

The impact of large terrestrial carnivores on Pleistocene ecosystems

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Large mammalian terrestrial herbivores, such as elephants, have dramatic effects on the ecosystems they inhabit and at high population densities their environmental impacts can be devastating. Pleistocene terrestrial ecosystems included a much greater diversity of megaherbivores (e.g., mammoths, mastodons, giant ground sloths) and thus a greater potential for widespread habitat degradation if population sizes were not limited. Nevertheless, based on modern observations, it is generally believed that populations of megaherbivores (>800 kg) are largely immune to the effects of predation and this perception has been extended into the Pleistocene. However, as shown here, the species richness of big carnivores was greater in the Pleistocene and many of them were significantly larger than their modern counterparts. Fossil evidence suggests that interspecific competition among carnivores was relatively intense and reveals that some individuals specialized in consuming megaherbivores. To estimate the potential impact of Pleistocene large carnivores, we use both historic and modern data on predator–prey body mass relationships to predict size ranges of their typical and maximum prey when hunting as individuals and in groups. These prey size ranges are then compared with estimates of juvenile and subadult proboscidean body sizes derived from extant elephant growth data. Young proboscideans at their most vulnerable age fall within the predicted prey size ranges of many of the Pleistocene carnivores. Predation on juveniles can have a greater impact on megaherbivores because of their long interbirth intervals, and consequently, we argue that Pleistocene carnivores had the capacity to, and likely did, limit megaherbivore population sizes.

Keywords: predator ; megafauna ; hypercarnivore ; Carnivora ; megaherbivore

Large mammalian terrestrial herbivores have dramatic effects on the ecosystems they inhabit and at their highest population densities the environmental impact of the largest modern herbivores, such as elephants, can be devastating (1, 2). At the end of the Pleistocene, loss of large mammalian herbivores has been implicated in major shifts in vegetation structure and ecosystem processes, including transitions between biomes, such as from grass-dominated steppe biomes to moss-dominated tundra (3, 4). What prevented widespread habitat destruction during the Pleistocene, when ecosystems sustained multiple species of megaherbivores (>800 kg), including mammoths, mastodons, and giant ground sloths?

Although the role of predators in structuring terrestrial ecosystems is widely acknowledged (5, 6), it is also widely assumed that the large body size of megaherbivores renders their populations exempt from “top-down” limitation by predators both now and in the past (7–10). However, the impact of large terrestrial predators on Pleistocene ecosystems may be difficult to appreciate, because these carnivores interacted within much more species-rich guilds than exist today. In addition, these Pleistocene guilds included extinct species (such as sabertooth cats and very large hyenas) for which we have no close living

analogs, making their prey preferences a matter of inference, rather than observation.

In this article, we estimate the predatory impact of large (>21 kg, ref. 11) Pleistocene carnivores using a variety of data from the fossil record, including species richness within guilds, population density inferences based on tooth wear, and dietary inferences based on stable isotope ratios as well as carnivore-produced bone accumulations. In addition, we use both historical and recent data on the relationships between masses of extant predators and prey to estimate the prey size preferences of prehistoric carnivores and compare these to the estimated sizes of their potential prey, specifically, juvenile and young adult mammoths and mastodons. On the basis of these data we suggest that Pleistocene carnivores had the capacity to, and likely did, limit megaherbivore population sizes through predation on juvenile and subadult individuals.

Guilds of Large Carnivores Then and Now

The importance of carnivores in shaping Pleistocene terrestrial ecosystems is readily underestimated because carnivore species diversity and body size are much reduced in modern communities. Using several localities with well-preserved fossils representing both the early (1–1.5 million years B.P.) and late (<500,000 y B.P.) Pleistocene of the Old and the New World, we compared the diversity in species size and richness among fossil and contemporary carnivore guilds (Fig. 1 and *SI Appendix*, Table S1). In general, Pleistocene guilds tended to have more

Significance

At very high densities, populations of the largest herbivores, such as elephants, have devastating effects on the environment. What prevented widespread habitat destruction in the Pleistocene, when the ecosystem sustained many species of huge herbivores? We use data on predator–prey body mass relationships to predict the prey size ranges of large extinct mammalian carnivores, which were more diverse and much larger than living species. We then compare these prey size ranges with estimates of young mammoth sizes and show that juvenile mammoths and mastodons were within predicted prey size ranges of many of the Pleistocene carnivores. From this and other fossil evidence we argue that, by limiting population sizes of megaherbivores, large carnivores had a major impact on Pleistocene ecosystems.

Author contributions: B.V.V., M.W.H., W.J.R., C.M., and V.L.R. designed research, performed research, analyzed data, and wrote the paper.

The authors declare no conflict of interest.

