



Research



Cite this article: Masoero G, Gencheva KG, Ioset N, Bersier L, Tettamanti F, Bize P. 2024 Shrinking Alpine chamois: higher spring temperatures over the last 27 years in Switzerland are linked to a 3 kg reduction in body mass of yearlings. *R. Soc. Open Sci.* **11**: 231295.

<https://doi.org/10.1098/rsos.231295>

Received: 30 August 2023

Accepted: 9 February 2024

Subject Category:

Ecology, conservation, and global change biology

Subject Areas:

ecology

Keywords:

climate change, *climwin*, ungulates, life stages, temperature, elevation

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Electronic supplementary material is available online at <https://doi.org/10.6084/m9.figshare.c.7095864>.

Shrinking Alpine chamois: higher spring temperatures over the last 27 years in Switzerland are linked to a 3 kg reduction in body mass of yearlings

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Although climate change is considered to be partly responsible for the size change observed in numerous species, the relevance of this hypothesis for ungulates remains debated. We used body mass measurements of 5635 yearlings (i.e. 1.5 years old) of Alpine chamois (*Rupicapra rupicapra*) harvested in September in the Swiss Alps (Ticino canton) from 1992 to 2018. In our study area, during this period, yearlings shrank by *ca* 3 kg while temperatures between May and July rose by 1.7°C. We identified that warmer temperatures during birth and the early suckling period (9 May to 2 July in the year of birth) had the strongest impact on yearling mass. Further analyses of year-detrended mass and temperature data indicate that this result was not simply due to changes in both variables over years, but that increases in temperature during this particularly sensitive time window for development and growth are responsible for the decrease in body mass of yearling chamois. Altogether, our results suggest that rising temperatures in the Alpine regions could significantly affect the ecology and evolution of this wild ungulate.

1. Introduction

As global changes induced by human activities accelerate, many species are undergoing phenotypic changes to adapt to their new environment [1], often measured by changes in distribution, abundance and phenology. An increasingly reported response to climate warming is the change in animal body size and shape [2–4], as morphology affects thermoregulation [5]. Indeed, a shrinking in body size leads to a larger surface-area-to-volume ratio, which, in turn, allows animals to dissipate heat more efficiently in warmer environments (Bergmann's rule; e.g. [6]). Global warming is also likely to impact body size due to changes in food availability and quality [7]. For example, the increase in frequency and length of summer droughts is expected to reduce forage quality and palatability for grazing species [8–10]. In mammals, high ambient temperature, together with the limits to the ability to dissipate heat, has also been previously shown to constrain maternal milk production and offspring growth [5,11], limiting adult size.

The size an individual reaches as an adult has critical consequences for reproductive success and overall fitness [12]. As adult size and mass are primarily determined by early growth conditions and juvenile size (e.g. [13]), it is fundamental to investigate how individuals are affected by climate in early-life stages [14]. Because juveniles usually have low energy reserves and must use a large portion of them for growth [15,16], they are sensitive to external biotic and abiotic factors [17–19]. In mammals, early growth is divided into three phases: gestation, suckling and post-weaning. However, the three phases might not be equally sensitive to climate warming. For example, the suckling phase might be the most sensitive, as offspring growth is the fastest during suckling, and the mother's milk production is constrained by high ambient temperatures [5,11].

We investigated the effect of weather conditions on the yearling size (i.e. 1.5 years of age) of Alpine chamois (*Rupicapra rupicapra*) using hunting data collected in the southern Swiss Alps from 1992 to 2018. We aimed to describe the body mass decrease in yearlings and identify which temporal period of their growth is most sensitive to temperature conditions. The Alpine chamois is the most abundant ungulate of the European Alps [20], and its morphology and physiology are adapted to high-elevation (cold) environmental conditions [21]. Accordingly, most previous studies on the Alpine chamois have revealed a gradual shrinking in chamois body mass (e.g. [19,22,23], but see [24]). As observed previously by these studies, we expected a temporal decrease in the body mass of yearling chamois and a negative relationship with the temperature during spring–summer in the first two years of life (e.g. [19]). To our knowledge, no study has yet precisely identified which developmental window during early life is most sensitive to climate warming and whether the shrinking in body mass over time is associated with an increase in temperature during this critical developmental window.

