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Seasonal changes in energy expenditure, body temperature and activity patterns in llamas (*Lama glama*)

Alexander Riek¹, Lea Brinkmann¹, Matthias Gauly², Jurcevic Perica³, Thomas Ruf³, Walter Arnold³, Catherine Hambly⁴, John R. Speakman^{4,5} & Martina Gerken¹

Mammals typically keep their body temperature (T_b) within a narrow limit with changing environmental conditions. There are indications that some wild ungulates can exhibit certain forms of energy saving mechanisms when ambient temperatures are low and/or food is scarce. Therefore, the aim of the study was to determine if the llama, one of the most extensively kept domestic livestock species, exhibits seasonal adjustment mechanisms in terms of energy expenditure, T_b and locomotion. For that purpose llamas ($N = 7$) were kept in a temperate habitat on pasture. Locomotor activity, T_b (measured in the rumen) and the location of each animal were recorded continuously for one year using a telemetry system. Daily energy expenditure was measured as field metabolic rate (FMR). FMR fluctuated considerably between seasons with the lowest values found in winter ($17.48 \pm 3.98 \text{ MJ d}^{-1}$, $402 \text{ kJ kg}^{-0.75} \text{ d}^{-1}$) and the highest in summer ($25.87 \pm 3.88 \text{ MJ d}^{-1}$, $586 \text{ kJ kg}^{-0.75} \text{ d}^{-1}$). Llamas adjusted their energy expenditure, T_b and locomotor activity according to season and also time of day. Thus, llamas seem to have maintained the ability to reduce their energy expenditure and adjust their T_b under adverse environmental conditions as has been reported for some wild ungulates.

Endothermic mammals usually have to keep their body temperature (T_b) in a narrow limit of 37–39 °C with changing ambient temperatures (T_a)^{1,2}, which comes at a high energetic cost. Endotherms therefore are confronted with the challenge to maintain a comparatively high energy intake in the face of seasonal variation in food quantity and quality to keep the T_b in that aforementioned narrow limit³. Many small animals (<1000 g) counter this challenge by employing energy saving mechanisms such as torpor or hibernation i.e. decreasing metabolic rate and T_b substantially, during times of adverse environmental conditions^{4–6}. With the exceptions of bears and badgers, larger animals (>5000 g) are not known to use these energy saving mechanisms. Recent studies, however, revealed that some ungulates can decrease their metabolic rate and T_b to some degree and thus adjust their daily energy expenditure (DEE) according to season. So far, this has been shown for the red deer (*Cervus elaphus*)⁷, Przewalski horses (*Equus przewalski*)⁸, Alpine ibex (*Capra ibex*)⁹ and Shetland ponies (*Equus caballus*)¹⁰. Additionally, less comprehensive studies mainly based on respirometry of captive specimens have suggested seasonal variation in the metabolism of several cervid species (reviewed in Mauget *et al.*)¹¹ and other ruminants such as muskoxen (*Ovibos moschatus*)¹² or Arabian oryx (*Oryx leucoryx*)¹³.

In its native region of South America the llama is used as a wool and meat producer, but also as a beast of burden. The llama as such is an integral part of the life of the rural population in the High Andes and contributes to a large extent to their overall income¹⁴. In the last two to three decades the llama also gained popularity in North America, Europe and Australia, mainly as pet animal, wool producer or for landscape management purposes¹⁵. However there is still a lack of scientific literature especially in the fields of nutrition and energetics^{16–18}.

¹Department of Animal Sciences, University of Göttingen, Albrecht-Thaer-Weg 3, 37075, Göttingen, Germany.

²Faculty of Science and Technology, Free University of Bolzano, Bolzano, Italy. ³Research Institute of Wildlife Ecology, Department of Integrative Biology and Evolution, University of Veterinary Medicine, Vienna, Austria. ⁴Institute of Biological and Environmental Sciences, University of Aberdeen, Aberdeen, AB24 2TZ, UK. ⁵Institute of Genetics and Developmental Biology, State Key Laboratory of Molecular Developmental Biology, Chinese Academy of Sciences, 100101, Beijing, PR China. Correspondence and requests for materials should be addressed to A.R. (email: arielk@uni-goettingen.de)