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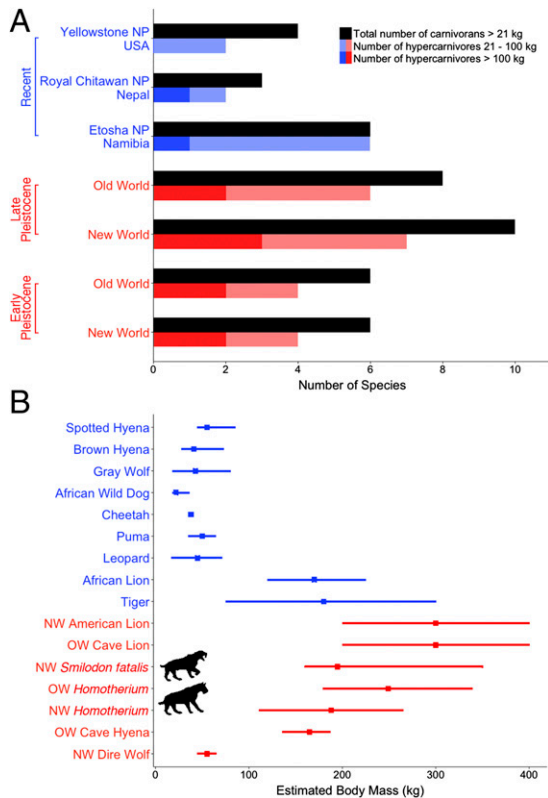


Fig. 1. (A) Predator guild composition for four Pleistocene (red) and three extant (blue) communities. Indicated for each guild are the total number of species of carnivores (hypercarnivores and omnivores, e.g., ursids) with masses >21 kg (black), the subset of these that are hypercarnivores (two-toned bar), and the subset of these that are hypercarnivores with masses >100 kg (dark blue or red). (B) Estimated body masses (mean and range) of extant (blue) and extinct (red) hypercarnivores. Silhouettes are provided only for the sabertooth cats because they lack modern analogs. For details on the localities and species compositions, as well as body mass estimations, see *SI Appendix, SI Materials and Methods*, section 1.

species with masses greater than 21 kg, and these species tended to be larger than equivalent extant species (Fig. 1). Averaging across the species found within single communities, the mean size of large hypercarnivores (species whose diets consist of >80% meat) in the extant guilds ranges from 53 to 63 kg, whereas it spanned 96–135 kg in the fossil guilds. Although guilds in the most diverse modern African communities are similar to those in the late Pleistocene in containing five to six large hypercarnivores, they include only one hypercarnivore that exceeds 100 kg, the lion (*Panthera leo*). At present (and excluding polar bears, who feed on marine resources), there are only two hypercarnivores that exceed 100 kg in mass, the aforementioned lion and the tiger (*Panthera tigris*), and these are not found in sympatry. In the late Pleistocene, there were four to five more large hypercarnivores and it was typical to find two to three in sympatry (Fig. 1). For example, there were massive sabertooth felids (*Smilodon* sp. and *Homotherium* sp.) in addition to much larger relatives of the extant lion (*Panthera leo spelaea* and *Panthera atrox*) in both the Old and the New World, as well as huge spotted hyenas (*Crocuta crocuta spelaea*) in the Old World and a relatively carnivorous, enormous bear (*Arctodus simus*) in the New World. These Pleistocene giants were at least a third to more than twice the mass of their extant relatives (Fig. 1). Moreover, as noted above, some of these species were sabertooth cats, an ecomorph without a close modern analog. These imposing felids possessed a suite of adaptations that enhanced their ability

to kill large prey, including enlarged knife-like upper canines, a long, thick neck, and robust, heavily muscled forelimbs (12). Notably, nearly all Pleistocene predator guilds found outside of Australia included at least one and often two species of large sabertooth cat.

What could have supported such a high diversity of coexisting, large predators? Among most extant communities, large predator species richness is more closely linked to prey richness than either primary productivity or climate and this relationship is more pronounced within sets of similar sized predators and prey (13). To further examine this relationship, we surveyed 181 present day mammalian faunas (*SI Appendix, SI Materials and Methods*, section 2, and Table S6) that include at least one species each of large hypercarnivore and megaherbivore (species >800 kg), and we found that, as the number of megaherbivore species increases, so does the likelihood of finding three or more coexisting hypercarnivores. For example, of the 28 faunas that include one megaherbivore, only 9 (32%) include three or more large hypercarnivores. By contrast, this percentage climbs to 52% (11/21) when two megaherbivores are present, and further still to 91% (31/34) when three or more megaherbivores are present. The maximum richness of six coexisting large hypercarnivores is found only in communities with three or more megaherbivores. This association between hypercarnivore diversity and megaherbivore diversity suggests that the presence of huge herbivores promotes, or at least permits, coexistence among big predators. The correlation could arise from a variety of causes. Given the size of their carcasses, megaherbivores could be a significant food resource for scavenging and hunting predators (14). Megaherbivores themselves may modify the environment in ways that increase hunting success by creating more edge habitats that favor ambush predators such as lions, or by shifting woodland and forest toward grassland, thus improving the habitat for cursorial hunters such as African wild dogs and spotted hyenas (9, 15). A dynamic cycle could arise where vegetation patterns shift with the relative dominance of megaherbivores or large predators. However, it is unclear to what extent such habitat modification would occur if the large carnivores exerted strong top-down pressure on the megaherbivores.