2. Material and methods

2.1. Study system

The Alpine chamois is a medium-sized ungulate with an asymptotic body mass of 22.3 kg in females and 29.6 kg in males [25–27]. In the Alps, chamois give birth in May [28] and suckle their offspring from May to July. These months also correspond to the growth peak of yearlings in their second year of life. Chamois are weaned between three and six months of age [29]. Vegetation in the Alps usually begins growing right after snow-melt in April, peaking in July, thus providing an abundant and protein-rich food source for a relatively brief period of time [30]. Alpine chamois are distributed over a broad altitudinal range (500–3100 m.a.s.l.; e.g. [31,32]) and generally follow resource availability and climate conditions [23,33].

Data on chamois size were extracted from hunting records in the southern Alps of Switzerland (canton Ticino) between 1992 and 2018. The study area covers 2700 km² with an elevation range of 250–2700 m.a.s.l. In Ticino, hunting starts at the beginning of September and is completed within 10 days. As required by Swiss law, animals were eviscerated by hunters, and their entrails were left at the kill site. Carcasses were then brought to checkpoints in the Ticino canton within 48 hours, where trained gamekeepers and wildlife wardens sexed and aged the animals before recording their weight with an electronic precision scale to the nearest 0.1 kg (eviscerated body weight). Age is estimated by counting the growth rings of horns [34]. The use of the eviscerated body weight is a good proxy for the total mass

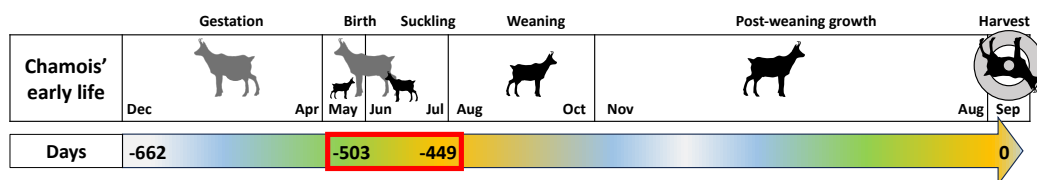


Figure 1. Timeline of the life stages of a yearling Alpine chamois from gestation to harvest. The timeline also features the time window reference days that have been used in the *climwin* analyses. Day -662 corresponds to 1 December of the year before the birth. The critical window identified by *climwin* spans from -503 to -449 between 9 May and 2 July of the birth year. Day 0 is 24 September of the harvest year, which corresponds to the end date of the 10-day hunting period.

(skeletal, muscle and fat store) of the animal, as it excludes the variation due to recently ingested food. Coordinates of the kill locations were provided by hunters at the official checkpoints when weighing their hunt, and elevation was extracted *a posteriori*. Of 34 017 animals harvested during the hunting period (age range: 0.5–22.5 years old), 5635 were yearlings (1.5 years old; 2491 females and 3144 males).

Daily mean ambient temperatures ($^{\circ}\text{C}$) from 1990 until 2018 (all years needed for analysis) were obtained from a Swiss meteorological station in the city of Lugano (273 m.a.s.l.), located within the harvesting area. Other stations at higher elevations had incomplete data but showed a high correlation with Lugano data (all Pearson $r > 0.8$; electronic supplementary material, S1), indicating that Lugano provides a good proxy for temperature in our study area.

2.2. Statistical analysis

We investigated the body mass of yearlings in relation to mean ambient temperature using the package *climwin* [35] in R v. 4.2.1 [36]. This package allows the detection of the precise time window when a biological variable is most strongly affected by an environmental variable by comparing the support of the data for competing hypotheses and formalizing them into regression models [37]. Competing models are based on a baseline model (not including weather effects) and ranked using the ΔAICc (difference in terms of the Akaike information criterion values calculated for a small sample size between the baseline model and the model of interest). The baseline model should include non-climatic variables that have been shown to explain variation in the response variable [37]. Therefore, ours was a multiple linear regression model with a body mass of yearling chamois as the response variable and elevation (continuous) and sex (two-level factor: male versus female) as the explanatory variables. The function *slidingwin* creates a set of competing models testing windows of different lengths for the weather variable of interest. We present results relative to the mean daily ambient temperature, but similar models using the minimum and maximum daily ambient temperatures provide similar time windows and comparable results (electronic supplementary material, S3). Nonlinear effects of temperature on body mass were investigated by testing for linear and quadratic trends. As parameters in *slidingwin*, we set an absolute time window with 24 September (the last date of harvest) as the reference day (figure 1). We looked for windows of all possible lengths, and start and end dates, between the reference day and 662 days prior (1 December of 2 years before) to include the critical periods in a young chamois' life: gestation, suckling, first winter and yearling. Detailed information on our analyses using *climwin* is provided in electronic supplementary material, S2.