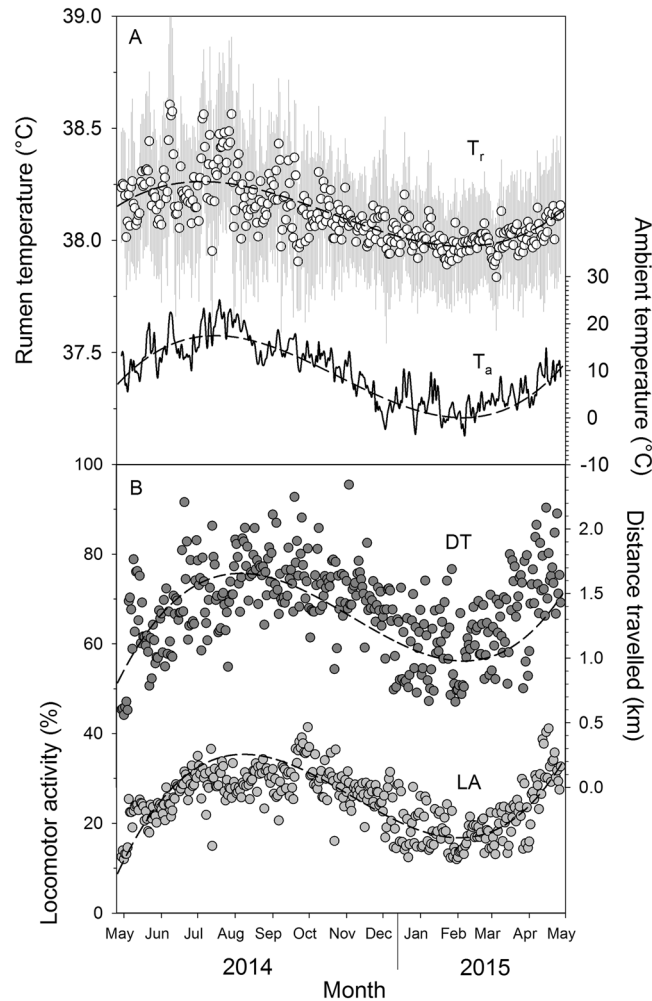


Figure 1. Yearly rhythm of rumen and ambient temperature as well as locomotor activity and distances travelled in llamas. Data are daily average rumen temperature (T_r , white dots with average minima and maxima, $N = 7$) and daily average ambient temperature (T_a , black solid line; **A**) as well as daily average locomotor activity (LA, light grey dots) and distances travelled (DT, dark grey dots; **B**) in adult non-pregnant llama dams during one year (29 April 2014–28 April 2015). Black dashed lines are best fit models for each parameter in A and B (see text for model equations). Values for T_r , LA and DT are means of seven animals over a total of 2321 recording days.

Furthermore so far there are no studies investigating seasonal adjustment mechanisms with regard to energy expenditure and T_b in South American camelids. In this regard the llama is a particular well suited model animal because it is one of the most robust livestock species, which can be described as a primary population¹⁹.

Therefore, the aim of our study was to determine if the llama, one of the most extensively kept livestock species, exhibits seasonal and ultradian adjustment mechanisms in terms of energy expenditure, T_b and locomotion under Central European climatic conditions. In particular we wanted to determine if differences in energy expenditure during different seasons exist and how T_a is correlated with T_b and locomotor activity (LA).

Results

Climatic conditions. The climatic conditions during the year of our study (29 Apr 2014–28 Apr 2015) were in the normal range expected for this Central European location. The T_a fluctuated substantially over the course of the study with daily averages ranging from 25.0 ± 9.28 °C in summer to -3.5 ± 2.1 °C in winter (Fig. 1A). The course of the average daily annual T_a was best described by a third degree polynomial (T_a , °C = $4E-06x^3 - 0.002x^2 + 0.27x + 7.97$, $R^2 = 0.81$, $P < 0.001$; where x is the day of study). Rainfall occurred on 157 of the 365 study days. The highest daily rainfall occurred on 29 July with 34.6 mm. The total annual rainfall over the one year study period was 867 mm. During winter there were 32 days with snow heights over 1 cm and during the entire study period there were 69 days with frost. The mean daily averages, mean daily maxima and mean daily minima for T_a and RH during the time of the four FMR measurements in spring, summer, autumn and winter are presented in Table 1.

Body temperature. The daily mean T_r during the entire study of one year was 38.11 ± 0.41 °C. There was a distinct fluctuation of daily mean T_r over the course of the study with the lowest daily mean T_r of 37.64 ± 0.36 °C

Variable		Spring	Summer	Autumn	Winter
Daily ambient temperature	(°C)	9.90 ± 2.15	17.53 ± 1.94	7.98 ± 2.00	0.73 ± 1.35
Daily minimum	(°C)	2.18 ± 2.90	11.52 ± 1.06	3.83 ± 2.86	-1.40 ± 2.41
Daily maximum	(°C)	17.52 ± 3.98	25.01 ± 4.20	13.39 ± 2.35	3.01 ± 1.87
Daily range	(°C)	15.33 ± 5.79	13.49 ± 4.48	9.56 ± 3.96	4.41 ± 2.82
Daily relative humidity	(%)	78.60 ± 9.17	81.62 ± 4.63	96.10 ± 3.74	97.28 ± 3.01
Daily minimum	(%)	48.28 ± 17.47	50.72 ± 10.97	77.80 ± 11.46	87.00 ± 8.80
Daily maximum	(%)	102.38 ± 4.08	101.22 ± 2.43	104.78 ± 1.22	103.85 ± 2.15
Daily range	(%)	54.01 ± 18.49	50.50 ± 12.59	26.98 ± 12.03	16.75 ± 9.08

Table 1. Ambient temperature and relative humidity during two weeks at four different seasons (spring, summer, autumn and winter, see text for details) under central European climatic conditions. Values are means ± sd.

Variable		Spring	Summer	Autumn	Winter	SEM	F-value	Season P-value
Body mass	(kg)	150.4 ^d	154.4 ^c	163.7 ^a	159.1 ^b	9.13	36.56	<0.001
Body condition score	(points)	3.11 ^b	3.14 ^{bc}	3.38 ^a	3.36 ^{ac}	0.21	6.25	0.004
Field metabolic rate	(MJ d ⁻¹)	18.33 ^{bc}	25.87 ^a	20.70 ^{abc}	17.48 ^c	1.53	6.14	0.005
	(kJ kg ^{-0.75} d ⁻¹)	401 ^b	586 ^a	427 ^b	402 ^b	37.5	5.87	0.006
Total body water	(%)	60.97	63.49	63.94	63.92	1.61	2.31	0.112
Total water intake	(L d ⁻¹)	6.55 ^b	9.17 ^a	7.55 ^{ab}	4.22 ^c	0.44	23.35	<0.001
Daily activity	(%)	32.80 ^a	29.32 ^{ab}	26.31 ^b	16.08 ^c	1.03	54.90	<0.001
Distance travelled	(km d ⁻¹)	1.64 ^b	1.74 ^a	1.50 ^c	1.01 ^d	0.03	185.6	<0.001
Daily body temperature	(°C)	38.55 ^{ab}	38.62 ^a	38.53 ^b	38.40 ^c	0.03	20.58	<0.001
Daily minimum	(°C)	37.49 ^{ab}	37.44 ^{ac}	37.60 ^b	37.40 ^c	0.05	17.45	<0.001
Daily maximum	(°C)	38.64 ^{ab}	38.75 ^b	38.52 ^a	38.47 ^a	0.06	9.39	0.001
Daily range	(°C)	1.15 ^{ab}	1.30 ^a	0.92 ^b	1.06 ^b	0.08	7.94	0.002