In contrast to the present, all four of the Pleistocene fossil communities we examined (*SI Appendix, Table S1*) had two to six megaherbivores and four to seven large hypercarnivores, two to three of which exceeded 100 kg in mass. In the Old World, the megaherbivores usually included mammoth (*Mammuthus* sp.), rhinoceros (*Stephanorhinus* sp. or *Coelodonta* sp.), and a giant bovid (*Praeovibos* sp., *Bison* sp., *Symbos* sp., or *Bos* p.) (16, 17). In North America, the species richness of megaherbivores was even greater, in part due to the immigration of giant ground sloths from South America at least 2.5 million years ago (18). At the late Pleistocene site of Rancho La Brea, California, there were six megaherbivores: two proboscideans (*Mammuthus columbi* and *Mammot americanum*), giant camel (*Camelops hesternus*), extinct bison (*Bison antiquus*), and two ground sloths (*Megalonyx jeffersoni* and *Paramylodon harlani*) (19). Not surprisingly, this site also exhibits the greatest richness of hypercarnivores >21 kg ($n = 7$) across all our fossil guilds.

Prey body size tends to increase with predator size (10, 20). Given the greater prevalence of very large (>100 kg) hypercarnivores in association with multiple megaherbivores in Pleistocene communities, it seems likely that predation pressure on megaherbivores was greater in the past than in modern mammalian communities. However, this idea is not so easily accepted, given the observation that even the largest of living carnivores, tigers and lions, rarely tackle adult elephants, hippos, or rhinos. But what about juveniles?

Predation on Extant Megaherbivores

Modern day megaherbivores may appear largely immune to the effects of predation because their bodies are massive and maternal protection of juveniles is strong (9, 10). Juvenile elephants do not stray far from their mother's side until they are about 5–7 years of age (*SI Appendix, SI Materials and Methods*, section 6). Nevertheless, our review of the literature reveals that young African elephants are taken regularly, especially at the end of the dry season when they may be more susceptible (*SI Appendix, Table S2*). For example, 74 elephants were killed by lions over a 4-y period in Chobe National Park, Botswana, with nearly two-thirds of the kills on juveniles and subadults 9 y old or younger (21). Elsewhere in Africa, 44 kills of elephants by lions were observed in Zimbabwe over a 6-y span, with juveniles less than 8 years of age being targeted (22), and smaller numbers of kills have been documented in both the Central African Republic (23) and Kenya (24). In one study, elephant made up 20–23% of the total biomass consumed by lions annually, and exceeded the biomass contributions made by all other prey except buffalo (25). Young rhinoceros are also not immune to predation by lions as evidenced by three kills of subadults that were made over 3 mo in Etosha National Park, Namibia (26). Goddard (27) estimated that 16% of black rhinos younger than 2 y old were killed by lions and spotted hyenas in East Tsavo Park, Kenya. Spotted hyenas were also observed to kill five young elephants in Hwange National Park, Zimbabwe in a single year (28). Thus, it is clear that lions, and to a lesser extent, spotted hyenas, are fully capable of killing juvenile and subadult megaherbivores that can weigh as much as 1,500 kg.

Most often, the killing of megaherbivores is accomplished by a group of individuals working together. Hunting in groups facilitates the killing of large prey and accordingly, the success rate for lions taking elephants appears to be enhanced by large pride size. In Botswana, lions were observed to regularly use a strategy in which one to two lions leapt onto and bit the back of the victim while others on the ground worked to sever the relatively thin flexor muscles of the hindlimb, resulting in rapid immobilization (29). Of 18 such attempts on elephants, 4 were successful, all of which involved more than 27 of the 30 lions in the pride. Similarly, the pride that took more than 70 elephants in 3 years in Botswana was also large, consisting of 18 individuals (21). These numbers suggest that large prides are predisposed to attack large megaherbivores. If so, predation on elephants by lions may be less frequent now than in the past because of declines in pride size due to human persecution and reductions in prey numbers (29, 30). With the advent of large-scale human hunting, larger prides would have provided more conspicuous targets. The decline in lion numbers in Africa over the past 100 y is well known; whereas there were perhaps 500,000 lions on the continent in 1950, there are now fewer than 30,000 (31). Whether or how closely pride size should follow population size is unclear, but in Etosha National Park, a reduction in the mean number of adults per pride from 10 to 6 accompanied a 33% population decline over 12 y (30). Our review of African fauna historical records in the period 1835–1950 suggests that modern perceptions of typical lion pride size may reflect only what has been observed over the past 60 y (*SI Appendix, Table S3*). A published survey of pride size across 27 African reserves between 1997 and 2007 found a mean of 9 (± 4) adults (32), but older records include multiple reports of prides of 35–40 individuals, and in some instances such sightings were not unusual. For example, Sikes (ref. 33, p. 253) commented on predator group sizes in the period between 1901 and 1931, writing “In the days when such well-known personalities as Lord Delamere habitually encountered prides of up to 36 lions on their ranches in Kenya, when hunters all too frequently stumbled on prides of between 17 and 40 on a kill or when the lone horseman found himself surrounded by packs

