Computing, Vienna, Austria. URL <https://www.R-project.org/>.

Rezende EL, Cortés A, Bacigalupo LD, Nespolo RF, Bozinovic F (2003) Ambient temperature limits above-ground activity of the subterranean rodent *Spalacopus cyanus*. *J Arid Environ* 55: 63–74.

Riccio AP, Goldman BD (2000) Circadian rhythms of locomotor activity in naked mole-rats (*Heterocephalus glaber*). *Physiol Behav* 71: 1–13.

Root T (1988) Energy constraints on avian distributions and abundances. *Ecology* 69: 330–339.

Root TL, Price JT, Hall KR, Schneider SH, Rosenzweig C, Pounds JA (2003) Fingerprints of global warming on wild animals and plants. *Nature* 421: 57–60 10.1038/nature01333.

Rusak B (1982) Physiological models of the rodent circadian system. In J Aschoff, S Daan, G Groos, eds, *Vertebrate Circadian Systems*. Springer-Verlag, Berlin-Heidelberg.

Ruxton GD (2006) The unequal variance t-test is an underused alternative to Student's t-test and the Mann-Whitney U test. *Behav Ecol* 17: 688–690.

Schwaibold U, Pillay N (2006) Behavioral strategies of the African ice rat *Otomys sloggetti robertsi* in the cold. *Physiol Behav* 88: 567–574.

Shen M, Piao S, Cong N, Zhang G, Jassens IA (2015) Precipitation impacts on vegetation spring phenology on the Tibetan plateau. *Glob Chang Biol* 21: 3647–3656 10.1111/gcb.12961.

Sikes RS (2016) 2016 Guidelines of the American Society of Mammalogists for the use of wild mammals in research and education. *J Mammal* 97: 663–688.

Sinervo B et al. (2010) Erosion of lizard diversity by climate change and altered thermal niches. *Science* (80-) 328: 894–899.

Skliba J, Sumbera R, Chitaukali WN, Burda H (2007) Determinants of daily activity patterns in a free-living afrotropical solitary subterranean rodent. *J Mammal* 88: 1009–1016.

Tachinardi P, Bicudo JEW, Oda GA, Valentimuzzi VS (2014) Rhythmic 24 h variation of core body temperature and locomotor activity in a subterranean rodent (*Ctenomys* aff. *knighti*), the tuco-tuco. *PLoS One* 9: e85674 10.1371/journal.pone.0085674.

Tachinardi P, Valentinuzzi VS, Oda GA, Buck CL (2017) The interplay of energy balance and daily timing of activity in a subterranean rodent: a laboratory and field approach. *Physiol Biochem Zool* 90: 546–552.

Therburg A, Corso ML, Stamat M, Bottero C, Lizana P, Pietragalla et V (eds) (2019) *Síntesis de resultados de la evaluación de la degradación de tierras: 2012–2017*, Ed1st. IADIZA, Mendoza.

Time and Date AS (2018) *Sunrise and sunset calculator*. <https://www.timeanddate.com/sun/> (last accessed 9 October 2018).

Tomotani BM, Flôres DEFL, Tachinardi P, Paliza JD, Oda GA, Valentinuzzi VS (2012) Field and laboratory studies provide insights into the meaning of day-time activity in a subterranean rodent (*Ctenomys* aff. *knighti*), the tuco-tuco. *PLoS One* 7: e37918 10.1371/journal.pone.0037918.

Valentinuzzi VS, Oda GA, Araujo JF, Ralph MR (2009) Circadian pattern of wheel-running activity of a south American subterranean rodent (*Ctenomys* cf *knighti*). *Chronobiol Int* 26: 14–27.

van der Vinne V, Gorter JA, Riede SJ, Hüt RA (2015) Diurnality as an energy-saving strategy: energetic consequences of temporal niche switching in small mammals. *J Exp Biol* 218: 2585–2593.

van der Vinne V, Riede SJ, Gorter JA, Eijer WG, Sellix MT, Menaker M, Daan S, Pilorz V, Hüt RA (2014) Cold and hunger induce diurnality in a nocturnal mammal. *Proc Natl Acad Sci* 111: 15256–15260.

Vassallo AI (1998) Functional morphology, comparative behaviour, and adaptation in two sympatric subterranean rodents genus *Ctenomys* (Caviomorpha: Octodontidae). *J Zool* 244: 415–427.

Vlasatá T, Sklíba J, Lovy M, Meheretu Y, Sillero-Zubiri C, Sumbera R (2017) Daily activity patterns in the giant root rat (*Tachyoryctes macrocephalus*), a fossorial rodent from the afro-alpine zone of the Bale Mountains, Ethiopia. *J Zool* 302: 157–163.

Warnecke L, Turner JM, Geiser F (2008) Torpor and basking in a small arid zone marsupial. *Naturwissenschaften* 95: 73–78.

Williams CT, Buck CL, Sheriff MJ, Richter MM, Krause JS, Barnes BM (2017) Sex-dependent Phenological plasticity in an Arctic hibernator. *Am Nat* 190: 854–859.

Williams CT, Wilsterman K, Kelley AD, Breton AR, Stark H, Humphries MM, McAdam AG, Barnes BM, Boutin S, Buck CL (2014) Light loggers reveal weather-driven changes in the daily activity patterns of arboreal and semifossorial rodents. *J Mammal* 95: 1230–1239 10.1644/14-MAMM-A-062.

Williams CT, Wilsterman K, Zhang V, Moore J, Barnes BM, Buck CL (2016) The secret life of ground squirrels: accelerometry reveals sex-dependent plasticity in above-ground activity. *R Soc Open Sci* 3: 160404 10.1098/rsos.160404.

Zuur AF, Ieno EN, Walker N, Saveliev AA, Smith GM (2009) *Mixed Effects Models and Extensions in Ecology with R*. Springer New York, New York, NY