Table 2. Average physiological and behavioural variables in llama dams (n = 7) during two weeks at four different seasons (spring, summer, autumn and winter, see text for details) under central European climatic conditions. Values are LS-Means with the corresponding SEM, F- and P-value. ^{a,b,c,d}Means within a row not sharing the same superscript differ by $P < 0.05$.

recorded in winter and the highest of 38.86 ± 0.48 °C in summer. Similar to T_a , the course of the average daily T_r over the study period was best described by a third degree polynomial (T_b , °C = $5E-08x^3 - 3E-05x^2 + 0.003x + 38.18$, $R^2 = 0.54$, $P < 0.001$; Fig. 1A, where x is the day of study). The lowest absolute T_r recorded was 36.02 °C and the highest was 39.98 °C. During spring and summer daily mean minimum T_r ($T_{r,min}$) and daily mean maximum T_r ($T_{r,max}$) varied considerably. Towards autumn and winter however these high variations decreased. This trend was also evident when considering T_r during the four FMR measurements, i.e. the daily T_r range (amplitude) was significantly lower in winter and autumn than during summer (Table 2). Over the entire study period of one year, daily $T_{r,min}$ was lowest in December (37.08 °C) and highest in August (37.89 °C). Similarly, lowest daily $T_{r,max}$ were recorded in March (38.27 °C) and highest in August (39.49 °C). Typically $T_{r,max}$ and $T_{r,min}$ occurred around late afternoon and early morning, respectively over the entire study including the four FMR measurement points (Fig. 2). Average daily T_r followed the T_a pattern throughout the year (Fig. 1A) and had a highly significant positive relationship with average daily T_a (Fig. 3A). Average daily T_r was also highly correlated with daily maximum ($r = 0.75$, $P < 0.001$) and minimum T_a ($r = 0.74$, $P < 0.001$).

Energy expenditure and water turnover. Energy expenditure in llamas measured as field metabolic rate (FMR) varied considerably between the four measurements, i.e. at different seasons (Table 2, Fig. 4). In winter, when T_a 's were on average 0.73 ± 1.35 °C but decreased on some days to values below -6 °C, the FMR was significantly lower (17.38 ± 3.98 MJ d⁻¹, 402 kJ kg^{-0.75} d⁻¹) compared to values measured during summer (25.87 ± 3.88 MJ d⁻¹, 586 kJ kg^{-0.75} d⁻¹) when T_a 's were on average 17.53 ± 1.94 °C but reached on some days over 30 °C. Thus DEE was about 30% lower in winter compared to summer. FMR was also higher ($P < 0.001$) in summer compared to spring but there were no differences ($P > 0.05$) between summer and autumn or winter and autumn (Table 2). Furthermore, mass independent FMR had a significant positive relationship with T_a ($R^2 = 0.90$, $P < 0.001$, Fig. 4). After the effect of body mass was removed, average residual variation of DEE explained to some degree average T_b ($r = 0.78$, $P < 0.001$), LA ($r = 0.37$, $P < 0.05$) and distances travelled ($r = 0.63$, $P < 0.01$).

Total body water (TBW) ranged from 57 to 69% of body mass between individual animals but average TBW did not differ between seasons (Table 2). However, total water intake (TWI) differed significantly between seasons. In summer animals ingested more than double the amount of water (9.17 ± 1.65 l d⁻¹) compared to winter (4.22 ± 0.33 l d⁻¹; Table 2). The TWI was also higher ($P < 0.001$) in summer than in spring and in spring TWI was

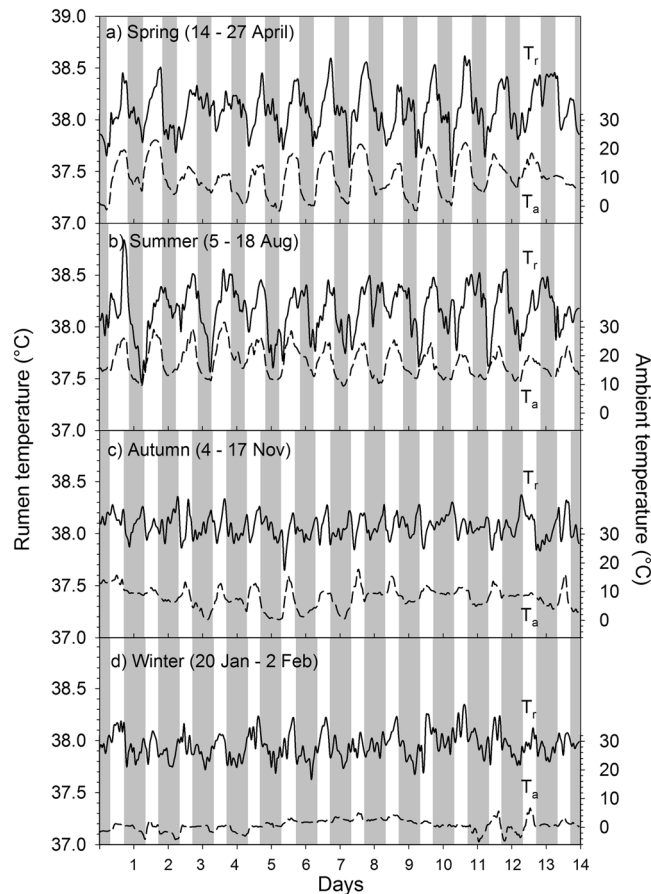


Figure 2. Rumen temperature in llamas. Data show the diurnal rhythm of ambient temperature (T_a , dashed line) and average rumen temperature (T_r , solid line) in adult non-pregnant llama dams ($N = 7$) during two weeks in four different seasons (spring, summer, autumn, winter) under central European climatic conditions. Grey shaded areas indicate night-phase. Axis range for T_r and T_a is the same for each season.

higher than in winter. However, no differences ($P > 0.05$) were detected between autumn and summer or autumn and spring.