of between 25 and 40 Cape hunting dogs, these animals were sufficiently numerous to keep elephant calf mortality at a high level.” If, as the historical record suggests, African lion prides were significantly larger in the past, predation on elephants may have been a more regular occurrence than is observed today.

Pleistocene Predator Group Size and Population Density

Typical group sizes for Pleistocene predators are difficult, if not impossible, to determine, but the behavior of extant predators suggests that conditions in the Pleistocene would have favored sociality and the existence of large prides, clans, and packs. The greatest diversity of social predator species today is found in African savannah woodlands, where lions, spotted hyenas, African wild dogs, and to a lesser extent cheetahs (*Acinonyx jubatus*) all hunt in groups. Beyond expanding the size of prey that they can kill, working in groups allows all but the cheetah to better defend their kills against carcass theft (kleptoparasitism). Moreover, in aggressive interactions over carcasses between lions and hyenas, or wild dogs and hyenas, group size is an important determinant of who wins (34, 35).

Given the multiple anthropogenic forces that currently limit large carnivore abundance, such as habitat loss, competition for prey, and direct persecution, it seems likely that predator densities and group sizes could have been much greater in the Pleistocene than in even the recent past (5). If so, attempts at kleptoparasitism were probably a common occurrence, and this behavior also would favor large group sizes (36, 37). Larger groups in turn would favor more complete consumption of carcasses, including bones. Among modern carnivores, more bone is eaten when prey are more difficult to acquire, and this activity increases both the number of teeth broken in life and rates of tooth wear (38). Elevated tooth fracture frequencies observed among multiple species of Pleistocene carnivores suggest ecosystems in which the densities of predators relative to prey were high, and thus competition for carcasses was intense, leading to more frequent intra- and interspecific confrontations over kills (39). In large New World predators of the Pleistocene, for example, rates of tooth fracture are as much as three to five times that of their modern counterparts (38). Very high rates of tooth fracture are also present in Pleistocene gray wolves from Great Britain dated between 50 and 85 thousand years ago (40) and late Pleistocene cave lions and cave hyenas from Zoolithen Cave, Germany (our data, *SI Appendix, Table S4*).

Estimating Pleistocene Predator–Prey Preferences

The prey preferences of extant large carnivores have recently been reviewed in a series of papers by Hayward and colleagues (*SI Appendix, SI Materials and Methods*, section 3). These data are used here to construct regression equations of accessible (typical) and largest prey body mass against predator body mass for extant species hunting alone or in groups, and the equations are extrapolated to predict the prey sizes of extinct species (Fig. 2 and *SI Appendix, Table S5* and Fig. S1). Based on their morphology and their extant relatives, several of the large Pleistocene hypercarnivores, such as the dire wolf (*Canis dirus*), gray wolf (*Canis lupus*), and cave hyena (*C. c. spelaea*) were probably social because all are large, cursorial predators that are unable to grapple with their prey, and instead must subdue prey with their jaws alone. When it is difficult for a solitary individual to kill prey much larger than itself, hunting in groups is favored. Felids are not so constrained by their anatomy and single individuals can kill relatively large prey. Nevertheless, hunting in groups does extend the size range of prey that can be killed and may increase hunting success on very large prey (see below), so it is possible that some or all of the large Pleistocene felids (*Homotherium*, *Smilodon*, *P. atrox*, *P. leo spelaea*) were social at times. Arguments have been made in favor of sociality in each of these (12, 41) but

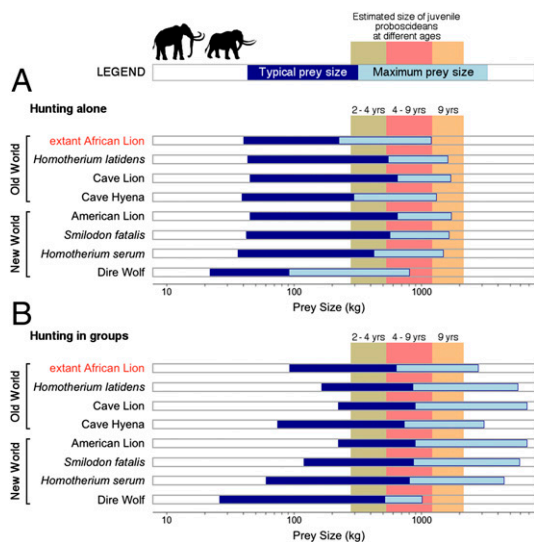


Fig. 2. Predicted typical (dark blue) and maximum (light blue) prey size ranges (horizontal bars) for the extant African lion (*Panthera leo*) and large extinct Pleistocene predators superimposed on the estimated sizes of juvenile proboscideans (mammoth and mastodons) at different ages (vertical stripes). Prey size ranges are estimated both for (A) solitary hunting and (B) group hunting. Size estimates for mammoths are based on data from living elephants. Prey size ranges for the predators were based on known relationships between prey size and predator body mass for extant large carnivores. For details, see *SI Appendix, SI Materials and Methods*, sections 3 and 6.

