






# Patterns of activity and body temperature of Aldabra giant tortoises in relation to environmental temperature

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## Abstract

We studied the temperature relations of wild and zoo Aldabra giant tortoises (*Aldabrachelys gigantea*) focusing on (1) the relationship between environmental temperature and tortoise activity patterns ( $n = 8$  wild individuals) and (2) on tortoise body temperature fluctuations, including how their core and external body temperatures vary in relation to different environmental temperature ranges (seasons;  $n = 4$  wild and  $n = 5$  zoo individuals). In addition, we surveyed the literature to review the effect of body mass on core body temperature range in relation to environmental temperature in the Testudinidae. Diurnal activity of tortoises was bimodally distributed and influenced by environmental temperature and season. The mean air temperature at which activity is maximized was 27.9°C, with a range of 25.8–31.7°C. Furthermore, air temperature explained changes in the core body temperature better than did mass, and only during the coldest trial, did tortoises with higher mass show more stable temperatures. Our results, together with the overall Testudinidae overview, suggest that, once variation in environmental temperature has been taken into account, there is little effect of mass on the temperature stability of tortoises. Moreover, the presence of thermal inertia in an individual tortoise depends on the environmental temperatures, and we found no evidence for inertial homeothermy. Finally, patterns of core and external body temperatures in comparison with environmental temperatures suggest that Aldabra giant tortoises act as mixed conformer–regulators. Our study provides a baseline to manage the thermal environment of wild and rewilded populations of an important island ecosystem engineer species in an era of climate change.

## KEYWORDS

Aldabra, ectotherm, giant tortoise, Testudinidae, thermoregulation

## 1 | INTRODUCTION

Activity and body temperature of reptiles depend on the external thermal fluctuations in the environment and are both drivers and

consequences of their physiological and behavioral biology, which ultimately affects their ecology (Heatwole, 1976; Huey & Stevenson, 1979; Lailvaux & Irschick, 2007; Van Damme, Bauwens, & Verheyen, 1991). Many physical processes can affect the thermal environment

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of reptiles, including for example fluxes of radiative heat, convection, conduction, and wind (Cossins & Bowler, 1987; Willmer, Stone, & Johnston, 2005). However, understanding this complex thermal environment alone does not allow an adequate description of a reptiles' activity patterns and core body temperature ( $T_{bc}$ ; Table 1 lists the terms adopted here and their definitions). Rather than being thermally passive, that is, with a body temperature driven only by fluctuations in environmental temperature, many reptiles have been shown to exhibit complex thermoregulatory behaviors and physiological processes to maintain their  $T_{bc}$  within a narrow range, albeit within limits determined by environmental conditions (Paladino, O'Connor, & Spotila, 1990; Seebacher & Franklin, 2001; Slip & Shine, 1988).

For example, when basking in sunny places, the common Puerto Rican ameiva (*Ameiva exsul*) can attain a  $T_{bc}$  that is higher than the air temperature ( $T_{air}$ ), which allows them to be active later while foraging in the shade (Rivera-Vélez & Lewis, 1994). In addition, salt water crocodiles (*Crocodylus porosus*) can employ shuttling behavior between water and land to regulate their  $T_{bc}$  (Seebacher, Grigg, & Beard, 1999). On the other hand, eastern bearded dragons (*Amphibolurus barbatus*) can exert physiological reactions of metabolism and circulation in response to varying  $T_a$  to control heating and cooling rates and maintain preferred  $T_{bc}$  (Bartholomew & Tucker, 1963). Furthermore, green iguanas (*Iguana iguana*) can display physiologically generated circadian rhythms in a constant environmental temperature, similar to those recorded in endotherms (Tosini & Menaker, 1995).

Another factor considered important in influencing fluctuations of  $T_{bc}$  in reptiles is their body size, as the surface-to-volume ratio influences the rate at which heat is exchanged with the environment. Hence, due to a relatively small surface-to-volume ratio for larger reptiles, their  $T_{bc}$  is expected to be less responsive to the thermal environment than that of smaller ones (i.e., more stable; Zimmerman & Tracy, 1989). Body size also affects the rate of heat absorption from the sun (Brattstrom, 1965), as larger bodies have a higher surface area,

which in part drives their external body temperature ( $T_{be}$ ). McNab and Auffenberg (1976) suggested that larger reptiles have a relatively low thermal conductance because their small surface-to-volume ratio and thick integuments could give them a substantial heat storage capacity, and because larger reptiles take longer to attain thermal equilibrium with the environment. For example, larger salt water crocodiles are able to attain not only higher but also more stable  $T_{bc}$  than smaller ones, at least in part due to thermal inertia (Seebacher et al., 1999).

Here, we describe activity and body temperature fluctuations of Aldabra giant tortoises (*Aldabrachelys gigantea* Schweigger 1812) in both their natural habitat and in captivity. Specifically, we focus on (1) the relationship between environmental temperature ( $T_a$ ) and activity patterns of wild tortoises to determine their optimal environmental temperature range ( $T_{a-opt}$ ); and (2) the body temperature fluctuations of captive and wild tortoises, including how their core and external body temperatures vary in relation to environmental temperatures, and whether body mass influences the response of tortoise core body temperatures to environmental temperatures. In addition, we surveyed the literature to investigate the effect of body mass on the body temperature ranges of Testudinidae in relation to air temperature.

## 2 | MATERIALS AND METHODS

### 2.1 | Study species and study sites

The Aldabra giant tortoise is endemic to Aldabra Atoll, Seychelles, with a stable population of an estimated 100,000 tortoises (Bourn et al., 1999; Turnbull et al., 2015). We studied wild tortoises on Picard Island, Aldabra, as well as captive ones in the Masoala Rainforest exhibit at Zurich Zoo, Switzerland. Aldabra is a raised coral atoll in the Western Indian Ocean and has a tropical climate with a wet season (November to April) and a dry season (May to October). The timing and distribution of rainfall vary greatly from year to year, which is unevenly

Term	Definition
$T_a$	Available environmental temperature envelope (e.g., available $T_{air}$ and $T_{sun}$ in our case) for temperature regulation (e.g., throughout a given time period or study; °C)
$T_{a\ mean}$	Mean environmental temperature (from $T_{air}$ and $T_{sun}$ ; °C)
$T_{air}$	Temperature recorded by loggers placed in a shaded area used by tortoises (equivalent to air temperature); and air temperature reported in studies for the review section; °C
$\Delta T_{air}$	Environmental temperature range given by the minimum and maximum temperatures recorded over a certain period from loggers placed in the shade (range in °C)
$T_{a-opt}$	Optimum mean air temperature at which activity is maximized (°C)
$T_{a-opt\ range}$	Optimum air temperature range given by the minimum and maximum temperatures at which the active state surpasses the inactive state of tortoises (°C)
$T_{bc}$	Core body temperature (i.e., gut temperature; °C)
$\Delta T_{bc}$	Core body temperature range given by the minimum and maximum temperatures recorded over a certain period (°C)
$T_{be}$	External (contact) body temperature (i.e., carapace, extremities and skin fold; °C)
$T_{sun}$	Temperature recorded by loggers placed in a sun-exposed area used by tortoises (includes radiative temperature; °C)

**TABLE 1** Terms related to thermoregulation ecology used in this article

Definitions based and modified from Cossins and Bowler (1987), Blatteis et al. (2001) and Willmer et al. (2005).

distributed across the atoll, and directly drive spatiotemporal patterns in vegetation productivity (Haverkamp et al., 2017; Shekeine et al., 2015). In Zurich Zoo, the Masoala Rainforest exhibit is an 11,000-m<sup>2</sup> greenhouse ecosystem covered with translucent foil permitting 75% of daylight and 50% of UV radiation to enter and has an artificial rain and fog systems and an air-circulating heating system with the aims of maintaining minimum temperatures of 24° during the day and 18°C during the night (Bauert, Furrer, Zingg, & Steinmetz, 2007).

## 2.2 | Environmental temperatures and tortoise activity patterns

The environmental temperature range at which activities such as locomotion, feeding, and mating occur in tortoises is narrow (Meek, 1984). As other reptiles, tortoises have to select environmental temperatures that allow them to maintain  $T_{bc}$ s at which manifold processes are optimized to increase their fitness, for example physiological, locomotory, and foraging ones. We examined activity patterns of wild tortoises in relation to air temperature ( $T_{air}$ ) to derive an approximation for the optimal environmental temperature at which activity is maximized ( $T_{a-opt}$ ). We derived activity based on accelerometer (ACC) data obtained from data loggers mounted on eight tortoises (06:00–24:00) for 2 years, as part of an ongoing long-term movement ecology study (we assumed no activity between 00:00 and 06:00). ACC data were recorded every 5 min for a burst of 5-s, during which 36 voltage readings were recorded. To assign a state of either active (1) or inactive (0) for/within each 5-min period, a rolling mean of the standard error (SE) of the ACC bursts was used to capture fluctuations in the ACC waveform. We set the threshold of the rolling average SE to five, to create binary data (similar to Nielsen, Pedersen, Herskin, & Munksgaard, 2010). The activity data were then coupled with  $T_{air}$  obtained from the weather station located at the research station on Picard, which was collected every 15 min.

## 2.3 | Body temperature of *Aldabrachelys gigantea*

We studied body temperature fluctuations in five captive tortoises from Zurich Zoo and four wild tortoises from Aldabra. The zoo tortoises were housed in a compartment within the Masoala Rainforest exhibit, while the wild tortoises, selected for their different body masses (Table 2), were temporarily housed in two 36 m<sup>2</sup> enclosures located on Picard. Data from the captive tortoises were obtained during summer 2009 (ZRH summer;  $n = 3$ ) and winter 2010 (ZRH winter;  $n = 5$ ), while data from the wild tortoises were obtained during April 2014 (Aldabra;  $n = 4$ ). In the course of the trials, all animals had ad libitum access to drinking water, food and access to both shade and sunlight. Food consisted of hay, freshly cut grass, and vegetables for the captive tortoises, and freshly cut leaves from native woody vegetation of known food species for the wild tortoises.