Finally, it has been reported that the climate–phenotype relationship can be potentially spurious when both variables also change across years [38]. Stronger statistical evidence for the climate–phenology relationship can be gathered by presenting a detrended relationship [38]. Therefore, we removed the temporal trends (i.e. detrending) in our climatic and phenotypic data by extracting the residuals of linear regressions between mass and year and between temperature (during the time windows previously identified using *climwin* that most strongly affect body mass) and year. We then ran a linear model with the residuals of body mass in relation to the residuals of temperature to establish that the decrease in body mass in relation to an increase in temperature was not confounded by changes in both variables across years.

3. Results

The final model included an effect of the sex of the individual, elevation and a linear and a quadratic effect of mean daily temperature averaged between days 503 and 449 from the reference day (24

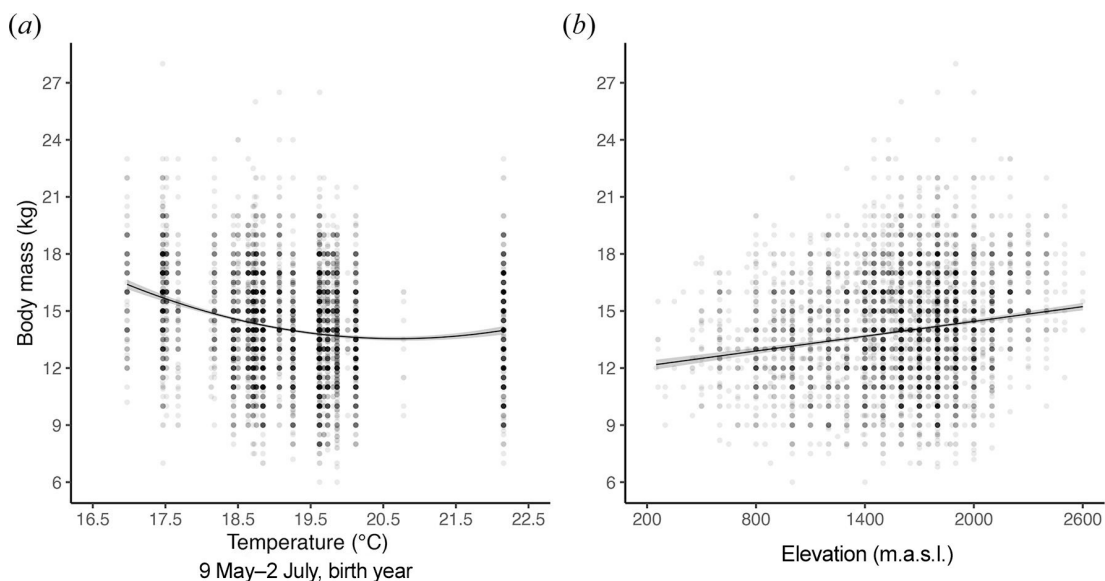


Figure 2. Relationship between body mass of harvested yearling Alpine chamois, (a) the annual average temperature between 9 May and 2 July of the birth year (suckling period) and (b) elevation (m.a.s.l.). Each dot is one observation (darker dots represent a higher number of observations); fitted lines in (a) and (b) show 95% confidence intervals (shaded areas).

Table 1. Results of the linear model showing the linear and quadratic effects of mean daily temperature (°C) averaged between 9 May and 2 July of the birth year (window: 503–449), harvest elevation (m.a.s.l.) and sex (males versus females) on body mass (kg) of harvested yearling Alpine chamois. Number of observations: 5635 in 27 years.

predictors	estimate	s.e.	t	p-value
intercept	11.867	0.153	77.78	<0.001
T (window: 503–449)	−31.948	2.581	−12.38	<0.001
T (window: 503–449) ²	14.294	3.262	4.38	<0.001
harvest elevation	0.001	0.000	14.31	<0.001
sex (M)	0.496	0.069	7.19	<0.001

September; figure 1, table 1, electronic supplementary material, S2). This climate window is equivalent to the period from 9 May until 2 July of the birth year. It is interesting to note that the body mass of yearling chamois decreases in a nonlinear fashion as the mean annual ambient temperature increases in this critical climate window, with a substantial decrease as temperatures increase up to 19.5°C followed by a plateau at higher temperatures (table 1, figure 2a). Yearling chamois mass was positively related to elevation (table 1, figure 2b), and males were heavier than females (mean \pm s.e. mass in kilograms of males: 14.2 ± 0.05 ; females: 13.6 ± 0.06 ; table 1).