Locomotor activity and distances covered. During our study mean daily LA followed a similar pattern as T_a (Fig. 1A) with the lowest daily average LA recorded in winter and the highest in summer (Fig. 1B). The course of average daily LA over the study period of one year was therefore, similar to T_a , best described by a third degree polynomial ($LA, \% = 5E-06x^3 - 0.003x^2 + 0.42x + 13.48$, $R^2 = 0.56$, $P < 0.001$, where x is the day of study). Daily average LA had thus a significant positive relationship with daily average T_a (Fig. 3B). This trend was also evident during the FMR measurement periods. The daily average LA in winter was with only $16.1 \pm 1.4\%$ significantly ($P < 0.001$) lower compared to autumn ($26.3 \pm 1.2\%$), summer ($29.3 \pm 2.6\%$) and spring ($32.8 \pm 4.4\%$, Table 2). As can be seen in Fig. 5, the diurnal LA rhythm varied between seasons, with many irregular activity peaks throughout the day in spring and summer, distinct daily peaks in autumn and infrequent activity during the measurement days in winter.

The average daily distances covered by the animals had a positive significant relationship with LA ($LA, \% = 14.50 * \text{Distance travelled, km} + 4.94$, $R^2 = 0.58$, $P < 0.001$; not shown). Similar to daily activity, animals covered on average significantly ($P < 0.001$) more distance in summer compared to winter and thus the model best describing the course of daily distances covered during the time of the study was again a third degree polynomial (Daily distances covered, km = $2E-07x^3 - 0.0001x^2 + 0.019x + 0.79$, $R^2 = 0.38$, $P < 0.001$; Fig. 1B, where x is the day of study). During the FMR measurements average daily distances covered differed significantly between each season and were highest in summer and lowest in winter (Table 2). Over the entire year the average daily distances covered per day ranged from 0.66 ± 0.12 km to 2.8 ± 0.43 km.

Discussion

Our study is the first in a camelid that combines the doubly labelled water method for measuring DEE with a telemetry system measuring locomotion and T_b , as well as estimating the distances covered by GPS. Furthermore, our study presents the first continuous long-term T_b and activity measurements in a South American camelid.

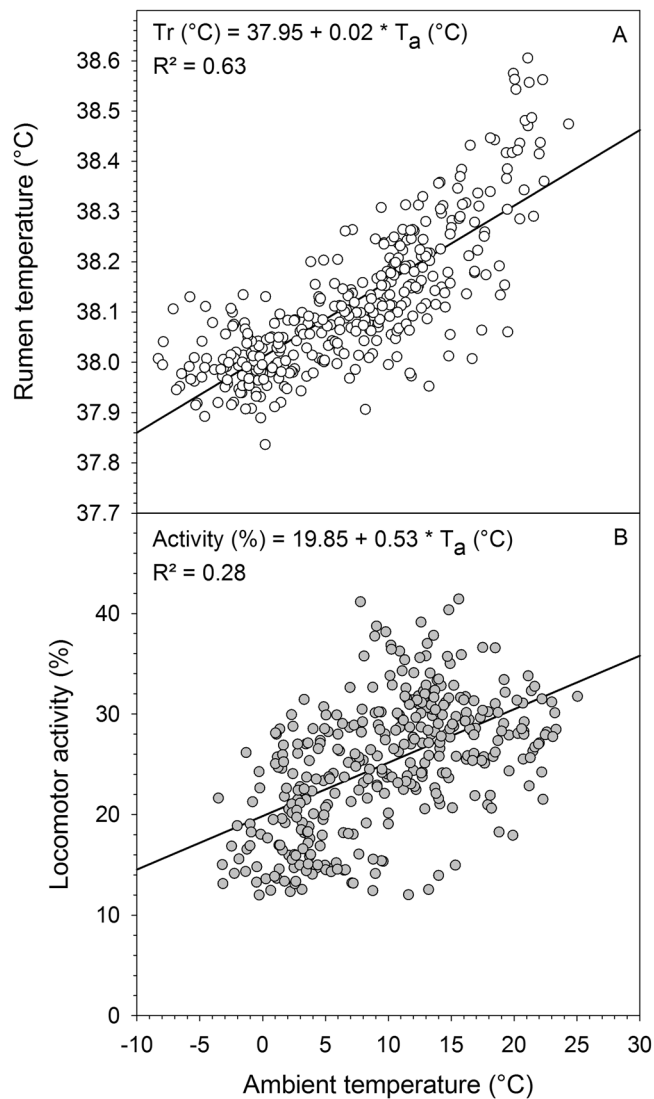


Figure 3. Rumen temperature and locomotor activity in llamas in relation to ambient temperature. Panels show the relationship between average daily ambient temperature (T_a) and average daily rumen temperature (T_r , **A**) and average daily locomotor activity (**B**) in adult non-pregnant llama dams ($N = 7$) during one year under Central European conditions.

The data show that llamas seem to have maintained the ability to reduce their DEE and activity and to a certain degree adjust their T_b according to season and time of the day as has been reported for some wild ungulates.

Some studies on wild and domestic ungulates have reported substantial reductions in DEE during adverse environmental conditions^{7,9,10,20}. In our study on llamas, we found similar adaptations in DEE. The DEE of the animals in our study varied considerably between seasons with FMR values in summer being on average 30% higher compared to winter (Table 2, Fig. 4). This can be to some degree explained by the higher activity levels during summer (Table 2, Fig. 5) when animals were grazing and also by a higher food intake. For example studies on alpacas²¹ and cervids^{7,22–24} have shown that the seasonality of metabolism is linked to the level of food intake.