Environmental temperatures were recorded with temperature loggers every 15 min for all study periods. In Zurich Zoo, we used 11-bit Thermocron HC temperature loggers ( $\pm 0.06^\circ\text{C}$  accuracy; OnSolution Pty Ltd., NSW, Australia), while on Aldabra, we used iButton®

temperature loggers ( $\pm 0.05^\circ\text{C}$  accuracy; Maxim Integrated, San Jose, CA, USA). In both sites, loggers were placed at two locations (shaded and directly exposed to sunlight) at a height of 0.3–0.5 m. Surface (external) body temperatures ( $T_{be}$ ) were measured using infrared temperature pistols: *Raytek Fluke 566* in Zurich Zoo (0.01°C accuracy; Raytek Corporation, Santa Cruz, USA) and *testo 810* on Aldabra (0.1°C accuracy; Testo SA & Co., Mönchaltorf, Switzerland). To test whether logger and pistol data could be reliably compared, logger temperatures were directly measured using the infrared temperature pistol in Zurich Zoo, and data were highly correlated ( $T_{air}$ :  $z = 150.68$ ,  $p < .001$ , Kendall  $\tau = 0.89$ ;  $T_{sun}$ :  $z = 84.05$ ,  $p < .001$ , Kendall  $\tau = 0.89$ ). We fed temperature loggers to the tortoises and recorded their internal temperature ( $T_{bc}$ ) every 15 min. Feces were examined daily for the loggers, which were voided 10–20 days after ingestion. In Zurich Zoo, three tortoises (100–180 kg; see Table 2 for individual body masses) were fed the data loggers during the summer trial and five tortoises for the winter trial (14–180 kg; same three individuals as in summer, plus two additional ones). On Aldabra, four tortoises (39–97 kg) were monitored. We additionally measured the following temperatures at 1–3-hr intervals for a 48–72-hr period, commencing 5 days after feeding the loggers to the tortoises: (1) surface of the carapace (the center of each of the 13 main scutes, vertebral and costal), (2) the four extremities (each extremity in the region of the metacarpal/metatarsal joints), and (3) the deep skin folds (the skin at the deepest point underneath the carapace between each fore extremity and the neck, and next to each hind extremity; i.e., four measurements).

## 2.4 | Body size and temperature in testudinidae

To investigate thermal inertia, and inertial homeothermy, we collated data on  $\Delta T_{bc}$ ,  $\Delta T_{air}$ , and on body mass in Testudinidae from the scientific literature. We searched the literature and selected studies that presented the aforementioned data for at least one Testudinid species, or summarized mean values for a group of individuals. When studies only showed results graphically, we extracted the data from figures using WebPlotDigitizer v. 3.10 (Rohatgi, 2017; <http://arohatgi.info/WebPlotDigitizer>). We then assessed whether  $\Delta T_{air}$  and/or mass significantly influenced  $\Delta T_{bc}$  using correlation analyses. We restricted our assessment to Testudinidae, rather than Testudines, as oceanic/aquatic and semiaquatic species are subject to different physical processes than terrestrial species (i.e., dissimilarities in heat dissipation caused by differences in convection and conduction properties of air and water).

## 2.5 | Statistical analyses

We performed all statistical analyses using R v. 3.3.0 (R Core Team, 2016) and report mean values and standard deviations ( $\pm$ SD). When plotting data, we fitted nonparametric locally weighted regressions using the nearest neighbor approach (loess; with t-based approximation 95% CI), using the package “ggplot2” (Wickham, 2016). We determined the temperature range at which tortoises maximize their activity ( $T_{a-opt}$ ) by performing kernel density estimation. We partitioned

**TABLE 2** Summary statistics of the environmental ( $T_{\text{air}}$  and  $T_{\text{sun}}$ ) and core body temperatures ( $T_{\text{bc}}$ ) of Aldabra giant tortoises (*Aldabrachelys gigantea*), with different body mass, exposed to different thermal environments. Temperatures are indicated in degrees celsius, and the acrophase is indicated in hours. The mean, maximum, and minimum temperatures refer to the overall temperatures during the trials, while the range refers to the mean daily temperature range

Environmental temperature		Study	Mean	$\pm$ SD	Min	Max	Daily range	Acrophase
Sun		ZRH Winter	18.0	3.0	14.4	32.1	10.3	14.0
Air (shade)		ZRH Winter	16.5	2.1	13.9	23.7	6.4	14.9
Mean		ZRH Winter	17.2	2.5	14.2	27.4	8.2	14.4
Sun		ZRH Summer	25.4	5.4	18.2	36.6	15.3	14.5
Air (shade)		ZRH Summer	22.0	3.1	17.1	28.6	8.7	15.3
Mean		ZRH Summer	23.7	4.2	17.7	32.6	11.8	14.7
Sun		Aldabra	31.6	7.3	23.0	56.5	18.5	14.0
Air (shade)		Aldabra	29.1	2.5	24.0	38.5	6.5	15.9
Mean		Aldabra	30.3	4.4	23.8	43.8	12.4	14.5
Tortoise	Mass (kg)	Study	Mean	$\pm$ SD	Min	Max	Daily range	Acrophase
JVS	14	ZRH Winter	20.8	1.6	17.0	26.0	3.1	19.04
JVL	19	ZRH Winter	21.5	1.3	18.6	25.5	2.7	20.26
HMA	100	ZRH Winter	22.2	0.9	20.1	24.6	1.4	20.98
SBY	140	ZRH Winter	21.1	1.0	19.2	23.9	1.6	20.28
BBY	180	ZRH Winter	20.5	0.8	18.4	23.0	1.5	21.53
HMA	100	ZRH Summer	29.9	2.1	25.1	34.1	5.0	19.63
SBY	140	ZRH Summer	30.1	2.4	24.7	34.2	5.7	20.03
BBY	180	ZRH Summer	29.7	2.0	24.2	33.2	4.7	20.57
BEL	39	Aldabra	29.9	1.6	26.0	34.5	4.7	18.13
UNM	48	Aldabra	29.9	1.7	26.0	34.0	4.9	17.86
CFK	61	Aldabra	30.2	1.6	26.5	35.0	4.4	18.34
LXD	97	Aldabra	31.0	1.7	26.0	34.5	5.0	17.86

the activity data in active and inactive states at a given temperature, which yielded a relative density distribution for each state with the area under the curve of the probability distributions adding to one. We visualized the kernel density estimation by partitioning the active and inactive states using R package “ggplot2” and expected the distribution of the activity probability in response to environmental temperature to show a bimodal distribution for the inactive state, with the active state exhibiting greater levels of activity in between. In addition to calculating the  $T_{\text{bc}}$  ranges of the Aldabra giant tortoises and that of  $T_{\text{air}}$  and  $T_{\text{sun}}$ , we also fitted cosines of the angles to the observed  $T_{\text{bc}}$  data with circular-linear regression (e.g., Jammalamadaka & Lund, 2006; Kinahan, Inge-moller, Bateman, Kotze, & Scantlebury, 2007) using the R package “psych” (Revelle, 2016) and calculated the acrophase (time period during which the peak of  $T_{\text{bc}}$  occurs).

We used the package “lme4” (Bates, Machler, Bolker, & Walker, 2015) to construct generalized linear mixed effects models with random factors (GLMMs) following Zuur, Ieno, Walker, Saveliev, and Smith (2009). For the activity data, we tested the effects of *year*, *time*, *season* (wet and dry),  $T_{\text{air}}$ , and the interactions between *time* and *season*, and  $T_{\text{air}}$  and *season* on tortoise *activity* (active or inactive state; using logistic regression analysis with binomial family and link “log”; from here on “activity model”). We added *individual* (tortoise

and *day* (date) as random factors to account for individual variation and repeated measures. The analysis was limited to 06:00–20:00, and the wet season comprised the months November–April and the dry season May–October. To account for the nonlinear relationship between *activity* and *time*, we discretised continuous time into four periods (I–IV), following the overall activity turning points through time, and comprising 06:00–08:00, 08:15–13:30, 13:45–17:30, and 17:45–20:00, respectively (see S1 in Supplementary Information). Furthermore, to assess the factors that influence  $T_{\text{bc}}$  of giant tortoises during our trials, we tested the following explanatory variables: *trial*, *time*,  $T_{\text{air}}$ , *mass*, and the interaction between  $T_{\text{air}}$  and *mass* (from here on “thermoregulation model”). We included a random factor with individual tortoises interacting with *trial* and a random factor with *day* (date) to account for individual variation and repeated measures among trials. Because they are correlated,  $T_{\text{air}}$  was selected over  $T_{\text{sun}}$  and  $T_{\text{a mean}}$  as an explanatory variable based on model selection ( $\Delta$ AIC). Moreover, and similar to the activity model, we discretised continuous time into three periods (I–III) following the turning points of  $T_{\text{bc}}$  through time for each independent trial and comprising the morning period when tortoises are cooling down, the morning–afternoon period when tortoises are heating up, and the night period when tortoises start to cool down, respectively,

to account for the nonlinear relationship between  $T_{bc}$  and time (see S2 in Supplementary Information). We obtained  $p$ -values for the explanatory variables using the Satterthwaite degrees of freedom approximation (implemented in package “lmerTest” for the thermo-regulation model; Kuznetsova, Brockhoff, & Christensen, 2016).

### 3 | RESULTS

#### 3.1 | Environmental temperatures and tortoise activity patterns

Diurnal activity of Aldabra giant tortoises on Aldabra was bimodally distributed, with the highest activity levels occurring during the morning, and during the late afternoon, albeit at comparatively lower levels (Figure 1). The mean  $T_{air}$  during the active state was 27.9°C ( $\pm 2.6$ ; 25%–75% quartile = 26.0–29.6°C). Notably, the probability of activity rapidly decreased as the environmental temperature increased above ca. 31–32°C (Figure 2a). Moreover, the kernel frequency distribution of the active state highlights that the temperature range of the active state in tortoises is 25.8–31.7°C (Figure 2b). In the activity model, the activity patterns of tortoises were significantly influenced by  $T_{air}$ , as well as time, season, and the interactions between  $T_{air}$  and time, and  $T_{air}$  and season ( $p < .001$ ), but not by year ( $p = .87$ ; see S3 in Supplementary Information for model statistics and S4 for seasonality plot).