Over the course of the study, the mean temperature between 9 May and 2 July increased by 0.6°C per decade ($\pm 0.2^\circ\text{C}$, $t = 2.9$, $p = 0.007$; figure 3a), leading to a 1.7°C increase in 27 years. During the same time period, the mean body mass of yearling chamois decreased by 1.12 kg per decade (± 0.06 kg, $t = -17.81$, $p < 0.001$; figure 3b), leading to an overall decrease of 2.92 kg during the study period. The analysis of the relationship between body mass and mean temperature between 9 May and 2 July after detrending the data for changes across years showed a significant nonlinear negative association between both variables (both $p < 0.001$; figure 3c, electronic supplementary material S2), thus providing strong statistical support for an effect of temperature on body mass.

4. Discussion

We found that the mass of yearling chamois is influenced by a temperature window between 2 May and 21 July during the spring and summer of their first year of life, which corresponds to the end of

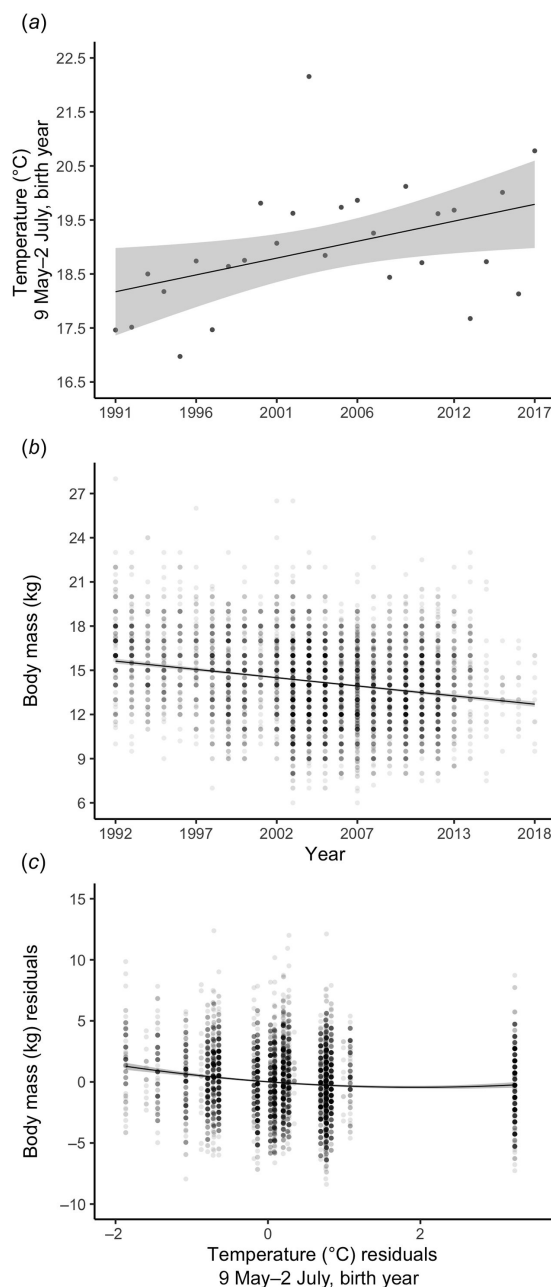


Figure 3. Annual trend of (a) mean daily temperature averaged between 9 May and 2 July of the birth year (between 1991 and 2017), (b) body mass of harvested yearling Alpine chamois between 1992 and 2018 and (c) year-detrended relationship between body mass and temperature. Detrended values in (c) are residuals from linear models in (a) and (b). Each dot is one observation (darker dots represent a higher number of observations in (b) and (c)); fitted lines show 95% confidence intervals (shaded areas).

gestation and the start of lactation. Further, we showed that the significant decrease in body mass of yearling chamois over recent decades [19] was significantly correlated with the increase in temperature between 2 May and 21 July. Our results revealed a decrease in body mass of almost 3 kg and an increase in average ambient temperature of *ca* 1.7°C between late spring and early summer during this 27-year study (1992–2018). The analysis of detrended data confirmed that this climate–phenotype relationship was not spurious and was simply explained by the changes in both variables across years. Our study adds to the previous research on the decrease in body mass in adults (data from this population and from Italy [19]) and yearlings (data from Italy [22] and Austria [23]). A recent study, however, did not find any change in body mass or size in chamois and three other ungulate species in the eastern Swiss Alps between 1991 and 2013 [24]. Several possible explanations are mentioned by the authors, including less extreme local changes in temperatures and forage quality in their population compared with other regions of the Alps.