The T_a fluctuations in our study with an average daily maximum of 25.0 ± 4.2 °C in summer and a daily minimum of -1.4 ± 2.41 °C in winter are typical for temperate regions like our Central European study site (Table 2). Compared to the High Andes of South America, the native region of the llama, where T_a can fluctuate by 40–50 °C per day, the seasonal and daily T_a fluctuations in our study can be considered as moderate. Nevertheless, during lower T_a 's, e.g. in winter, thermoregulatory costs for endothermic animals usually increase to keep the T_b within a narrow limit^{1,2} and thus resulting in an increased DEE²⁵. In our study however we observed the opposite, i.e. decreased DEE in winter when T_a 's were low and increased DEE in summer when T_a 's were high. The thermal neutral zone, which is the range of T_a in which T_b in an animal is only regulated by the control of sensible heat loss and thus does not require additional energy for thermoregulation^{26,27}, has so far not been directly measured in the llama. However data from guanacos (*Lama guanicoe*) which is the wild ancestor of the llama, suggest that the breadth of the thermal neutral zone seems to be quite large with a lower critical temperature being in the range of -10 to -15 °C and an upper critical temperature of about 20 °C^{27,28}. This large thermal neutral zone is most likely

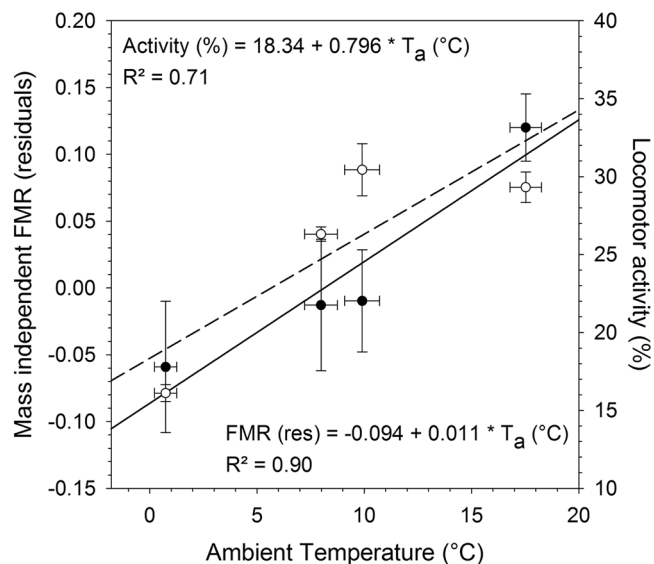


Figure 4. Field metabolic rate (FMR) and locomotor activity in llamas. Data are average FMR (●) and activity (○) values of llama dams (N = 7) with the corresponding fit lines (FMR: solid line; Activity: dashed line) measured at four different seasons, i.e. ambient temperature ranges. FMR values are given as residuals from the regression of FMR on body mass.

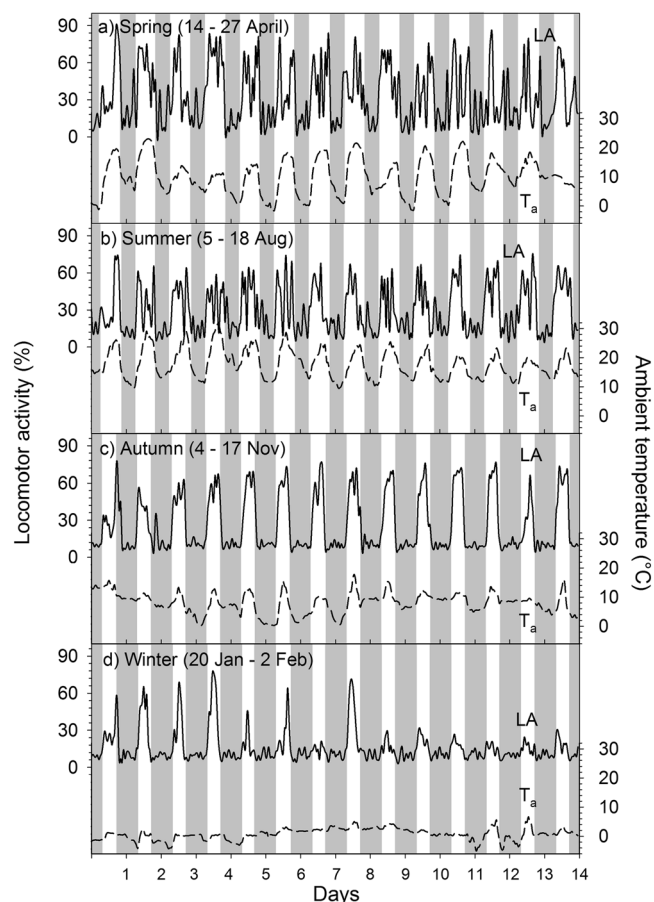


Figure 5. Locomotor activity in llamas. Data show the diurnal rhythm of ambient temperature (T_a , dashed line) and mean daily locomotor activity (LA, solid line) in adult non-pregnant llama dams (N = 7) during two weeks in four different seasons (spring, summer, autumn, winter) under central European climatic conditions. Grey shaded areas indicate night-phase. Axis range for LA and T_a is the same for each season.