some workers have disagreed (42). Because it is difficult to be confident of the social behavior of extinct species, we estimated prey sizes for all species as both solitary and group hunters. These prey sizes are compared with our estimates of body sizes of proboscideans (the largest of the megaherbivores) in the most vulnerable age classes (*SI Appendix, SI Materials and Methods*, section 6).

With the exception of the dire wolf, estimated typical prey size ranges of the Pleistocene species exceed that of extant African lions (Fig. 2). Whereas we infer that the typical range of prey for a solitary African lion would not include 2- to 4-y-old mammoths, we predict that all four of the Pleistocene felids could have included them as typical prey, even without considering the special weaponry of the sabertooths. Our predictions of the maximum prey sizes for the fossil cats also exceed that shown for the extant lion, with each species, we infer, being capable of killing 9-y-old subadult proboscideans. Hunting in groups increases the upper range of available prey sizes, and the difference between the extant lion and Pleistocene species is most apparent in terms of the predicted maximum size of prey. In groups, the extinct cats are estimated to have been able to kill adults with masses between 5,700 kg (*Homotherium* spp.) and 6,700 kg (*P. atrox* and *P. l. spelaea*), thus encompassing the size of female adult and male young adult proboscideans (Fig. 2 and *SI Appendix, Fig. S 2*). These data suggest that juvenile proboscideans, rhinos, and ground sloths would all have been well within the realm of possibility for many of these extinct hypercarnivores. Adult megaherbivores appear to have been outside the typical prey size range of Pleistocene hypercarnivores, but would have been accessible to most species hunting in groups. This situation suggests that, if predators did limit their populations, it would have been mainly through predation on younger individuals. In Africa today, predation on elephants by nonhuman predators is observed, but not top-down regulation, in large part because maternal defense of juveniles appears to greatly inhibit successful attacks by modern predators that hunt in groups that are comparatively small (see above).

The fossil record provides some limited data indicating that Pleistocene carnivores did in fact consume megaherbivores. Studies of carbon and nitrogen stable isotope ratios in a number of large species suggest that most individuals were generalists that consumed a mix of large ungulates but some individuals of gray wolves and cave hyenas specialized on mammoth (*SI Appendix, SI Materials and Methods*, section 7). In addition to the stable isotope data, there is evidence of a preference for megaherbivores from several fossilized den sites of both cave hyenas (43, 44) and the sabertooth cat, *Homotherium serum* (45). These den sites include numerous tooth-marked bones of juvenile woolly mammoth and rhinoceros, in the case of the cave hyenas, and juvenile Columbian mammoth, in the case of the sabertooth cat. The predominance of juvenile prey suggests that most or all of these individuals were killed rather than scavenged (44, 45).

Could Pleistocene Carnivores Limit Megaherbivore Populations?

Others before us have viewed the large hypercarnivores of the Pleistocene as capable of killing megaherbivores (e.g., refs. 29, 46), but few have addressed the issue of whether this capability resulted in limiting megaherbivore populations (39). However, when the issue has been explicitly addressed, the prevailing opinion seems to be similar to that of Owen-Smith (9) who wrote, “prior to human arrival, populations of mammoths, mastodon, and ground sloth would have existed at saturation levels where further increase was prevented by food limitation,” or Sinclair et al. (10), who said, “A threshold occurs at prey body sizes of 150 kg, above which ungulate species have few natural predators and exhibit food limitation,”—statements based on observations of living systems. The conclusion that megaherbivores were immune to the effects of predation seems improbable given the greater size of the Pleistocene hypercarnivores. As we describe below, selective predation on juveniles would have intensified the effects of these carnivores on prey populations, given that species with the low reproductive rates typical of megaherbivores are susceptible to population reduction under conditions of relatively low predation pressure (47). Among extant large predators, the proportion of prey that are juveniles increases with prey size; living spotted hyenas tend to take juveniles of smaller- and medium-sized ungulates in proportion to their abundance, but shift to taking mostly or all juveniles of very large prey such as giraffes, black rhinos, and elephants (17). Clearly, this change in preference is a consequence both of their body sizes and of the much greater challenge of killing adults.