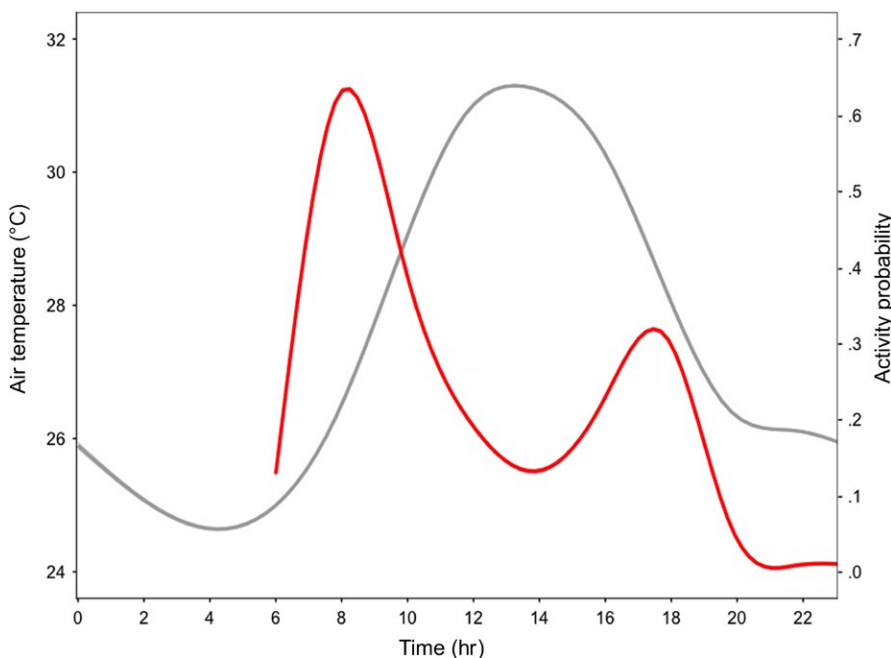
#### 3.2 | Body temperature of *Aldabrachelys gigantea*

Temperatures measured in shaded and sunny areas in each of the trials had similar daily minima but different maxima (Table 2), with considerable day-to-day variation in all trials (Figure 3). On Aldabra, we recorded relatively higher temperatures, in some cases above 40°C, in the sunny areas. The general daily patterns of  $T_{bc}$  lagged behind those of  $T_a$ ,

Tortoises on Aldabra seemed to behave as thermoconformers at the beginning of the trial (first 5 days, where the  $T_{bc}$  of tortoises followed  $T_a$  closely). At the beginning of the ZRH summer trial, the 100-kg individual seems to have avoided basking in the sun. During the trials, there were some perturbations in the  $T_a$  cycles, and the ability of tortoises to cope with these depended on the direction of the perturbation. Tortoises were able to maintain a stable  $T_{bc}$  when  $T_a$  increased above normal levels (e.g., Aldabra) but had difficulties doing so when  $T_a$  decreased sharply (e.g., ZRH summer and the last day at Aldabra). Overall, tortoises were able to maintain their  $T_{bc}$  above low mean  $T_a$  and below high mean  $T_a$ . Tortoises maintained their mean  $T_{bc}$  at 30.1°C  $\pm$  1.9 during the ZRH summer and Aldabra trials (for all tortoises combined,  $n = 7$ ; Table 2). In contrast, during the ZRH winter trial, tortoises maintained a mean  $T_{bc}$  of 21.2°C  $\pm$  1.3, albeit higher than mean  $T_a$ . The mean daily  $T_{bc}$  of tortoises varied by 3.7°C  $\pm$  0.9 on Aldabra, 4.7°C  $\pm$  1.2 during ZRH summer, and 4.9°C  $\pm$  1.0 during ZRH winter. Moreover, only during the winter trial, where juveniles were included, did we observe that the rate of heat loss during the night increased as mass decreased (Figure 3). On the other hand, only on Aldabra when  $T_a$  was above 25°C did the  $T_{bc}$  of tortoises reach values very close to the minimum  $T_a$ .

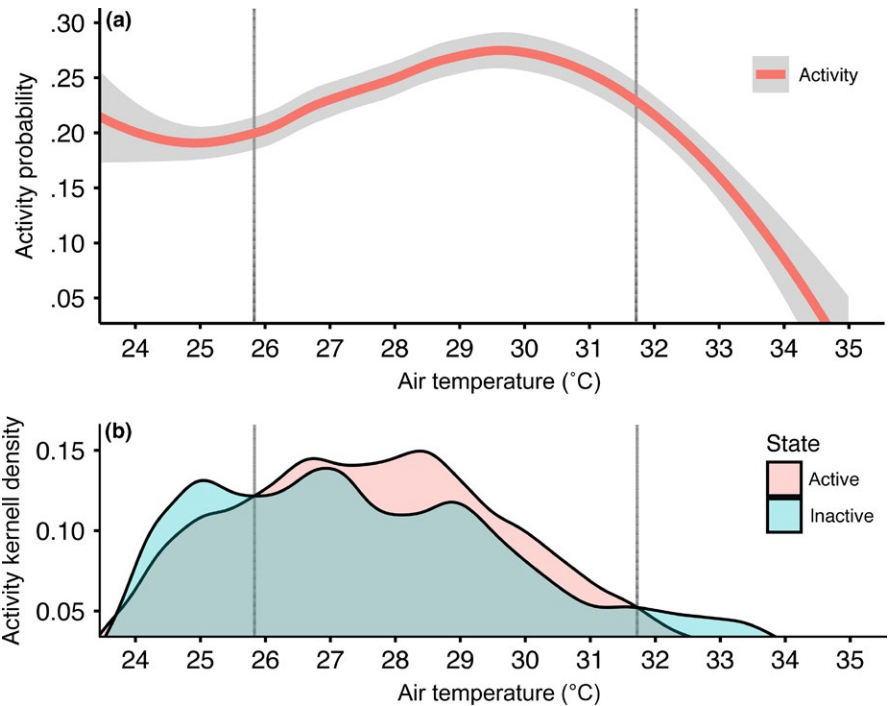
Viewing the aggregated variability in the tortoises'  $T_{bc}$  over a 24-hr period, the variable dependence of the effects of mass on  $T_a$  (due to different  $T_a$  ranges available) and their influence on  $T_{bc}$  became clear (Figure 4). During winter, the magnitude of the response of  $T_{bc}$  to daily changes in  $T_a$  decreased with size, and only the smallest tortoises were able to briefly reach  $T_{bc}$  close to  $T_{a-opt}$ . However, as  $T_{a-mean}$  reached values and temperature ranges closer to  $T_{a-opt}$ , larger tortoises were able to increase and maintain their mean  $T_{bc}$  close to the upper range of  $T_{a-opt}$  (i.e., ZRH summer and Aldabra trials). In general, as  $T_{a-mean}$  and minimum  $T_a$  increased, so did the  $T_{bc}$  of the tortoises (Table 2), but most of  $T_{bc}$  readings remained above  $T_{a-mean}$ .

The magnitude of daily  $\Delta T_{bc}$  differed among trials (Table 2). Overall, there was a positive correlation between the daily  $\Delta T_{bc}$  and



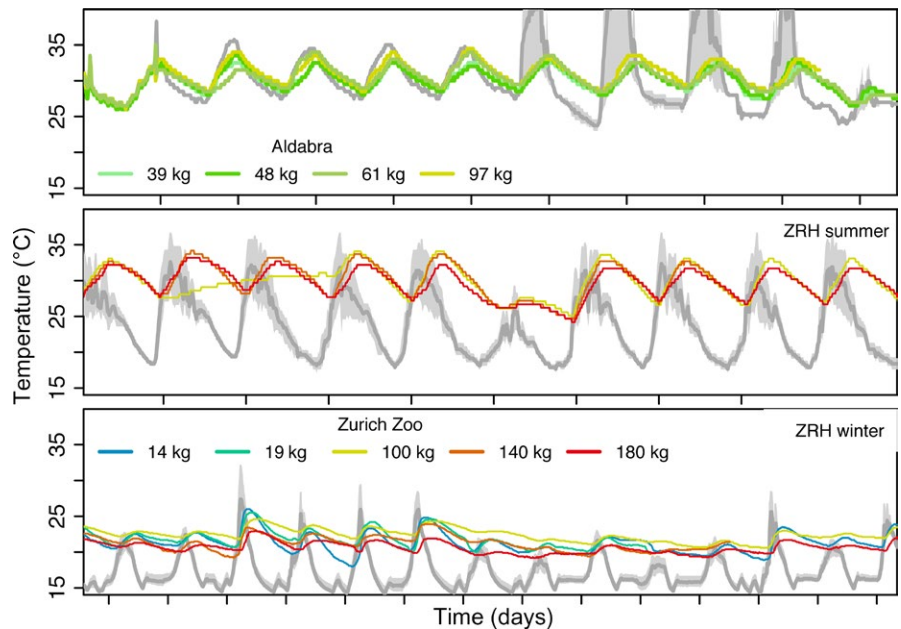
**FIGURE 1** Bimodal distribution of the probability of activity (red line) of wild Aldabra giant tortoises (*Aldabrachelys gigantea*) and environmental temperature (gray line) per hr/day. Line smoothing by local regression loess





**FIGURE 2** Probability of Aldabra giant tortoise (*Aldabrachelys gigantea*) activity as a response to air temperature ( $T_{\text{air}}$ ) (a) and kernel density plot of active and inactive states for a given  $T_{\text{air}}$  (b). Line smoothing by local regression loess (shading indicates the 95% CI). Vertical lines delimit the temperature range at which activity is maximised.  $T_{\text{air}}$  was recorded from Picard Island weather station daily along with tortoise activity data from eight individuals during 2013–2014