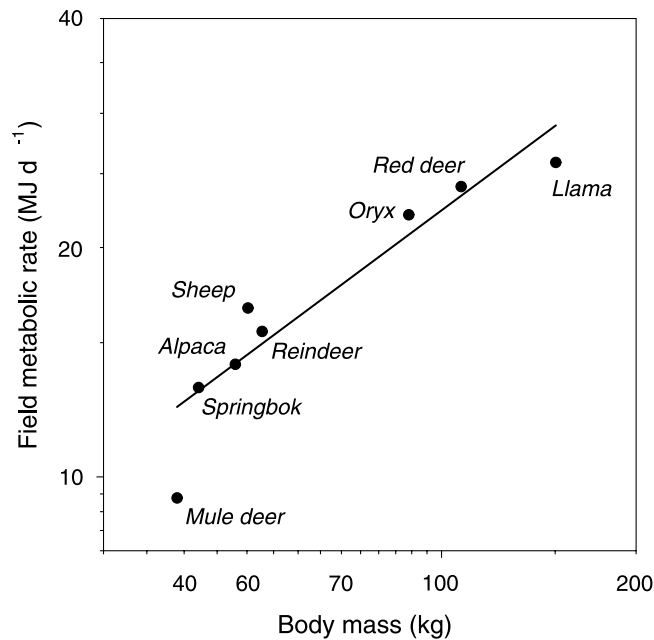


Figure 6. Relationship between daily energy expenditure and body mass in artiodactyls. Data are measurements of field metabolic rate (FMR) using the doubly labelled water method and body mass (BM) for eight artiodactyls (including data on llamas from the present study). The phylogenetically controlled allometric relationship is $FMR (MJ d^{-1}) = 1.23 BM^{0.63 \pm 0.12}$ ($F_{1,7} = 26.2$, $P < 0.001$, $R^2 = 0.81$, $ML \lambda = 0.64$) (see text for details).

due to the ability of guanacos and llamas to use peripheral vasoconstriction and local heterothermy. Furthermore guanacos and llamas employ behavioural adjustments to minimise heat loss during cold exposure²⁹. In endothermic animals the energetic costs for thermoregulation increase when T_a decreases below the lower critical temperature or increase above the upper critical temperature, thus defining the thermal neutral zone of an animal¹. It can be assumed that our llamas were within their thermal neutral zone during the cooler seasons, i.e. winter, autumn and spring when T_a did not exceed 20 °C or fell below –10 to –15 °C (Table 1). Thus suggesting that no additional energy was needed for thermoregulation during these seasons since the decrease of T_r during winter shifted the zone of thermoneutrality to a lower temperature indicating that our animals were under conditions of thermoneutrality for most of the winter. During the summer months however when T_a exceeded 20 °C, the DEE was significantly higher compared to winter or spring. Furthermore LA also differed between summer and winter. Therefore the lower DEE measured in winter compared to summer is most likely due to a reduction in LA and T_b , evidenced by a positive and significant relationship between FMR and LA as well as T_b . Furthermore animals were within their thermal neutral zone and thus did not need the extra cost for thermoregulation. Conversely, the increased DEE in summer can be explained by an increased LA and an increased cost for thermoregulation because the T_a most likely exceeded the upper critical temperature at least on some days during the summer measurement. Similar results have been also found for red deer⁷, Przewalski horses⁸ and ibexes⁹ indicating that during low T_a animals reduce their activity and thus lower their energy expenditure. This has been also observed in smaller mammals such as red squirrels (*Tamiasciurus hudsonicus*)³⁰ and least weasels (*Mustela nivalis*)³¹ living in temperate or arctic regions. Interestingly other studies of animals living in warmer regions such as kangaroo rats (*Dipodomys merriami*) or white footed mice (*Peromyscus leucopus*) do not show this kind of adaptation^{25, 32–34}.

Very few studies on artiodactyls are available that measured DEE using the doubly labelled water method. So far DEE has been measured in eight artiodactyls (Mule deer, *Odocoileus hemionus*³⁵; reindeer, *Rangifer tarandus*³⁶; springbok, *Antidorcas marsupialis*³⁷; red deer, *Cervus elaphus*³⁸; Arabian oryx, *Oryx leucoryx*¹³; sheep, *Ovis aries*³⁹; alpacas, *Lama pacos*⁴⁰). Comparing average DEE between these species, llamas have a DEE of 28.9 MJ/d predicted from a phylogenetic corrected regression equation for artiodactyls ($FMR, MJ d^{-1} = 1.23 BM^{0.63 \pm 0.12}$, Fig. 6). This is about 10% higher than the actual DEE measured during summer (25.9 MJ/d), which was the highest average DEE measured during the four measurement points of the present study and more than 30% higher than the value measured in winter (17.48 MJ/d) which was the lowest average DEE measured during our study. This suggests that llamas seem to have on average a lower DEE relative to other artiodactyls (with the exception of the mule deer). Similar suggestions have been also made for other camelids¹⁸. Interestingly, the exponent of the phylogenetic corrected regression equation for the relationship between FMR and body mass in artiodactyls (0.63, 95% CI 0.32–0.91) is very close to the one found for Metatheria (0.60)⁴¹. However these results need to be treated with caution as only very few data from artiodactyls are available and thus no definite conclusions can be reached or valid comparisons be made until further systematic studies on other artiodactyl species are available. Furthermore, predicting values for missing species that have not been measured from phylogenetic regression

equations, as it is sometimes done in comparative analysis, are likely to be incorrect as the fit-lines are phylogenetically controlled and thus will not account for the phylogenetic history of the missing species.

The TWI calculated from isotope turnover rates differed between seasons (Table 2). During summer animals ingested with 9.17 l/d on average more than double the amount of water compared to winter (4.22 l/d). Most likely, these differences can be attributed to an increased drinking water ingestion during summer when animals displayed higher physical activities and the average daily maximum T_a was above 25 °C (Table 1). Furthermore animals consumed more pasture in summer, containing a higher percentage of water compared to hay, which was the main food source in winter. As explained before, when T_a increases above the upper critical temperature it triggers energy dependent thermoregulative mechanisms for dissipation of metabolic heat and thus keeping T_b within a narrow limit⁴². Therefore, substantial amounts of water will be dissipated with increasing T_a especially via the thermal windows at the ventral regions of the body in llamas, resulting in increased water ingestion and water turnover⁴³. Additionally, the increased DEE of the animals during the summer months would have resulted in a greater production of metabolic water. These changes in water turnover between seasons however did not affect the TBW. The average TBW of 63.1% of body mass of all animals did not differ between seasons (Table 2) and was in the range of reported TBW values for ungulates (for comparison see Table 2 in ref. 10).