Our results support the hypothesis that spring–summer temperatures are more important than winter temperatures with respect to phenotypic changes in seasonal environments [14,39]. Our study also identifies a specific critical window for chamois growth and development. Interestingly, we found a quadratic relationship with temperature in the critical time window that seems to indicate the presence of a temperature threshold for the growth of young chamois, with body mass being larger at increasingly low temperatures (note that the quadratic model is heuristic and does not imply that the relationship is parabolic over the whole range of temperatures). Future studies should further investigate this threshold in other populations and related taxa to better understand potential nonlinear relationships between climate and morphology.

Climate change can affect chamois growth in several ways. First, chamois births no longer coincide with the highest peak of vegetation growth due to rising temperatures altering the phenology of the plants they feed on [40]. The mismatch is linked to the fact that the annual birth peaks of herbivores are mainly influenced by day length, not by resource availability [41]. The lack of resources for mothers during the lactation period might influence how much energy can be allocated to nursing, with cascading effects on offspring growth. Second, although climate warming can increase plant productivity, Alpine plants are sensitive to heat stress and droughts, and climate warming is expected to decrease forage quality and palatability in Alpine grassland systems [8,9] for nursing mothers and their offspring. Third, ambient temperatures can also strongly influence the nutrient intake of yearling chamois during growth by altering the feeding activity [22] in young and adults. During warm days, chamois have been shown to reduce heat-generating activities (including foraging), probably in an attempt to avoid thermal stress [7]. Fourth, climate conditions may affect the body reserves of mothers, which in turn can affect offspring growth during gestation [42] and suckling. Ambient temperatures can also directly affect the quantity and quality of milk production [43], with increasing temperatures leading to lower milk yields in domesticated ungulates [44,45]. Climate change can also affect milk composition, with a significant decline in milk protein and fat content in response to warmer temperatures [46,47]. Finally, changes in body mass can also be driven by density-dependent effects if, for instance, chamois populations have increased in size during the study period, leading to more intense competition for access to grazing sites. This scenario is, however, unlikely as the populations of chamois have decreased in size in our study area [48] and in other parts of the Alps [40].

Our results support previous studies showing the importance of climatic conditions for ungulate growth at high elevations and latitudes [17,19,23]. Although ungulates can modify their behaviours by eating at night [49] or shifting their range to higher elevations [7] to cope with warming spring and summer temperatures, these changes in behaviour cannot cope with the negative effects of climatic conditions on forage quality and palatability [8,9], which persist throughout the day and can be observed at a wide range of elevation. Hence, studies on the effects of climate warming on the quantity and quality of forage during the time window most sensitive for ungulate growths (e.g. between 9 May and 2 July for chamois in our study population) could provide insightful information on the role of nutrition as an underlying mechanism linking climate warming to changes in body size [50]. Because body mass growth in the first few years of life can strongly impact an individual survival and future reproductive performance [12], with important consequences for population dynamics (and possibly explaining the decline in population size observed in this species [40,48]), more research is needed on the effects of climate change on body mass in the first few years of life rather than in adulthood. Consequently, it remains to be understood whether, for example, the decrease in body mass in response to the increase in summer temperatures may be detrimental to surviving the harsh variation in winter weather still present at high elevations. Research is also needed into the long-term fitness consequences of the changes in early growth on the whole life history of Alpine ungulates, from age at first reproduction to the onset of senescence and death, in order to adequately model the demographic consequences of climate change on these species.

Ethics. This work did not require ethical approval from a human subject or animal welfare committee.

Data accessibility. All data and code used for statistical analysis and plots are provided via the Open Science Framework and were shared with editor and reviewers at first submission [51].

Supplementary material is available online [52].

Declaration of AI use. We have not used AI-assisted technologies in creating this article.

Authors' contributions. G.M.: conceptualization, formal analysis, investigation, methodology, visualization, writing—original draft, writing—review and editing; K.G.G.: formal analysis, investigation, methodology, writing—original draft, writing—review and editing; N.I.: data curation, writing—review and editing; L.-F.B.: data curation, writing—review and editing; F.T.: data curation, writing—review and editing; P.B.: conceptualization, formal analysis, investigation, methodology, supervision, writing—review and editing.

All authors gave final approval for publication and agreed to be held accountable for the work performed therein.

Conflict of interest declaration. We declare we have no competing interests.

Funding. This project has received funding from the European Union's Horizon 2020 research and innovation programme under the Marie Skłodowska-Curie grant agreement no. 101025938 to G.M.