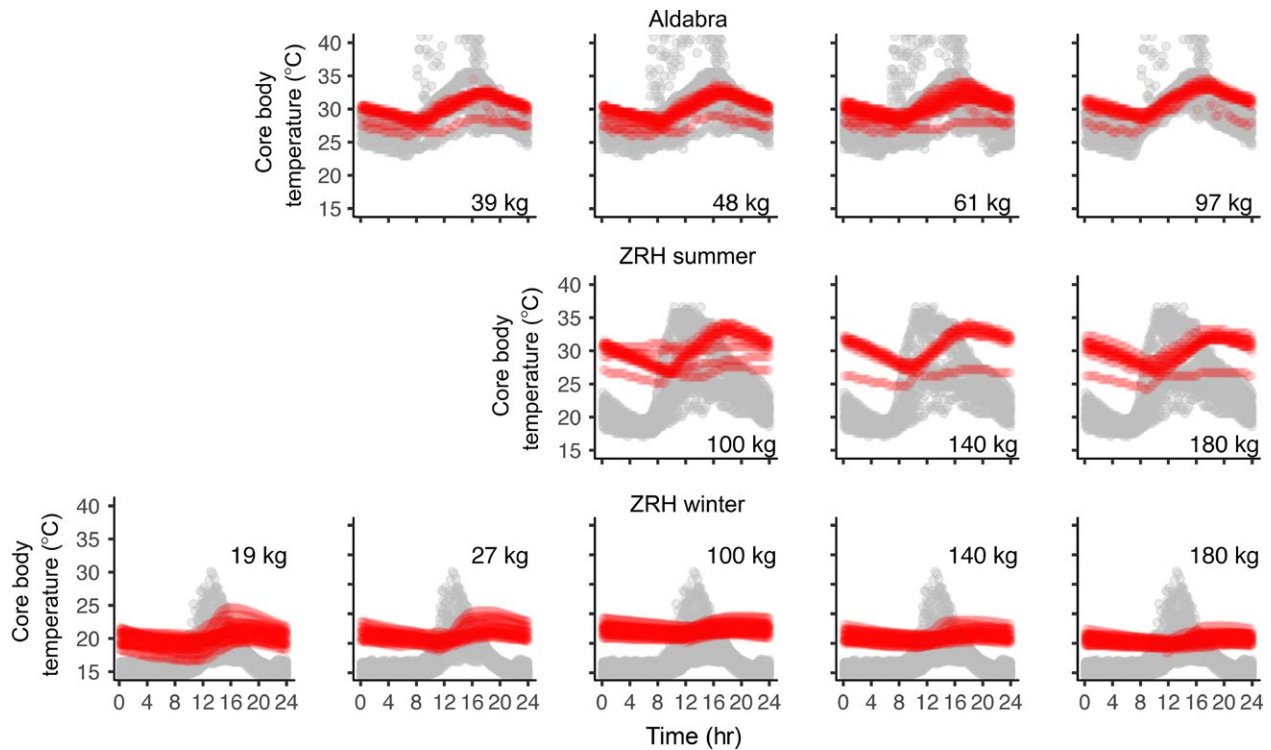
**FIGURE 3** Temporal fluctuations in the environmental temperature range ( $T_{\text{air}}$  and  $T_{\text{sun}}$ ; gray fill) and core temperature recorded by data loggers in the gastrointestinal tract of Aldabra giant tortoises (*Aldabrachelys gigantea*) in different trials. The dark gray line depicts  $T_{a \text{ mean}}$ , and cool to warm colored lines depict ascending mass range for individual tortoises (with mass given in the legend). One trial was performed on Aldabra Atoll with wild tortoises, and two trials were performed at Zurich Zoo (during winter and summer). Tick marks in the x-axis depict a 24-hr interval: peaks of environmental temperature indicate temperature at ~mid-day, and lowest point of the valleys indicates temperatures at ~midnight



$\Delta T_{\text{air}}$ , but it was only significant for the ZRH trials ( $z = 5.87$  and  $4.12$ ,  $p < .01$ , Kendall  $\tau = 0.43$  and  $0.55$  for the winter and summer trials, respectively). On Aldabra, there was no significant correlation ( $z = 1.46$ ,  $p = .14$ , Kendall  $\tau = 0.16$ ). For ZRH winter,  $\Delta T_{\text{bc}}$  decreased from the smallest to the largest animal ( $z = -4.76$ ,  $p < .001$ , Kendall  $\tau = -0.37$ ). In contrast,  $\Delta T_{\text{bc}}$  of tortoises during ZRH summer (without smaller, juvenile tortoises) and on Aldabra remained virtually the same between individuals of different mass ( $z = -0.73$  and  $0.28$ ,  $p = .47$  and  $p = .78$ , Kendall  $\tau = -0.10$  and  $0.03$ , respectively). Moreover, in ZRH winter, the  $\Delta T_{\text{bc}}/\Delta T_{\text{air}}$  quotient was negatively correlated with tortoise body mass ( $z = -5.87$ ,  $p < .001$ , Kendall  $\tau = -0.45$ ). However, we found no

correlation in the ZRH summer trial (where no juveniles were included) or the Aldabra trial ( $z = -0.71$  and  $0.12$ ,  $p = .48$  and  $0.91$ , Kendall  $\tau = -0.10$  and  $0.01$ , respectively).

There was greater variation in  $T_{\text{be}}$  than in  $T_{\text{bc}}$  (Figure 5; see S5 in Supplementary Information for summary statistics). Daily temperature fluctuations were greater on the carapace surface than on the surface of the extremities, which in turn were greater than in the skin folds or in the core body temperature. Temporal turning points of the temperature curves occurred first in the environment, followed by the carapace, the extremities, the skin folds, and, finally, the core. Temperatures of the carapace and extremities were also correlated



**FIGURE 4** Daily environmental temperatures (gray;  $T_{\text{air}}$  and  $T_{\text{sun}}$ ) and core body temperature (red;  $T_{\text{bc}}$ ) of Aldabra giant tortoises (*Aldabrachelys gigantea*) of different mass across different environmental temperature ranges (trials)

with  $T_{\text{bc}}$ , but to a lesser degree ( $z = 33.38$ ,  $p < .001$ , Kendall  $\tau = 0.70$ , and  $z = 51.28$ ,  $p < .001$ , Kendall  $\tau = 0.59$ , respectively), and remained lower than  $T_{\text{bc}}$  (Wilcoxon rank sum test,  $p < .001$  in both cases). The temperature recorded in the skin folds was highly correlated with but remained lower than  $T_{\text{bc}}$  ( $z = 37.77$ ,  $p < .001$ , Kendall  $\tau = 0.79$ ; Wilcoxon rank sum test,  $p < .001$ ).

All explanatory variables of the thermoregulation model, *trial*, *time*,  $T_{\text{air}}$ , *mass*, and the interaction between  $T_{\text{air}}$  and *mass*, influenced the variation in  $T_{\text{bc}}$  at  $p < .001$  (see S6 in Supplementary Information for model summary statistics). When considered independently,  $T_{\text{air}}$  better explained the variation in the  $T_{\text{bc}}$  of tortoises than *mass* ( $\Delta\text{AIC} = 1668$ ) or the interaction between  $T_{\text{air}}$  and *mass* ( $\Delta\text{AIC} = 1128$ ). The relationship between *mass* and  $T_{\text{bc}}$  in ZRH winter is bell-shaped, increasing until it reaches 100 kg and then decreasing again. In contrast,  $T_{\text{bc}}$  did not exhibit any clear pattern in relation to *mass* in the summer trial (where no juveniles were used, and hence, the body mass range was much smaller than during winter), while on Aldabra  $T_{\text{bc}}$  showed a slight increase with *mass* (Table 2). Similar patterns to those exhibited by the relationship between *mass* and  $T_{\text{bc}}$  for each trial were observed for the acrophase, and the time lag of  $T_{\text{bc}}$  to acrophase (i.e. the time difference at which  $T_{\text{bc}}$  reaches the acrophase in relation to  $T_{\text{a}}$ ).

For the relationship between  $T_{\text{a}}$  and  $T_{\text{bc}}$  for all tortoises in our study, most of the data point lay above  $T_{\text{a}}$  and slowly shifted toward lower values after  $T_{\text{a}}$  reached  $>30^{\circ}\text{C}$  (Figure 6). As before, the variation in  $T_{\text{bc}}$  decreased with increasing size only during the winter trial. Interestingly, the  $T_{\text{bc}}$  trend increased more sharply with increasing  $T_{\text{air}}$  (temperature of shaded areas where they seek refuge) in contrast to  $T_{\text{sun}}$ .

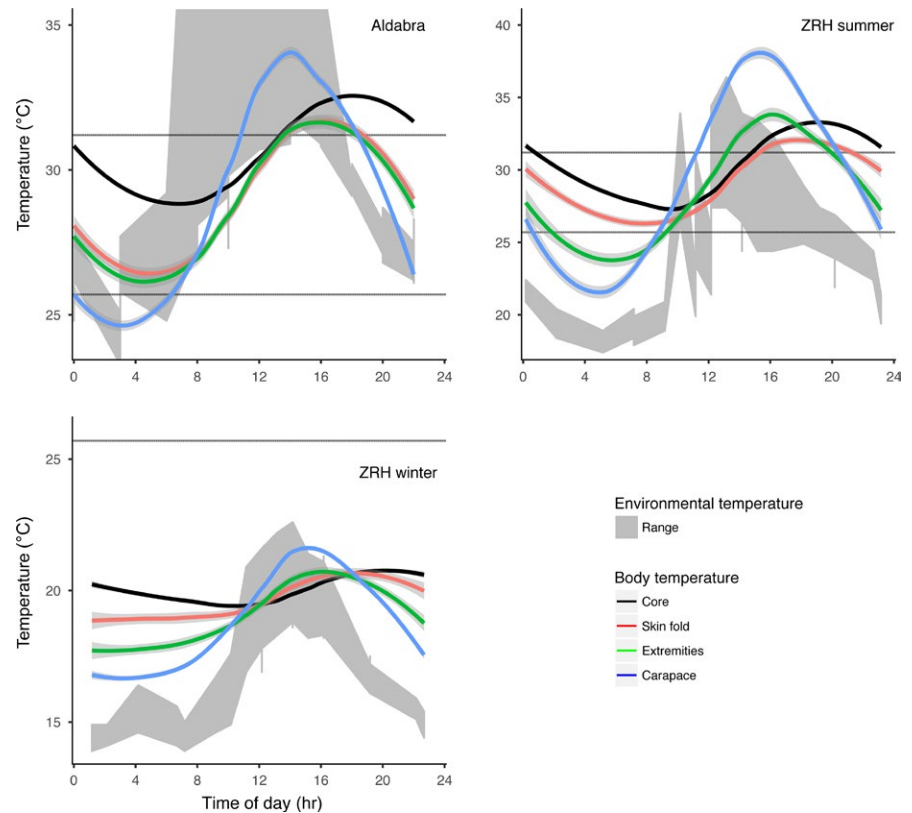
### 3.3 | Body size and temperature in testudinidae

In addition to our data, we were able to gather 22 measurements of six species in four genera, from six studies (Benedict, 1932; Huot-Daubremont, Grenot, & Bradshaw, 1996; Mackay, 1964; McMaster & Downs, 2013c; Meek & Jayes, 1982; Swingland & Frazier, 1979). Seven of these 22 data points contain estimated mass values from various individuals, and methods for measuring temperature varied (see S7 in Supplementary Information for details).