The daily T_b fluctuations, i.e. T_b decreasing during the night and increasing during the day were more pronounced in summer and spring when T_a fluctuations were higher, than in winter and autumn when T_a fluctuations were lower (Fig. 1A, Table 2) suggesting that animals were following a shallow daily hypometabolism. This adaptive mechanism to save energy by reducing the metabolic rate has been reported for many smaller animals (body mass < 5 kg) during the 24 h rhythm of rest and activity^{6,44}. For humans and larger animals similar results have been found^{7–9,45}. In our study average daily T_b amplitudes during the FMR measurement points were highest in summer (1.3 °C) and lowest in autumn (0.92 °C, Table 2). Similar average T_b amplitudes have been reported for alpacas (1.5 °C)⁴⁶, angora goats (1.4 °C)⁴⁷, blesbok (1.4 °C)⁴⁶, pronghorn (1.0 °C)⁴⁸ and impala (1.1 °C)⁴⁹. However our results are daily averages of seven animals over two weeks. Looking at the individual daily T_b variations over the study year the highest amplitudes found ranged from 1.8 to 3.2 °C and occurred all in summer (Fig. 1). Higher T_b amplitudes of 6–7 °C have been reported for springboks⁵⁰ and camels (*Camelus dromedarius*)⁵¹. Nevertheless the T_b amplitudes of our animals were higher than the normal circadian variations in T_b for llamas (37.5–38.6 °C)⁵² possibly using adaptive heterothermy to some degree to reduce DEE. Furthermore in our study daily T_b variations could be observed over the entire study period following the daily photoperiod as has also been reported for horses^{8,45,53}, red deer⁷ and ibex⁹. Usually the 24 h T_b rhythm is as such an endogenous rhythm and is synchronized by the external ‘Zeitgeber’ independent of exogenous factors. This could explain why on some days T_b decreased at night when T_a did not but instead stayed rather constant, e.g. during some winter days (Fig. 2d). However, T_b was correlated with T_a throughout the study (Fig. 3A) and thus following the general rhythm of the daily T_a cycle. Nevertheless animals sharply increased their activity in the morning possibly resulting in an increase of T_b . During the lower T_b variations in winter and autumn, i.e. when T_a was generally low, animals might have shifted from a short daily hypometabolism in spring and summer to a more intense hypometabolism to save energy, as has been recently shown for Shetland ponies¹⁰. Thus the decreased T_b amplitude during the cooler seasons could be explained by an adaptation to save energy. Similar observations have been recently described in grey kangaroos⁵⁴ and the oryx⁵⁵. Furthermore, the lowering of the T_b during night hours might reduce the energetic cost for thermoregulation by increasing the capacity to store heat during hot days^{3,50}. This adaptive heterothermy has been already demonstrated in a variety of other ungulates such as the eland⁵⁶, oryx^{42,55}, giraffe⁵⁷, Arabian sand gazelle⁵⁸ and Thomson’s as well as Grant’s gazelles⁵⁹. However other studies challenge these findings^{60,61} or suggest that heterothermy in large mammals could be a sub-optimal response to environmental challenges⁴⁶. Nevertheless in general llamas seem to lower their T_b under energetic constraints which must involve some trade-offs that are less energy demanding than keeping the T_b constant^{62,63}.

In conclusion we show that llamas, one of the most extensively kept livestock species, are able to reduce their energy expenditure under adverse environmental conditions by reducing their activity and adjusting their daily T_b variation. Thus llamas show distinct seasonal acclimatization similar to wild ungulates.

Methods

Animals and study site. The study was conducted at the research farm Relliehausen (41°46′ N, 9°41′ E) of the Department of Animal Sciences at the University of Göttingen (Göttingen, Germany) and involved seven non pregnant llama dams (age: 3–13 years, body mass: 113–174 kg). The measurements were carried out continuously for one year (29 Apr 2014–28 Apr 2015). Animals were kept on a pasture (3 ha) within a herd of 15 llama dams. All llamas had free access to a barn providing shelter from wind and rain. On pasture food consisted of natural vegetation and a mineral supplement provided by a salt lick (Eggersmann Mineral Leckstein, Heinrich Eggersmann GmbH & Co KG, Rinteln, Germany). Hay and water was available *ad libitum* for all animals throughout the experiment.

Measurements. *Climate.* The T_a (resolution: 0.0625 °C) and relative humidity (RH, resolution: 0.04%) were recorded continuously throughout the year with miniature data loggers at hourly intervals (i-Button, DS1923#F5, Maxim Integrated Products, Sunnyvale, CA, USA). Precipitation data were obtained from a nearby weather station at approx. 2 km distance to the farm.

Telemetry and body condition score. We equipped 7 animals with a telemetry system (GPS Plus-3 Store on Board collar, Vectronic Aerospace GmbH, Berlin, Germany) described in detail elsewhere⁶⁴. In brief, the telemetry system consists of two units, a ruminal unit (22 × 80 mm, 100 g) and a collar unit (450 g). The ruminal unit was administered *perorally* after animals were immobilized with an anaesthetic drug (Xylacin, Rompun®; Bayer HealthCare, Leverkusen, Germany, 1 ml/100 kg body mass). The ruminal unit measured T_r every 3 min, which

was transmitted via short-distance UHF link to a data logging system located in the collar unit⁹. Furthermore, locomotor activity (LA) was also recorded every 3 min with two different activity sensors. All data were recorded every 3 min for one year and stored in the collar unit and downloaded via a laptop. Additionally the position of each animal was recorded every 30 min using a GPS device located on the back of the collar (GPS Plus-3 Store on Board collar, Vecronic Aerospace GmbH, Berlin, Germany). The body condition score (BCS), a palpable and visual assessment of the degree of fatness (BCS scale: 0 = emaciated, 5 = obese), of individual animals was recorded during the four FMR measurement times according to the system described elsewhere⁶⁵.