Acknowledgements. We thank the managers of the hunting and fishing cantonal office of Ticino, Switzerland, and the Swiss federal office of meteorology and climatology (MeteoSchweiz) for collecting the data and making them available to us.

References

1. Hetem RS, Fuller A, Maloney SK, Mitchell D. 2014 Responses of large mammals to climate change. *Temperature* **1**, 115–127. (doi:10.4161/temp.29651)
2. Gardner JL, Peters A, Kearney MR, Joseph L, Heinsohn R. 2011 Declining body size: a third universal response to warming? *Trends Ecol. Evol.* **26**, 285–291. (doi:10.1016/j.tree.2011.03.005)
3. Sheridan JA, Bickford D. 2011 Shrinking body size as an ecological response to climate change. *Nature Clim. Change* **1**, 401–406. (doi:10.1038/nclimate1259)
4. Ryding S, Klaassen M, Tattersall GJ, Gardner JL, Symonds MRE. 2021 Shape-shifting: changing animal morphologies as a response to climatic warming. *Trends Ecol. Evol.* **36**, 1036–1048. (doi:10.1016/j.tree.2021.07.006)
5. Speakman JR, Król E. 2010 Maximal heat dissipation capacity and hyperthermia risk: neglected key factors in the ecology of endotherms. *J. Anim. Ecol.* **79**, 726–746. (doi:10.1111/j.1365-2656.2010.01689.x)
6. Bergmann C. 1847 Über die Verhältnisse der Wärmeökonomie der Thiere zu ihrer Grösse. *Abgedruckt Aus Den Göttinger Studien*. **3**, 595–708.
7. Brivio F, Bertolucci C, Tettamanti F, Filli F, Apollonio M, Grignolio S. 2016 The weather dictates the rhythms: Alpine chamois activity is well adapted to ecological conditions. *Behav. Ecol. Sociobiol.* **70**, 1291–1304. (doi:10.1007/s00265-016-2137-8)
8. Mainetti A, Ravetto Enri S, Pittarello M, Lombardi G, Lonati M. 2023 Main ecological and environmental factors affecting forage yield and quality in Alpine summer pastures (NW-Italy, Gran Paradiso National Park). *Grass Forage Sci.* **78**, 254–267. (doi:10.1111/gfs.12609)
9. Dibari C et al. 2021 Climate change impacts on the Alpine, Continental and Mediterranean grassland systems of Italy: a review. *Ital. J. Agronomy* **16**, 1843. (doi:10.4081/ija.2021.1843)
10. Zamin TJ, Côté SD, Tremblay JP, Grogan P. 2017 Experimental warming alters migratory caribou forage quality. *Ecol. Appl.* **27**, 2061–2073. (doi:10.1002/eap.1590)
11. Simons MJP, Reimert I, van der Vinne V, Hambly C, Vaanholt LM, Speakman JR, Gerkema MP. 2011 Ambient temperature shapes reproductive output during pregnancy and lactation in the common vole (*Microtus arvalis*): a test of the heat dissipation limit theory. *J. Exp. Biol.* **214**, 38–49. (doi:10.1242/jeb.044230)
12. Beauplet G, Guinet C. 2007 Phenotypic determinants of individual fitness in female fur seals: larger is better. *Proc. Biol. Sci. B* **274**, 1877–1883. (doi:10.1098/rspb.2007.0454)
13. Festa-Bianchet M, Jorgenson JT, Réale D. 2000 Early development, adult mass, and reproductive success in bighorn sheep. *Behav. Ecol.* **11**, 633–639. (doi:10.1093/beheco/11.6.633)
14. Garel M, Gaillard JM, Jullien JM, Dubray D, Maillard D, Loison A. 2011 Population abundance and early spring conditions determine variation in body mass of juvenile chamois. *J. Mammal.* **92**, 1112–1117. (doi:10.1644/10-MAMM-A-056.1)
15. Hudson RJ, White RG. 1985 *Bioenergetics of wild herbivores*. Boca Raton, FL: CRC Press.
16. Gaillard JM, Festa-Bianchet M, Yoccoz NG, Loison A, Toigo C. 2000 Temporal variation in fitness components and population dynamics of large herbivores. *Annu. Rev. Ecol. Syst.* **31**, 367–393. (doi:10.1146/annurev.ecolsys.31.1.367)
17. Forchhammer MC, Clutton-Brock TH, Lindström J, Albon SD. 2001 Climate and population density induce long-term cohort variation in a northern ungulate. *J. Anim. Ecol.* **70**, 721–729. (doi:10.1046/j.0021-8790.2001.00532.x)
18. Herfindal I, Solberg EJ, Saether BE, Høgda KA, Andersen R. 2006 Environmental phenology and geographical gradients in moose body mass. *Oecologia* **150**, 213–224. (doi:10.1007/s00442-006-0519-8)
19. Rugghetti M, Festa-Bianchet M. 2012 Effects of spring-summer temperature on body mass of chamois. *J. Mammal.* **93**, 1301–1307. (doi:10.1644/11-MAMM-A-402.1)
20. Corlatti L, Lorenzini R, Lovari S. 2011 The conservation of the chamois *Rupicapra* spp. *Mamm. Rev.* **41**, 163–174. (doi:10.1111/j.1365-2907.2011.00187.x)
21. Ascenzi P et al. 1993 Functional, spectroscopic and structural properties of haemoglobin from chamois (*Rupicapra rupicapra*) and steinbock (*Capra hircus ibex*). *Biochem. J.* **296**, 361–365. (doi:10.1042/bj2960361)
22. Mason TH, Apollonio M, Chirichella R, Willis SG, Stephens PA. 2014 Environmental change and long-term body mass declines in an alpine mammal. *Front. Zool.* **11**, 69. (doi:10.1186/s12983-014-0069-6)
23. Reiner R, Zedrosser A, Zeiler H, Hackländer K, Corlatti L. 2021 Forests buffer the climate-induced decline of body mass in a mountain herbivore. *Glob. Change Biol.* **27**, 3741–3752. (doi:10.1111/gcb.15711)