The  $\Delta T_{\text{bc}}$  of different Testudinid species was negatively correlated with their *mass* ( $z = -3.11$ ,  $p = .002$ , Kendall  $\tau = -0.38$ ; Figure 7a). However, the  $\Delta T_{\text{bc}}$  of tortoises was more strongly positively correlated with  $\Delta T_{\text{air}}$  ( $z = 4.60$ ,  $p < .001$ , Kendall  $\tau = 0.57$ ; Figure 7b). When corrected for  $\Delta T_{\text{air}}$  (using the ratio of the of  $\Delta T_{\text{bc}}$  to  $\Delta T_{\text{air}}$ ), there was a weaker negative correlation between the  $\Delta T_{\text{bc}}/\Delta T_{\text{air}}$  and *mass* ( $z = -2.42$ ,  $p = .02$ , Kendall  $\tau = -0.29$ ; Figure 7c).

## 4 | DISCUSSION

Abiotic, biotic, physiological, and behavioral factors play an important role in the regulation of body temperature of reptiles. Here, we described the activity patterns of Aldabra giant tortoises in relation to the environmental temperature, the optimum environmental temperature at which tortoises maximize their activity, and how different gradients of environmental temperatures and body mass influenced the variation in their internal and external body temperatures. Moreover, we found evidence of thermal inertia, but this effect seemed to be



**FIGURE 5** Average daily fluctuations of environmental temperature and Aldabra giant tortoise (*Aldabrachelys gigantea*) body temperatures on Aldabra Atoll and the Zurich Zoo (winter and summer). Horizontal lines depict the temperature range at which tortoise activity is maximised. Note that the y-axis is scaled independently for each trial. The range intervals of  $T_a$  (maxima & minima) are depicted in gray shading in the background. The shading around the lines represents the 95% CI based on the line smoothing by local regression-loess. For the Aldabra trial, sun temperatures go beyond the y-axis limits

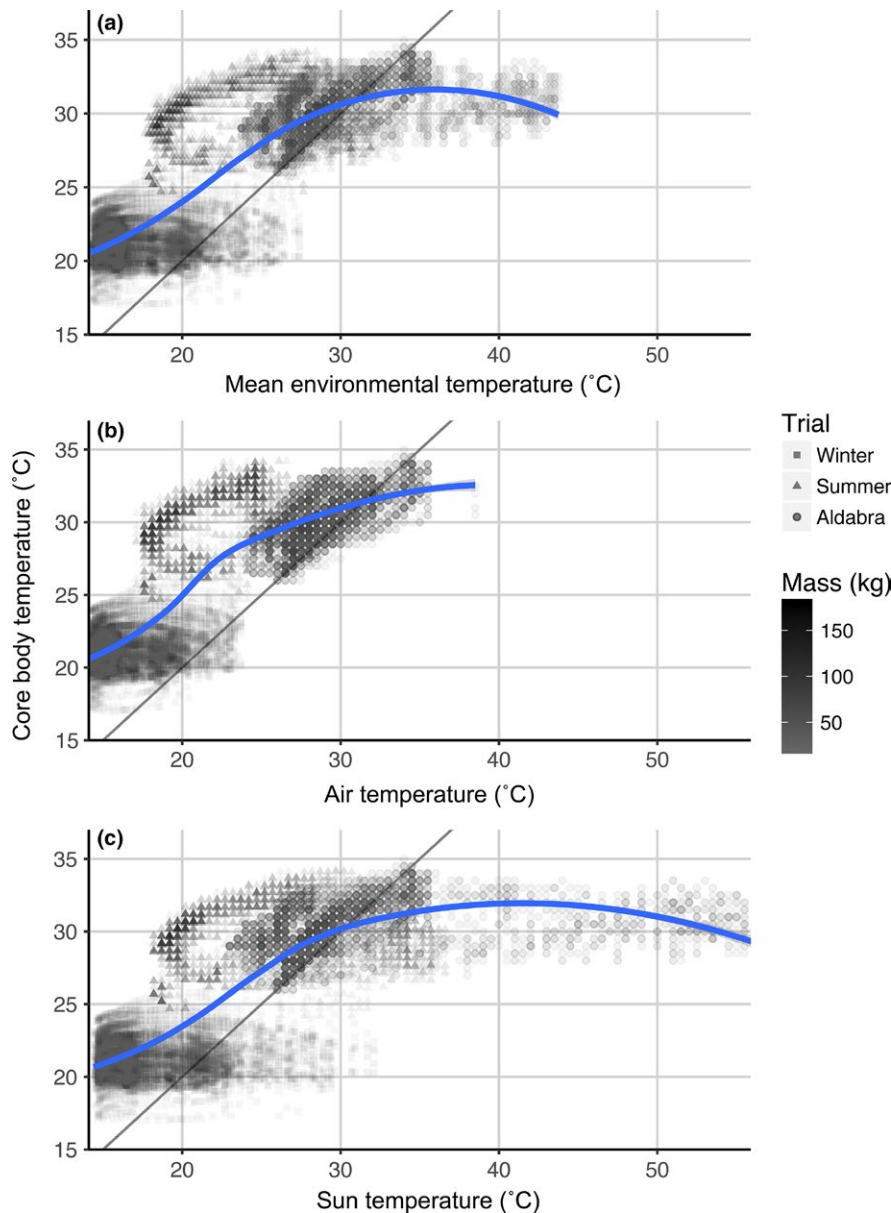
context dependent in terms of the environmental temperatures experienced by the tortoises, and the overall Testudinidae dataset indicated little effect of mass on the temperature stability of tortoises.

#### 4.1 | Environmental temperatures and tortoise activity patterns

We used activity patterns of Aldabra tortoises in response to  $T_{air}$  to identify the temperature range at which activity is maximized and use this as a proxy for optimal environmental temperature range,  $T_{a-opt}$ . The activity of Aldabra giant tortoises is similar to that of other tortoise species, including southern Africa's largest tortoise, the leopard tortoise (*Stigmochelys pardalis*; Monadjem, McCleery, & Collier, 2013; McMaster & Downs, 2013a,c). Similar to Aldabra giant tortoises (R. P. Baxter et al., unpubl. data.), leopard tortoises exhibit a transition in daily activity patterns across seasons, being unimodal during the dry-cold season, and shifting progressively toward bimodal as the season transitions toward hot and wet. In addition, under similar  $T_a$  conditions as those experienced by Aldabra giant tortoises (i.e., during the Nama-Karoo South African summer), leopard tortoises started becoming inactive when  $T_a$  increased above 30.5–32°C between 10:00 and 11:00 hr (McMaster & Downs, 2013a), when presumably tortoises seek shade to cool down. Moreover, Lambert (1981) found similar relationships between temperature and activity in the spur-thighed tortoise (*Testudo graeca*), although these tortoises were active at a lower temperature range of 18–28°C and inactive when  $T_a$  was below 18°C. The  $T_a$  threshold for switching from a unimodal to bimodal activity pattern in *T. graeca* was 28°C.

Aldabra giant tortoises maximize their activity ( $T_{a-opt}$ ) in the temperature range of 25.8–31.7°C. This is within the preferred temperature range (i.e., the range of  $T_{bc}$  within which an ectotherm seeks to maintain itself by behavioral means) of other Testudinid species, with reported ranges of 25–31°C (*Gopherus agassizii*, Woodbury & Hardy, 1948; *G. agassizii* and *Testudo hermanni*, Brattstrom, 1965; *T. marginata* and *T. hermanni*, Panagiota & Valakos, 1992; Huot-Daubremont et al., 1996). Moreover, the range of mean  $T_{bc}$  maintained by wild Galápagos (*Chelonoidis nigra*) and Aldabra giant tortoises was within  $T_{a-opt}$  (Mackay, 1964; Swingland & Frazier, 1979; this study). Rather than preferred body temperatures, we calculated  $T_{a-opt}$ , and our environmental temperature measurements were performed in the shade. Because tortoises may be exposed to the sun when active, our measured optimal environmental temperature range may be biased to the lower limits of  $T_{a-opt}$  (i.e.,  $T_{a-opt}$  may actually be higher). However, the pronounced drop in activity probability once temperature in the shade ( $T_{air}$ ) increases above 31°C (upper limit of  $T_{a-opt}$  range), as well as the clear thermoregulatory pattern exhibited by tortoises when  $T_{bc}$  reaches this temperature, suggests that the upper limit of the preferred core body temperature range is around 31°C. This is in accordance with the findings of Swingland and Frazier (1979), who reported the maximum critical temperature for Aldabra giant tortoises in the wild to be 36–38°C (measured in tortoises dying or that recently died from overheating in the wild). Therefore, the  $T_{a-opt}$  range in combination with the mean  $T_{bc}$  of wild tortoises may serve as a reference for a high quality thermal environment for husbandry and captive care practices (e.g., McArthur & Barrows, 2008) and conservation efforts