In the extreme, it is possible for relatively low rates of selective predation on juveniles to lead to extinction (47). As shown in an analysis of extinction patterns among nine groups of mammals (185 species) that lost three or more species at or near the end of the Pleistocene (47), species with reproductive rates of less than one offspring per female per year were much more likely to have gone extinct than those with faster reproductive rates. Some species of extinct megafauna, such as mammoths, mastodons, and ground sloths were not included in that analysis, but it is almost certain that all these species had very low reproductive rates. Using the PanTHERIA database for extant mammals (esapubs.org/archive/ecol/e090/184/metadata.htm), we find that 22 of the 29 ungulate species with masses greater than 200 kg have interbirth intervals that exceed 1 y, and of these, 9 of the 12 species with masses greater than 600 kg have interbirth intervals that exceed 2 y (*SI Appendix, Fig. S 4*). Indeed, for woolly mammoths, weaning age has been estimated from stable isotope analysis and tooth wear to have been at least 1.5 y and in one case more than 5 y (*SI Appendix, SI Materials and Methods, section 6*), and patterns of tusk growth in female mastodons point to typical calving intervals of 3–4 y (e.g., ref. 48). Although there are small mammals with relatively slow reproductive rates, such as echidnas, there are no very large mammals with relatively high reproductive rates (47);

thus it is safe to assume that extinct megaherbivores had interbirth intervals that exceeded 1 y, and could have been 4 y or more, as is typical of African elephants (*SI Appendix, SI Materials and Methods*, section 6 for more details). Given such low fecundity, the number of deaths due to predation and other causes, such as ep-isodic droughts or wild fires, would not have had to be high to keep mortality rates above recruitment rates and thereby limit population growth.

Providing further confirmation, a recent study used a discrete, stochastic model (49) analogous to a life-table analysis to examine the long-term impact of age-specific mortality on African elephant population growth (50). Drawing values of life-history parameters from multiple studies of wild elephant populations, the authors estimated the age-specific mortality needed to achieve 0% population growth for simulations spanning hundreds of years. Model projections showed that annual mortality of just 17% of juveniles aged 0–9 y would be enough to halt population growth; by comparison, the death annually of 10.5% of all adults, aged 10–60, would be needed to yield the same effect (50). We find no data on percentages of juvenile elephants taken by extant lions in Africa, but in other large mammal predator–prey systems, annual percentages of juveniles killed easily exceed 17%. For example, in Wood Buffalo National Park, Canada, gray wolves regularly prey on bison, a species at the upper limit of their capabilities, and are documented to kill more than a third of all juveniles each year (51). Similarly, bears in Yellowstone National Park remove 40% of the elk calves annually (52). Like their modern counterparts, Pleistocene carnivores probably preyed preferentially on juvenile rather than adult megaherbivores, all of which fell within their predicted range of typical prey size.

Theoretical evidence has supported the idea that populations of medium to large sized herbivores were limited by large Pleistocene hypercarnivores (53), but it has been less clear whether the theory applies to megaherbivores. Based on studies of modern African elephants, it is not clear that they can effectively self-regulate and maintain their own numbers at levels that allow for a sustainable existence of a healthy population. Gough and Kerley (54), for example found no evidence for density-dependent regulation in a South African elephant population they studied between 1976–1979 and 1996–2006. Birth rates and overall population growth rate did not slow as elephant density rose, despite serious declines in plant biomass and biodiversity. Of course, it might be argued that human activities have limited elephants to reserves that are too small, and in the past, they would have moved from areas of low forage quality to areas of better quality as needed. However, it seems likely that before the expansion of modern and especially industrialized humans any appropriate habitats would have been fully occupied by elephants, thus limiting their ability to expand their foraging range. Africa was reportedly home to five species of elephants during the Pleistocene, with two or more inhabiting some regions (55). It is hard to imagine how they partitioned their shared resources, but it certainly suggests a crowded system, in which top-down forcing was probably essential to ensure long-term stability.

We suggest that large hypercarnivores must have limited at least the proboscideans, especially given the impressive impact these species have on vegetation structure and quality. Of course, predation would not have been the sole factor; periodic droughts can produce substantial mortality in modern elephant and rhinoceros populations, especially among juveniles and subadults (56). It is noteworthy that Pleistocene large mammal community composition is remarkably stable at a continental scale over at least the last 1 million years in both the Old and the New Worlds, despite glacial–interglacial fluctuations in climate (57–59). The apparently long-term and persistent stability suggests the existence of rich and complex communities that included multiple species at different trophic levels playing similar roles