Field metabolic rate. The FMR, TBW and TWI were determined during two weeks in the European summer (5–18 Aug 2014), autumn (4–17 Nov 2014), winter (20 Jan – 2 Feb 2015) and spring (14–27 April 2015), for each animal using the doubly labelled water (DLW) method^{66,67}. At the beginning and at the end of the FMR measurements, body mass was recorded for each llama using a mobile scale (Weighing System MP 800, resolution: 0.1 kg, Patura KG, Laudenbach, Germany) and a blood sample of 5 ml was drawn from the *Vena jugularis* of every animal to estimate the background isotopic enrichment of ²H and ¹⁸O in the body fluids (method D⁶⁸). After taking the background sample, each llama was injected intravenously with approximately 0.16 g of DLW per kg body mass, (65% ¹⁸O and 35% ²H). The individual dose of each llama was determined prior to the injection according to its body mass. The actual dose given was gravimetrically measured by weighing the syringe before and after administration to the nearest 0.001 g (Sartorius model CW3P1–150IG-1, Sartorius AG, Göttingen, Germany). The llamas were then held in the stable with no access to food or water for an 8-h equilibration period, after which a further 5 ml blood sample was taken. After dosing, additional blood samples were taken at 7 and 14 days to estimate the isotope elimination rates.

All blood samples were drawn into blood tubes containing sodium citrate. Whole blood samples were pipetted into 0.7 ml glass vials and stored at 5 °C until determination of ¹⁸O and ²H enrichment. Blood samples were vacuum distilled⁶⁹, and water from the resulting distillate was used to produce CO₂⁷⁰ and H₂⁷¹. The isotope ratios ¹⁸O: ¹⁶O and ²H: ¹H were analysed using gas source isotope ratio mass spectrometry (Isochrom μ G and Isoprime respectively, Micromass Ltd, Manchester, UK). Samples were run alongside five lab standards for each isotope (calibrated to the IAEA International standards: SMOW and SLAP) to correct delta values to ppm. Isotope enrichments were converted to values of CO₂ production using a two pool model as recommended for this size of animal⁷². We chose the assumption of a fixed evaporation of 25% of the water flux, since this has been shown to minimize error in a range of applications^{73,74}. Specifically carbon dioxide production rate (r_{CO_2}) per day in mols was calculated using equation A6 from Schoeller *et al.*⁷⁵. The daily amount of energy expended measured as FMR was calculated from carbon dioxide production by assuming a respiration quotient of 0.85. Total body water (mols) was calculated using the intercept method⁶⁷ from the dilution spaces of both oxygen and hydrogen under the assumption that the hydrogen space overestimates TBW by 4% and the oxygen-18 space overestimates it by 1%⁷⁵. The TWI (l/day) that consists of drinking water, preformed water ingested in food and metabolic water was estimated as the product of the deuterium space and the deuterium turnover rate⁷⁶.

Analysis. The measurements of T_r had non-physiological declines that could be attributed to the ingestions of water and cold food⁶⁴. These data points were removed by visually checking the raw data. In this cleaned data set, T_r values ranged from 36.02 to 39.98 °C. In total 2321 individual days were available for data analysis of LA and T_r . Hourly and daily means were calculated using R 3.3.2⁷⁷.

In order to compare FMR values with measured physiological variables during the time of FMR measurements a mixed model was used with animal as a random factor and season (i.e. FMR measurement periods) as a fixed factor to compare animals between seasons using the MIXED procedure in SAS version 9.2 (SAS, Inst. Inc., Cary, NC). The model residuals were normally distributed. Data are expressed as LS-Means or means \pm S.D where appropriate. Furthermore FMR values were also expressed as mass independent FMR by calculating the residuals of the regression of FMR on body mass. Daily distances between continuous GPS locations for each animal were calculated with the program package ‘geosphere’⁷⁸ in R⁷⁷. Furthermore Spearman correlations were calculated between different variables.

To test for the generality of the relation between body mass and FMR in artiodactyls, published data^{35–40} and our results, using the summer measurements, were assessed using the phylogenetic generalized least squares approach (PGLS)^{79,80} in order to account for the potential lack of independence between species, because of their shared evolutionary history^{81,82}. The statistical procedure has been described in detail elsewhere⁸³. In brief, the phylogeny was derived from a published mammalian supertree which includes 4510 species with updated branch lengths derived from dated estimates of divergence times⁸⁴. The supertree for mammals was pruned to include only the species of concern, i.e. artiodactyls ($n = 8$), using the ‘Analysis in phylogenetics and evolution’ (APE) package⁸⁵ and the ‘Analysis of evolutionary diversification’ (GEIGER) package⁸⁶ in R⁷⁷. The method of PGLS was implemented for the trait data using the ‘Comparative analyses of phylogenetics and evolution’ (CAPER) package⁸⁷ in R⁷⁷. PGLS analysis allows more flexibility than ordinary least square or independent contrasts methods through the use of a parameter (lambda, λ). The parameter λ is determined by maximum likelihood (ML) and can range between 0 (no phylogenetic signal, similar to ordinary least squares analysis) and 1 (pattern of trait data variation is fully explained by the phylogeny) and thus indicates how strong the phylogenetic signal for a certain trait or the relationship between two traits is. Intermediate values of λ indicate that the trait evolution is phylogenetically correlated, but does not follow fully a Brownian motion model⁸⁸. A more in depth description and further mathematical details on PGLS analysis can be found in detail elsewhere^{79,89,90}.

Ethics. Procedures performed in our study were in accordance with the German animal ethics regulations and approved by the State Office of Lower Saxony for Consumer Protection and Food Safety (Ref. No.: 33.4-42502-05-13A393).

Data availability. The data analysed during the current study are available from the corresponding author on reasonable request.

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Author Contributions

A.R. conceived the experiment, A.R. and L.B. conducted the experiment, M.G. contributed to the administering of the boli, W.A., T.R. and P.J. contributed to the analysis of the telemetry data. J.R.S. and C.H. conducted the doubly-labelled water analysis and calculated the FMR values. A.R. wrote the manuscript and all authors reviewed the manuscript.

Additional Information

Competing Interests: The authors declare that they have no competing interests.

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