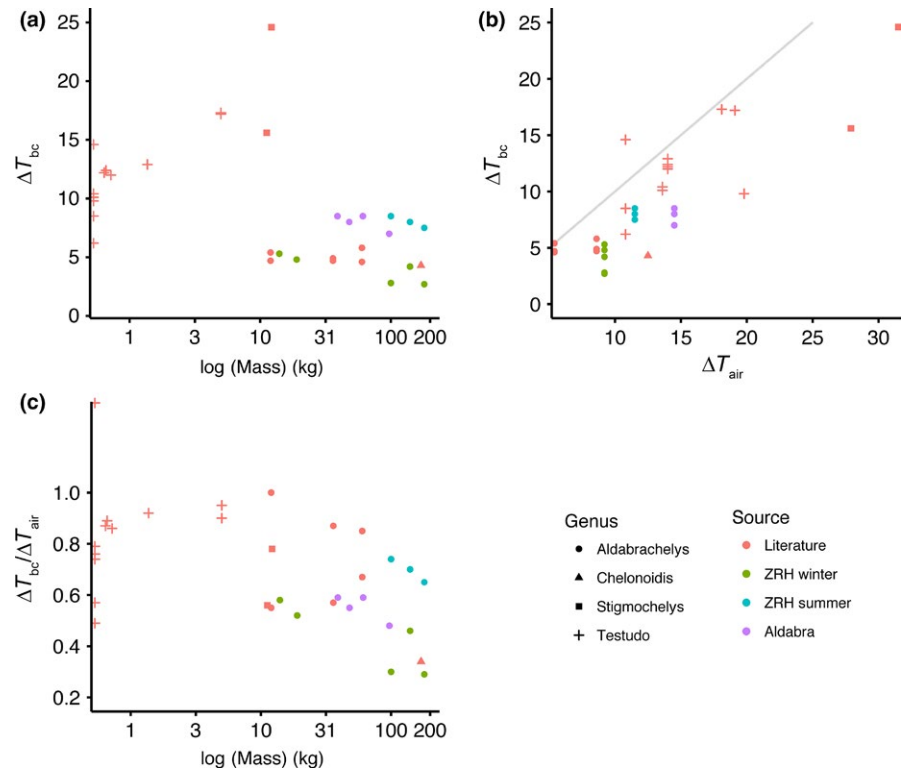


**FIGURE 6** Aldabra giant tortoise (*Aldabrachelys gigantea*) core body temperature ( $T_{bc}$ ) in relation to mean environmental temperatures ( $T_a$ ), air temperature ( $T_{air}$ , in the shade; b), and in the sun-exposed areas ( $T_{sun}$ ; c). Points represent the  $T_{bc}$  of tortoises, with the gray-color gradient indicating body mass, and trials indicated by shape. The solid-line represents a 1:1 response, while the blue lines represent the response of the tortoises'  $T_{bc}$  (all animals per trial combined) to environmental temperatures based on smoothing by local regression–loess (the shading represents the 95% CI)

with regard to climate change and thermal refuge availability (e.g., Moulherat et al., 2014; Swingland & Frazier, 1979).

Although the activity of tortoises was affected by seasonality, our results are likely to capture the full extent of environmental temperatures at which Aldabra tortoises maximize their activity because we include activity and temperature data from a 2-year period. However, the data show the flexibility of tortoises with regard to their activity patterns and available  $T_{air}$ , depicted by the variation in the seasonal activity patterns in relation to temperature. Other tortoises also show flexibility in terms of their activity in relation to seasonal changes in temperature (among other factors, see McMaster & Downs, 2013a). Rainfall and temperature have been shown to be important factors influencing the activity of tortoises (Kazmaier, Hellgren, & Synatzske, 2011). In our activity model, the interaction between the dry season and  $T_{air}$  negatively affected activity. We thus hypothesize that the changes in activity patterns dependent on seasons in Aldabra giant

tortoises may also be driven by changes in water balance and water conservation. Water balance is important in tortoises because it affects their food consumption, diet, daily behavior, osmoregulation, and body mass (Nagy & Medica, 1986). Tortoises have been shown to suffer significant evaporative water loss from their integuments and through respiration (Schmidt-Nielsen & Bentley, 1966). Moreover, increasing temperatures and drought conditions have been shown to increase water loss through evaporation in tortoises (Cloudsley-Thompson, 1968; Minnich, 1977). The expected main mechanism for thermoregulation in tortoises is to change their daily activity levels and behavior (because the rigid shell limits the effectiveness of postural changes during behavioral thermoregulation; McMaster & Downs, 2013c). To conserve water, tortoises may decrease their activity levels and stay in the shade during and after the warmest part of the day in the dry season. If our hypothesis is correct, we can expect that the increasing frequency of drought periods on Aldabra



**FIGURE 7** Core body temperature ranges ( $\Delta T_{bc}$ ) of Testudines in relation to the environmental temperature range ( $\Delta T_{air}$ ) and their mass. Relationship between  $\Delta T_{bc}$  and mass (a),  $\Delta T_{bc}$  and  $\Delta T_{air}$  (b), and between the  $\Delta T_{bc}$  and  $\Delta T_{air}$  ratio (to control for environmental temperature; c). Point shapes indicate the genus, and color indicates the source of the data. In Panel c, one of the points is  $>1$ , probably because the air temperature did not capture the environmental temperature at which the tortoise was exposed (e.g., if the tortoise was mainly basking in the sun)

(Haverkamp et al., 2017) will have negative impacts on the activity of giant tortoises.

It should be noted that temperature is not the only factor influencing the activity of tortoises. For example, Gibson and Hamilton (1983) hypothesized that seasonal changes in the activity of Aldabra giant tortoises were in response to food availability. Moreover, precipitation, solar radiation, and wind velocity also seem to play a role in determining the bimodal to unimodal activity patterns and the activity levels of Aldabra giant tortoises (unpubl. data). Further research is being undertaken to determine the environmental drivers of the activity of Aldabra giant tortoises on Aldabra.

#### 4.2 | Body temperature of *Aldabrachelys gigantea*

Fluctuations in  $T_{bc}$  lagged behind those of  $T_a$ , and in general tortoises heated more rapidly during the day than they lost heat during the night, when  $T_{bc}$  fell slowly. During the ZRH winter trial,  $T_{bc}$  was always higher than  $T_{air}$ ; only the temperature logger placed in direct sunlight recorded temperatures that were higher than  $T_{bc}$ , especially during the middle of the day.  $T_{air}$  better explained  $T_{bc}$  of tortoises, and in general, they were able to maintain their  $T_{bc}$  above low  $T_{air}$ , and below high  $T_{air}$ , and their  $T_{bc}$  was affected by the range of available environmental temperatures. Moreover, the temperatures measured on the surface of the carapaces were notably higher than  $T_{air}$  during the morning, which is evidence for thermoregulation via basking behavior (e.g., Crawford, Spotila, & Standora, 1983; Lambert, 1981). While basking, reptiles reach higher core temperatures than air temperatures, and  $T_{bc}$  correlates positively with time spent basking (Boyer, 1965; Rivera-Vélez & Lewis, 1994). Correspondingly,  $T_{bc}$  dropped in cloudy days, when

basking was not possible, and environmental temperatures dropped by 3–5°C. During ZRH summer and on Aldabra, the mean and the range of  $T_a$  were higher, and tortoises were able to maintain their  $T_{bc}$  close to  $T_{a-opt}$ . On Aldabra, we recorded some extreme high temperatures in the sun during the second half of the study, probably due to clear skies and virtually no wind, but tortoises were able to maintain a stable  $T_{bc}$  nonetheless. Swingland and Frazier (1979) reported very similar patterns of  $T_{bc}$  for *A. gigantea* to those exhibited by our tortoises, but in the southeast of the atoll on Grande Terre Island, where shade is more limited and where the size dimorphism of tortoises is much less pronounced. Additionally, Mackay (1964) observed similar patterns of  $T_{bc}$  in two Galápagos giant tortoises (65 and 170 kg), which were able to maintain their  $T_{bc}$  within ~28–32°C when the mean  $T_a$  was ca. 28°C and fluctuated between ca. 23–35°C, despite the difference in their mass.

As found in leopard tortoises (McMaster & Downs, 2013b), the differential variation in  $T_{bc}$  of different body surfaces and as well as that of  $T_{bc}$  indicated that there are large thermal gradients within the bodies of Aldabra giant tortoises. For example, the maximum carapace temperature of Aldabra giant tortoises in our study sometimes greatly exceeded that of their recorded  $T_{bc}$ , to the point that it surpassed the maximum critical temperature (of  $T_{bc}$ ) recorded for the species. Studying the thermoregulation of Galápagos giant tortoises, Mackay (1964) proposed, based on the temperature differential between the core body temperature and the carapace, that heat flows through a limited region with high resistance when compared to that of the material absorbing the heat. We found the same pattern in Aldabra giant tortoises, where the integral of the difference between the  $T_{bc}$  and the carapace temperature (mean of the vertebral and costal scutes) followed the same temporal pattern as that of  $T_{bc}$ . While in the shade, the temperature of the scutes

on the carapace of the tortoises remained above shaded environmental temperatures as well as above the  $T_{bc}$ , indicating that they lost heat to the environment. In contrast, at night, carapace temperatures dropped below  $T_{bc}$  and were closer to  $T_a$  than that of the extremities or skin folds (which were also below  $T_{bc}$ ). Thus, our results suggest that Aldabra giant tortoises employ different behavioral and physiological mechanisms to use their carapace as a heat exchanger: a heat collector in the mornings, a radiator (i.e., cooling system) during the warmest part of the day, and an insulator during the coldest part (see McMaster & Downs, 2013b, and references therein, for a discussion of the differential temperature of  $T_{bc}$  and  $T_{be}$  in tortoises and possible control mechanisms).

It is worth noting that Aldabra hosted a population of introduced goats until 2012, when an eradication program was completed (Bunbury et al., 2013). In 1985, during the same time as the tortoise population decline from an estimated 130,000 to around 100,000 (Bourn et al., 1999), Coblenz and Vuren (1987) estimated that there were as many as 1,300 goats on Aldabra. They suggested that the major impact of the goats was their negative effect on shade resources through overbrowsing, rather than direct competition with tortoises for food. Our results suggest a strong role of available shade for structuring tortoise activity and in body temperature regulation, supporting the likelihood of overbrowsing by goats having had a negative impact on Aldabra's giant tortoises.