24. Büntgen U, Jenny H, Galván JD, Piermattei A, Krusic PJ, Bollmann K. 2020 Stable body size of Alpine ungulates. *R. Soc. Open Sci.* **7**, 200196. (doi:10.1098/rsos.200196)
25. von Hardenberg A, Bassano B, Peracino A, Lovari S. 2000 Male Alpine Chamois Occupy Territories at Hotspots Before the Mating Season. *Ethology* **106**, 617–630. (doi:10.1046/j.1439-0310.2000.00579.x)
26. Bassano B, Perrone A, Von Hardenberg A. 2003 Body weight and horn development in Alpine chamois, *Rupicapra rupicapra* (Bovidae, Caprinae). *Mammalia* **67**, 65–74. (doi:10.1515/mamm.2003.67.1.65)
27. Garel M, Loison A, Jullien JM, Dubray D, Maillard D, Gaillard JM. 2009 Sex-specific growth in Alpine Chamois. *J. Mammal.* **90**, 954–960. (doi:10.1644/08-MAMM-A-287.1)
28. Rughetti M, Festa-Bianchet M. 2011 Effects of early horn growth on reproduction and hunting mortality in female chamois. *J. Anim. Ecol.* **80**, 438–447. (doi:10.1111/j.1365-2656.2010.01773.x)
29. Scornavacca D, Cotza A, Lovari S, Ferretti F. 2018 Suckling behaviour and allonursing in the Apennine chamois. *Ethol. Ecol. Evol.* **30**, 385–398. (doi:10.1080/03949370.2017.1423115)
30. Pettorelli N, Pelletier F, Von Hardenberg A, Festa-Bianchet M, Côté SD. 2007 Early onset of vegetation growth vs. rapid green-up: impacts on juvenile mountain ungulates. *Ecology* **88**, 381–390. (doi:10.1890/06-0875)
31. Shackleton DM. 1997 *Wild sheep and goats and their relatives: status survey and conservation action plan for Caprinae*. Gland, Switzerland: IUCN.
32. Spitzenberger F, Bauer K, Mayer A, Weis E, Preleuthner M, Sackl P, Sieber J. 2001 Die Säugetierfauna Österreichs. In *Bundesministerium für Land- und Forstwirtschaft, Umwelt und Wasserwirtschaft*. Graz Austria: Bundesministerium für Land- und Forstwirtschaft Umwelt und Wasserwirtschaft.
33. Nesti I, Posillico M, Lovari S. 2010 Ranging behaviour and habitat selection of Alpine chamois. *Ethol. Ecol. Evol.* **22**, 215–231. (doi:10.1080/03949370.2010.502316)
34. Schroder W, Elsner-Schack IV. 1985 Correct age determination in chamois. In *The biology and management of mountain ungulates* (ed. S Lovari), pp. 65–70, London, UK: Croom Helm.
35. Bailey LD, van de Pol M. 2016 climwin: An R Toolbox for Climate Window Analysis. *PLoS One* **11**, e0167980. (doi:10.1371/journal.pone.0167980)
36. R Core Team. 2023 *R: A language and environment for statistical computing*. Vienna, Austria: R Foundation for Statistical Computing.
37. van de Pol M, Bailey LD, McLean N, Rijdsdijk L, Lawson CR, Brouwer L. 2016 Identifying the best climatic predictors in ecology and evolution. *Methods Ecol. Evol.* **7**, 1246–1257. (doi:10.1111/2041-210X.12590)
38. Iler AM, Inouye DW, Schmidt NM, Høye TT. 2017 Detrending phenological time series improves climate-phenology analyses and reveals evidence of plasticity. *Ecology* **98**, 647–655. (doi:10.1002/ecy.1690)