(redundancies), thus enhancing their resilience in the face of environmental perturbations. Environmental reconstructions of late Pleistocene interglacial environments in the United Kingdom, for example, reveal an abundant, diverse large herbivore guild associated with a mosaic of vegetation structures that promoted biodiversity (60). In extant large mammal communities that lack big apex predators, large herbivores often experience rapid population expansions. For example, in Eurasia and North America, cervid densities were on average nearly six times greater in areas without wolves compared with areas with wolves (61). These impressive herbivore irruptions can have very negative impacts on vegetation and ecosystem services and can produce declines in floral and faunal biodiversity if they are persistent or occur repeatedly (62, 63). If megaherbivores had not been predator limited, the Pleistocene might be expected to have experienced a long-term decline in ecosystem stability but there is no evidence of such a gradual decline. Instead, megafaunal extinctions are concentrated close to the Pleistocene–Holocene transition, associated with the presence of humans (64), and potentially linked to the effects of human hunting/scavenging in addition to ongoing predation by large carnivores (39). The negative impact of human hunting on megaherbivore numbers could have been especially large if the prey species were already under pressure due to top-down forcing by large carnivores (39) and episodic environmental stressors, such as severe drought and wild fires.

Implications for the Future

Why should we care about the role of extinct predators in their ecosystems? What bearing does it have on current struggles to preserve biodiversity? One answer is that many of the species we are most concerned about preserving evolved during or before the Pleistocene, and thus did so under very different conditions from the present. As a result, aspects of their behavior and morphology may be better explained as a response to ancient rather than current selection pressures. Secondly, studies of the Pleistocene reveal that the planet was capable of sustaining many more species-rich communities that included a greater proportion of megafauna than are found today (65). It appears that the complexity of these communities and their trophic depth, especially the presence of large apex predators, contributed to their stability, and the same would apply to the many, more ancient communities that included megaherbivores before the Pleistocene. Recreating these communities is not possible, but their record of success compels us to maintain the diversity we have and rebuild it where feasible (e.g., rewilding). Then as now, it is likely that large predators influenced their communities via processes that favored biodiversity by creating increased scavenging opportunities, refuges from herbivory for plants, and enhanced environmental heterogeneity and stability (5, 6, 66, 67). The late Pleistocene extinction of the largest of the hypercarnivores almost certainly resulted from the disappearance of their preferred prey, including large equids, bovids, and we argue, young megaherbivores. It is probably not a coincidence that spotted hyenas and lions have persisted in Africa alongside megaherbivores, while disappearing from more northern latitudes. With a growing awareness of the prevalence of top-down forcing, we are just beginning to understand the ecological and evolutionary linkages among these large mammals, and studies of their interactions on deeper timescales are an important piece of the puzzle.

ACKNOWLEDGMENTS. We thank the organizers of the Megafauna and Ecosystem Function conference for inviting this paper, and thank C. Badgley, J. Damuth, C. Sandom, P. Shipman, the Behavioral Ecology and Morphology discussion group at Duke University, students in the B.V.V. laboratory, and two anonymous reviewers for their comments. M. Balisi and C. Brown helped with illustrations and the National Science Foundation provided funding from EAR 1237928 (to B.V.V.).

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SUPPORTING Information

The Impact of Large Terrestrial Carnivores on Pleistocene Ecosystems

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SI Materials and Methods (subsections numbered and indicated with §)

§ 1) Extant and extinct guilds of large carnivores with body masses >21 kg (Table S1).

The species composition, body masses and diets for four Pleistocene guilds and three extant guilds are listed in Table S1. The sources for the species lists are as follows: Scladina Cave (1); Venta Micena (2); Rancho La Brea, <http://www.nhm.org/site/research-collections/rancho-la-brea/rlb-mammals-list> ;Leisey Shell Pit, <http://www.flmnh.ufl.edu/vertpaleo/Leiseyshellpit1A.htm>; Yellowstone (3),; Chitawan NP, <http://www.chitwannationalpark.gov.np/index.php/biodiversity>; Etosha NP (4). Body masses for the extinct species were estimated using previously published regressions of various skeletal or dental measures against body mass in extant carnivores as noted in Table S1.

§ 2) Analysis of the diversity of megaherbivores and hypercarnivores >21 kg in modern mammal communities (Table S6).

We extracted data on megaherbivore and large hypercarnivore diversity for all mammal communities (outside of Australia and Sri Lanka) that included at least one large hypercarnivore and one megaherbivore (n=181) from the Absolut Extant Database that is being compiled by J. Damuth and C. Badgley. This database is a compilation of published literature records of over 300 mammalian local or small-regional faunas, worldwide in scope, begun originally as a project at the National Center for Ecological Analysis and Synthesis in Santa Barbara, California. The data are currently being prepared for publication. Because the database is not yet publicly available, the data we used are presented here (Table S6).