### 4.3 | Body size and temperature in testudinidae

Thermal inertia likely explains why after basking, tortoises at the Zurich Zoo had higher  $T_{bc}$  than minimum  $T_a$  during early mornings on the next day, and cooling rates appeared to decrease with size. However, the effect of mass on temperature stability of Aldabra giant tortoises ( $\Delta T_{bc}$ ) differed by trial. When the mean  $T_a$  was 17.2°C during the ZRH winter trial,  $T_{bc}$  stability increased with increasing size. This was not the case for the mean  $T_{bc}$ . In ZRH winter, the tortoise of intermediate size (100 kg) had the highest mean  $T_{bc}$ , suggesting that both the fast cooling rate of small animals and the slow heating rate of larger animals influenced their mean  $T_{bc}$ . On the other hand,  $\Delta T_{bc}$ , as well as the mean core body temperatures in ZRH summer and on Aldabra, did not seem to vary with mass. Furthermore, there was a tendency for the acrophase of tortoises (time at which  $T_{bc}$  peaked), and of the thermal lag of  $T_{bc}$  to  $T_a$ , to increase with mass in the Zurich tortoises, but there was no apparent trend on Aldabra. Also, the effect size of mass in our thermoregulation model was rather small, and  $T_{air}$  better explained changes in the  $T_{bc}$  of Aldabra giant tortoises. Moreover, the analysis on the overall Testudinidae dataset appears to indicate that within the body size range of the tortoise species studied, large individuals are subject to similar fluctuations in body temperature as smaller ones once variation in air temperature has been taken into account. Thus, the presence of thermal inertia in tortoises seems to depend on the environmental temperatures. However, it is notable that behavioral thermoregulation and acclimatization can potentially override the effects of mass on the  $T_{bc}$  of tortoises and thus provide an alternative and possibly synergistic explanation of the apparent context-dependent effect of mass on  $T_{bc}$ .

In addition, the notion that large ectotherms may maintain a high (30°C) and stable  $T_{bc}$  within a narrow range (2°C) due to mass-dependent thermal inertia, similar to homeothermic endotherms, is often referred to as "inertial homeothermy" (McNab & Auffenberg, 1976; Seebacher, 2003). Despite being "giants," even under stable conditions, the range of  $T_{bc}$  in Aldabra giant tortoises (as well as other species of smaller tortoises) was much larger than 2°C. These results indicate that that inertial homeothermy is not possible in tortoises with the range of body masses studied. These findings support the conclusion of Grigg, Beard, and Augee (2004), who found that inertial homeothermy over the course of a single day is only found in large ectotherms above 500 kg of body size.

## 5 | CONCLUSIONS

Despite the large sizes that Aldabra giant tortoises can attain, and the presence of thermal inertia, our results suggest that tortoises are incapable of regulating their core body temperatures within a range narrow enough to be considered inertial homeotherms. Rather, the interplay between the mass of the tortoises and the variation of  $T_a$ , in combination with behavioral thermoregulation, limits the degree to which these ectotherms can attain core body temperatures close to their presumed optimum  $T_{bc}$ . We found evidence of thermoregulation, where tortoises were able to maintain  $T_{bc}$ s independent of  $T_a$ . We also found instances of thermoconformity, and when we evaluated  $T_{bc}$  in response to  $T_a$ , our results indicate that giant tortoises act as mixed conformer-regulators (Willmer et al., 2005). However, although the relationship between  $T_{bc}$  and  $T_a$  suggests that giant tortoises can maintain a stable  $T_{bc}$  when the mean  $T_a$  is above the lethal temperature (>36°C), evaluating the components of  $T_a$  independently (the temperature of shade and sun-exposed loggers) shows that Aldabra giant tortoises have a limit: when shade temperature ( $T_{air}$ ) surpasses ca. 31°C, the  $T_{bc}$  seems to keep increasing rather than reaching a plateau. While some tortoises can adjust their behavior to survive extreme environmental temperatures (e.g., *Gopherus agassizii* can remain active even when  $T_a$  reaches ca. 60°C by adjusting the time spent in burrows; Zimmerman et al., 1994), larger animals such as *A. gigantea* have limited options. The plasticity and intraspecific and interspecific variation in tortoises, and other reptiles, certainly allow for the animals to respond to the selective pressures imposed by the environment. However, it is likely that climate change will accentuate thermoregulatory pressures (Barrows, 2011; Gunderson & Stillman, 2015), especially on larger species.

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## DATA ACCESSIBILITY

The data used for this article are accessible through Dryad (<http://datadryad.org/resource/doi:10.5061/dryad.b4v22>).

## CONFLICT OF INTEREST

None declared.

## AUTHORS' CONTRIBUTIONS

WF, RB, DH designed the Aldabra part of this study and collected the data and SF, MB, JMH, MC designed the Zurich part of this study; SF and MC collected the data. WF, MC, DH, and AO analyzed the data; WF, MC, DH performed the literature review and drafted a version of the manuscript that then received input from all coauthors.


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## REFERENCES

- Barrows, C. W. (2011). Sensitivity to climate change for two reptiles at the Mojave-Sonoran Desert interface. *Journal of Arid Environments*, 75(7), 629–635. <https://doi.org/10.1016/j.jaridenv.2011.01.018>
- Bartholomew, G. A., & Tucker, V. A. (1963). Control of changes in body temperature, metabolism, and circulation by the agamid lizard, *Amphibolurus barbatus*. *Physiological Zoology*, 36(3), 199–218. <https://doi.org/10.1086/physzool.36.3.30152307>
- Bates, D., Machler, M., Bolker, B., & Walker, S. (2015). Fitting linear mixed-effects models using lme4. *Journal of Statistical Software*, 67(1), 1–48. <https://doi.org/10.18637/jss.v067.i01>
- Bauert, M. R., Furrer, S. C., Zingg, R., & Steinmetz, H. W. (2007). Three years of experience running the Masoala Rainforest ecosystem at Zurich Zoo, Switzerland. *International Zoo Yearbook*, 41(1), 203–216. <https://doi.org/10.1111/j.1748-1090.2007.00012.x>
- Benedict, F. G. (1932). *The physiology of large reptiles: With special reference on the heat production of snakes, tortoises, lizards and alligators*. Washington, DC: Carnegie Institution of Washington.
- Blatteis, C., Boulant, J., Cabanac, M., Cannon, B., Freedman, R., Gordon, C. J., ... Kozyreva, T. (2001). Glossary of terms for thermal physiology. *The Japanese Journal of Physiology*, 51(2), 245–280. <https://doi.org/10.2307/1467052>
- Bourn, D., Gibson, C., Augeri, D., Wilson, C. J., Church, J., & Hay, S. I. (1999). The rise and fall of the Aldabran giant tortoise population. *Proceedings of the Royal Society B-Biological Sciences*, 266(1424), 1091–1100. <https://doi.org/10.1098/rspb.1999.0748>
- Boyer, D. R. (1965). Ecology of the basking habit in turtles. *Ecology*, 46(1–2), 99–118. <https://doi.org/10.2307/1935262>
- Brattstrom, B. H. (1965). Body temperatures of reptiles. *The American Midland Naturalist*, 73(2), 376. <https://doi.org/10.2307/2423461>
- Bunbury, N., von Brandis, R., Currie, J., Jean-Baptiste, M., Accouche, W., Souyave, J., ... Fleischer Dogley, F. (2013). Goats eradicated from Aldabra Atoll. *Aliens: The Invasive Species Bulletin*, 33, 18–22.
- Cloudsley-Thompson, J. L. (1968). Thermoregulation in tortoises. *Nature*, 217(5128), 575. <https://doi.org/10.1038/217575a0>
- Coblentz, B. E., & Vuren, D. V. (1987). Effects of feral goats (*Capra hircus*) on Aldabra Atoll. *Atoll Research Bulletin*, 306, 1–6. <https://doi.org/10.5479/si.00775630.306.1>
- Cossins, A. R., & Bowler, K. (1987). *Temperature biology of animals*, 1st ed. New York, NY: Chapman and Hall. <https://doi.org/10.1007/978-94-009-3127-5>
- Crawford, K. M., Spotila, J. R., & Standora, E. A. (1983). Operative environmental temperatures and basking behavior of the turtle *Pseudemys scripta*. *Ecology*, 64(5), 989–999. <https://doi.org/10.2307/1937806>
- Gibson, C. W. D., & Hamilton, J. (1983). Feeding ecology and seasonal movements of giant tortoises on Aldabra atoll. *Oecologia*, 56(1), 84–92. <https://doi.org/10.1007/BF00378221>
- Grigg, G. C., Beard, L. A., & Auger, M. L. (2004). The evolution of endothermy and its diversity in mammals and birds. *Physiological and Biochemical Zoology*, 77(6), 982–997. <https://doi.org/10.1086/425188>
- Gunderson, A. R., & Stillman, J. H. (2015). Plasticity in thermal tolerance has limited potential to buffer ectotherms from global warming. *Proceedings of the Royal Society B-Biological Sciences*, 282(1808), 20150401. <https://doi.org/10.1098/rspb.2015.0401>
- Haverkamp, P. J., Shekeine, J., de Jong, R., Schaepman, M., Turnbull, L. A., Baxter, R., ... Schaepman-Strub, G. (2017). Giant tortoise habitats under increasing drought conditions on Aldabra Atoll—Ecological indicators to monitor rainfall anomalies and related vegetation activity. *Ecological Indicators*, 80, 354–362. <https://doi.org/10.1016/j.ecolind.2017.05.029>
- Heatwole, H. (1976). *Reptile ecology*. Brisbane, Qld: University of Queensland Press.
- Huey, R. B., & Stevenson, R. D. (1979). Integrating thermal physiology and ecology of ectotherms: A discussion of approaches. *Integrative and Comparative Biology*, 19(1), 357–366. <https://doi.org/10.1093/icb/19.1.357>
- Huot-Daubremont, C., Grenot, C., & Bradshaw, D. (1996). Temperature regulation in the tortoise *Testudo hermanni*, studied with indwelling probes. *Amphibia-Reptilia*, 17(2), 91–102. <https://doi.org/10.1163/156853896X00135>
- Jammalamadaka, S. R., & Lund, U. J. (2006). The effect of wind direction on ozone levels: A case study. *Environmental and Ecological Statistics*, 13(3), 287–298. <https://doi.org/10.1007/s10651-004-0012-7>
- Kazmaier, R. T., Helligren, E. C., & Synatzske, D. R. (2011). Patterns of behavior in the Texas tortoise, *Gopherus berlandieri*: A multivariate ordination approach. *Canadian Journal of Zoology*, 79(8), 1363–1371. <https://doi.org/10.1139/z01-092>
- Kinahan, A. A., Inge-moller, R., Bateman, P. W., Kotze, A., & Scantlebury, M. (2007). Body temperature daily rhythm adaptations in African savanna elephants (*Loxodonta africana*). *Physiology & Behavior*, 92(4), 560–565. <https://doi.org/10.1016/j.physbeh.2007.05.001>
- Kuznetsova, A., Brockhoff, P. B., & Christensen, R. H. B. (2016). *ImerTest: Tests in linear mixed effects models*. 2nd edn. Retrieved from <https://CRAN.R-project.org/package=ImerTest>.