39. Klein DR. 1965 Ecology of deer range in Alaska. *Ecol. Monogr.* **35**, 259–284. (doi:10.2307/1942139)
40. Chirichella R, Stephens PA, Mason THE, Apollonio M. 2021 Contrasting effects of climate change on Alpine chamois. *J. Wildl. Manag.* **85**, 109–120. (doi:10.1002/jwmg.21962)
41. Post E, Forchhammer MC. 2008 Climate change reduces reproductive success of an Arctic herbivore through trophic mismatch. *Philos. Trans. R. Soc. Lond. B. Biol. Sci.* **363**, 2369–2375. (doi:10.1098/rstb.2007.2207)
42. Hansen PJ. 2009 Effects of heat stress on mammalian reproduction. *Philos. Trans. R. Soc. Lond. B. Biol. Sci.* **364**, 3341–3350. (doi:10.1098/rstb.2009.0131)
43. Liu J, Li L, Chen X, Lu Y, Wang D. 2019 Effects of heat stress on body temperature, milk production, and reproduction in dairy cows: a novel idea for monitoring and evaluation of heat stress — a review. *Asian-Australas J. Anim. Sci.* **32**, 1332–1339. (doi:10.5713/ajas.18.0743)
44. Upadhyay RC, Singh SV, Kumar A, Gupta SK. 2007 Impact of climate change on milk production of Murrah buffaloes. *Ital. J. Anim. Sci.* **6**, 1329–1332. (doi:10.4081/ijas.2007.s2.1329)
45. Gorniak T, Meyer U, Südekum KH, Dänicke S. 2014 Impact of mild heat stress on dry matter intake, milk yield and milk composition in mid-lactation Holstein dairy cows in a temperate climate. *Arch. Anim. Nutr.* **68**, 358–369. (doi:10.1080/1745039X.2014.950451)
46. Knapp DM, Grummer RR. 1991 Response of lactating dairy cows to fat supplementation during heat stress. *J. Dairy Sci.* **74**, 2573–2579. (doi:10.3168/jds.S0022-0302(91)78435-X)
47. Gantner V, Mijić P, Kuterovac K, Solić D, Gantner R. 2011 Temperature-humidity index values and their significance on the daily production of dairy cattle. *Mljekarstvo: Časopis Za Unaprjeđenje Proizvodnje i Prerade Mlijeka*. **61**, 56–63.
48. Tettamanti F. 2016 Ricostruzione statistica della popolazione di camoscio nel Canton Ticino. Evoluzione durante il periodo 1991–2014. Ufficio della caccia e della pesca, Canton Ticino
49. Grignolio S, Brivio F, Apollonio M, Frigato E, Tettamanti F, Filli F, Bertolucci C. 2018 Is nocturnal activity compensatory in chamois? A study of activity in a cathemeral ungulate. *Mamm. Biol.* **93**, 173–181. (doi:10.1016/j.mambio.2018.06.003)
50. Rosenblatt AE, Schmitz OJ. 2016 Climate change, nutrition, and bottom-up and top-down food web processes. *Trends Ecol. Evol.* **31**, 965–975. (doi:10.1016/j.tree.2016.09.009)
51. Masoero G, Gencheva KG, Ioset N, Bersier LF, Tettamanti F, Bize P. 2024 Data from: Body size changes in yearling Chamois. OSF. (doi:10.17605/OSF.IO/P9C4M)
52. Masoero G, Gencheva KG, Ioset N, Bersier LF, Tettamanti F, Bize P. 2024 Data from: shrinking Alpine chamois: higher spring temperatures over the last 27 years in Switzerland are linked to a 3 kg reduction in body mass of Yearlings. Figshare. (doi:10.6084/m9.figshare.c.7095864)