- Lailvaux, S. P., & Irschick, D. J. (2007). Effects of temperature and sex on jump performance and biomechanics in the lizard *Anolis carolinensis*. *Functional Ecology*, 21(3), 534–543. <https://doi.org/10.1111/j.1365-2435.2007.01263.x>
- Lambert, M. R. K. (1981). Temperature, activity and field sighting in the mediterranean spur-thighed or common garden tortoise *Testudo graeca* L. *Biological Conservation*, 21(1), 39–54. [https://doi.org/10.1016/0006-3207\(81\)90067-7](https://doi.org/10.1016/0006-3207(81)90067-7)
- Mackay, R. S. (1964). Galapagos tortoise and marine iguana deep body temperatures measured by radio telemetry. *Nature*, 204(4956), 355–358. <https://doi.org/10.1038/204355a0>
- McArthur, S., & Barrows, M. (2008). General care of chelonians. In S. McArthur, R. Wilkinson, & J. Meyer (Eds.), *Medicine and surgery of tortoises and turtles* (pp. 87–108). Oxford, UK: John Wiley & Sons.
- McMaster, M. K., & Downs, C. T. (2013a). Seasonal and daily activity patterns of leopard tortoises (*Stigmochelys pardalis* Bell, 1828) on farmland in the Nama-Karoo, South Africa. *African Zoology*, 48(1), 72–83. <https://doi.org/10.3377/004.048.0110>
- McMaster, M. K., & Downs, C. T. (2013b). Thermal variability in body temperature in an ectotherm: Are cloacal temperatures good indicators of tortoise body temperature? *Journal of Thermal Biology*, 38(4), 163–168. <https://doi.org/10.1016/j.jtherbio.2013.02.002>
- McMaster, M. K., & Downs, C. T. (2013c). Thermoregulation in leopard tortoises in the Nama-Karoo: The importance of behaviour and core body temperatures. *Journal of Thermal Biology*, 38(4), 178–185. <https://doi.org/10.1016/j.jtherbio.2013.02.003>
- McNab, B. K., & Auffenberg, W. (1976). The effect of large body size on the temperature regulation of the komodo dragon, *Varanus komodoensis*. *Comparative Biochemistry and Physiology Part A: Physiology*, 55(4), 345–350. [https://doi.org/10.1016/0300-9629\(76\)90058-X](https://doi.org/10.1016/0300-9629(76)90058-X)
- Meek, R. (1984). Thermoregulatory behaviour in a population of Hermann's tortoise (*Testudo hermanni*) in southern Yugoslavia. *British Journal of Herpetology*, 6(11), 387–391.
- Meek, R., & Jayes, A. S. (1982). Body temperatures and activity patterns of *Testudo graeca* in North West Africa. *British Journal of Herpetology*, 6, 194–197.
- Minnich, J. E. (1977). Adaptive responses in the water and electrolyte budgets of native and captive desert tortoises, *Gopherus agassizii* to chronic drought. Las Vegas: Proceedings of the Desert Tortoise Council Symposium, pp. 102–129.
- Monadjem, A., McCleery, R. A., & Collier, B. A. (2013). Activity and movement patterns of the tortoise *Stigmochelys pardalis* in a subtropical savanna. *Journal of Herpetology*, 47(2), 237–242. <https://doi.org/10.1670/12-070>
- Moulherat, S., Delmas, V., Slimani, T., El Mouden, E. H., Louzizi, T., Lagarde, F., & Bonnet, X. (2014). How far can a tortoise walk in open habitat before overheating? Implications for conservation. *Journal for Nature Conservation*, 22(2), 186–192. <https://doi.org/10.1016/j.jnc.2013.11.004>
- Nagy, K. A., & Medica, P. A. (1986). Physiological ecology of desert tortoises in southern Nevada. *Herpetologica*, 42(1), 73–92. <https://doi.org/10.2307/3892239>
- Nielsen, L. R., Pedersen, A. R., Herskin, M. S., & Munksgaard, L. (2010). Quantifying walking and standing behaviour of dairy cows using a moving average based on output from an accelerometer. *Applied Animal Behaviour Science*, 127(1), 12–19. <https://doi.org/10.1016/j.applanim.2010.08.004>
- Paladino, F. V., O'Connor, M. P., & Spotila, J. R. (1990). Metabolism of leatherback turtles, gigantothermy, and thermoregulation of dinosaurs. *Nature*, 344(6269), 858–860. <https://doi.org/10.1038/344858a0>
- Panagiota, M., & Valakos, E. D. (1992). Contribution to the thermal ecology of *Testudo marginata* and *T. hermanni* (Chelonia: Testudinidae) in semi-captivity. *Herpetological Journal*, 2(2), 48–50.
- R Core Team (2016). *R: A language and environment for statistical computing*, 3rd ed. Vienna, Austria: R Foundation for Statistical Computing. Retrieved from <https://www.R-project.org>
- Revelle, W. (2016). *psych: Procedures for psychological, psychometric, and personality research*. Evanston, IL: Northwestern University. Retrieved from <https://CRAN.R-project.org/package=psych>
- Rivera-Vélez, N., & Lewis, A. R. (1994). Threshold temperatures and the thermal cycle of a heliothermic lizard. *Journal of Herpetology*, 28(1), 1. <https://doi.org/10.2307/1564672>
- Rohatgi, A. (2017). *WebPlotDigitizer*. 3rd edn. Retrieved from <http://arohatgi.info/WebPlotDigitizer>.
- Schmidt-Nielsen, K., & Bentley, P. J. (1966). Desert tortoise *Gopherus agassizii*: Cutaneous water loss. *Science*, 154(3751), 911. <https://doi.org/10.1126/science.154.3751.911>
- Seebacher, F. (2003). Dinosaur body temperatures: The occurrence of endothermy and ectothermy. *Paleobiology*, 29(1), 105–122. [https://doi.org/10.1666/0094-8373\(2003\)029<0105:DBTTOO>2.0.CO;2](https://doi.org/10.1666/0094-8373(2003)029<0105:DBTTOO>2.0.CO;2)
- Seebacher, F., & Franklin, C. E. (2001). Control of heart rate during thermoregulation in the heliothermic lizard *Pogona barbata*: Importance of cholinergic and adrenergic mechanisms. *Journal of Experimental Biology*, 204(24), 4361–4366.
- Seebacher, F., Grigg, G. C., & Beard, L. A. (1999). Crocodiles as dinosaurs: Behavioural thermoregulation in very large ectotherms leads to high and stable body temperatures. *Journal of Experimental Biology*, 202(1), 77–86.
- Shekine, J., Turnbull, L. A., Cherubini, P., de Jong, R., Baxter, R., Hansen, D., ... Schaeppman-Strub, G. (2015). Primary productivity and its correlation with rainfall on Aldabra Atoll. *Biogeosciences Discussion*, 12(2), 981–1013. <https://doi.org/10.5194/bgd-12-981-2015>
- Slip, D. J., & Shine, R. (1988). Thermoregulation of free-ranging diamond pythons, *Morelia spilota* (Serpentes, Boidae). *Copeia*, 1988(4), 984–995. <https://doi.org/10.2307/1445722?ref=search-gateway:5163e322701902f21d23da697ad3dba9>
- Swingland, I. R., & Frazier, J. G. (1979). The conflict between feeding and overheating in the Aldabran giant tortoise. In C. J. Amlaner, & D. W. McDonald (Eds.), *A handbook of biotelemetry and radio tracking* (pp. 611–615). Oxford: University of Oxford.
- Tosini, G., & Menaker, M. (1995). Circadian rhythm of body temperature in an ectotherm (*Iguana iguana*). *Journal of Biological Rhythms*, 10(3), 248–255. <https://doi.org/10.1177/074873049501000307>
- Turnbull, L. A., Ozgul, A., Accouche, W., Baxter, R., ChongSeng, L., Currie, J. C., ... Bunbury, N. (2015). Persistence of distinctive morphotypes in the native range of the CITES-listed Aldabra giant tortoise. *Ecology and Evolution*, 5(23), 5499–5508. <https://doi.org/10.1002/ece3.1764>
- Van Damme, R., Bauwens, D., & Verheyen, R. F. (1991). The thermal dependence of feeding behaviour, food consumption and gut-passage time in the lizard *Lacerta vivipara* Jacquin. *Functional Ecology*, 5(4), 507–517. <https://doi.org/10.2307/2389633?ref=search-gateway:7e8343f784f7b2edf8269aec5ede75c2>
- Wickham, H. (2016). *ggplot2: Elegant graphics for data analysis*. Cham: Springer. <https://doi.org/10.1007/978-3-319-24277-4>
- Willmer, P., Stone, G., & Johnston, I. A. (2005). *Environmental Physiology of Animals* (pp. 1–779). Malden, MA: Blackwell Publishing.
- Woodbury, A. M., & Hardy, R. (1948). Studies of the desert tortoise, *Gopherus agassizii*. *Ecological Monographs*, 18(2), 145–200. <https://doi.org/10.2307/1948638>
- Zimmerman, L. C., & Tracy, C. R. (1989). Interactions between the environment and ectothermy and herbivory in reptiles. *Physiological Zoology*, 62, 374–409. <https://doi.org/10.2307/30156176>
- Zimmerman, L. C., Zimmerman, L. C., O'Connor, M. P., O'Connor, M. P., Bulova, S. J., Bulova, S. J., ... Salice, C. J. (1994). Thermal ecology of desert tortoises in the Eastern Mojave Desert: Seasonal patterns of operative and body temperatures, and microhabitat utilization. *Herpetological Monographs*, 8, 45–59. <https://doi.org/10.2307/1467069>
- Zuur, A., Ieno, E. N., Walker, N., Saveliev, A. A., & Smith, G. M. (2009). *Mixed effects models and extensions in ecology with R*.



New York, NY: Springer Science & Business Media. <https://doi.org/10.1007/978-0-387-87458-6>

## SUPPORTING INFORMATION

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