§ 3) Prediction of prey size ranges for extinct carnivores (Table S5, Figure S1)

We used global reviews (5-14) of the diets of large, extant predators to derive equations of dietary parameters to predict those of extinct species. The large extant species were separated into predominant hunting strategy as follows: solitary cheetah *Acinonyx jubatus* [n = 3909 kill records]; leopard *Panthera pardus* [n = 8643]; snow leopard *P. uncia* [n = 1696], and tiger *P. tigris* [n = 3187] and group hunters (African wild dog *Lycaon pictus* [n = 4878]; dhole *Cuon alpinus* [n = 8816]; gray wolf *Canis lupus* [n = 13,348]; lion *Panthera leo* [n = 22,684]; and spotted hyena *Crocuta crocuta* [n = 3478]) (references below). These reviews provided data on the accessible prey weight range (i.e., those species most likely to be eaten); reference 11) and the largest species killed. We used $\frac{3}{4}$ of adult body mass estimates to account for sub-adults and young

killed by the predators following previous studies (reference 12). We used the equations derived from these data (Fig. S1; Table S5) to predict the same dietary parameters for extinct species based on their body masses estimates (Table S1).

§ 4) Survey of megaherbivore predation in modern faunas (Table S2).

We surveyed the literature for all records of predation on megaherbivores by carnivores that included some estimate of the age of the prey and in some cases, size of the predator group making the kills. The results are shown in Table S2. We were able to find these data for predation on African elephants and black rhinoceros, but not other megaherbivores such as giraffes and hippos.

§ 5) Historical data on large carnivore group sizes (Table S3)

We searched the literature for data on group sizes (e.g. lion prides, wild dog packs, spotted hyena clans) in the last 150 years. Data were recovered for lions and wild dogs, but not hyenas. The data span the years 1835 to 1997, and the following countries: Ethiopia, Kenya, Namibia, Tanzania, South Africa, and Uganda.

§ 6) Vulnerability of Proboscideans to Predation (Figures S2, S3)

To obtain the size ranges for juvenile proboscidean prey in the different age categories represented in Fig. 2 we combine evidence from behavior, growth, and predation rates of modern proboscideans with isotopic and fossil evidence from fossil mammoths and mastodons. Modern African bush elephants (*Loxodonta africana*) serve as our model because, among modern analogues, the best documentation of size, behavior, and predation at several localities (Table S2) are available for this species. We present ages and sizes as intervals, rather than point estimates, because body size and maturation schedules of both living and Late Pleistocene proboscideans vary among populations within species, with resource availability within species, and other variables such as age of the mother and sex of the offspring (15-19).

Age-specific behavior of juvenile African elephants and predation on them by lions: In Chobe National Park, Botswana, Joubert (20) tallied success rates of over 60% for lions attacking African elephants in age categories 2-4 and 4-9 years (y.), with the largest numbers of elephant kills in the 4 – 9 year old (y.o.) category (even adjusting for its longer duration). Known-age young elephants observed in Amboseli National Park (Kenya) reportedly began consuming plants at ~ 3 months of age, spent an increasing amount of time feeding independently between then and 24 months (the youngest calf observed to survive without milk was orphaned at 26 months), and between 2 and 4.5 y. of age leveled off in spending ~ 55% of the time daily in feeding (16). The mean distance between offspring and mother increased steadily with age, from 2m at age 12 months, to 6 m (for females) or 10 m (for males) in the age category of “old juveniles” 5-7 years old (21). We infer therefore that juvenile elephants’ exposure to predation rises from an age of 2 years onward as their dependence on milk diminishes and they increasingly stray from protection by their mothers. Concurrently, their vulnerability

decreases as they reach body sizes that are difficult for predators to handle. We use ages 2, 4, and 9 as reference points in the calculations below, and present ranges of estimated body sizes for age intervals 2-4, 4-9, and 9 years of age. Even within single populations, rates of growth in stature and the relationship between body mass and shoulder height vary (15). Size estimates for different age classes overlap, so we present size intervals rather than strict cutoff points.

Growth in stature and body mass estimates of juvenile African and Asian elephants: Shrader *et al.* (22) measured standing shoulder heights of 355 known-age elephants from Addo Elephant National Park (South Africa) and Amboseli National Park (Kenya) and derived sex-specific Von Bertalanffy growth curves for predicting age from shoulder heights, with confidence intervals generated through Monte Carlo simulation. Good sampling of youngest age classes allowed age estimation with high precision up to 15 years for females and 36 years in males. Growth curves for young males and females are indistinguishable up to an age of ~ 9 y (shoulder height ~190) at which point they begin to separate, with age predicted for females falling outside the confidence limits for males at heights of ~210 (ages ~ 9.5 and 12, for males and females respectively).

Figure S2A shows intervals of shoulder height (in cm) at which male (m) or female (f) African elephants are predicted by ref. 22 to be 2 (olive), 4 (blue), and 9 (purple) years of age (y.o.), respectively; paler shades represent 95% confidence intervals (c.i.) for those age estimates. (Note that c.i. for 2 and 4 y.o. males overlap; those for 2 and 4 y.o. females do not.) For the intervals shown in Fig. 4 (main text) we estimated body mass for the following shoulder heights: 115 cm (shortest stature at which an age prediction of 2 y. falls within the 95% c.i.), 140 cm (shortest stature at which an age prediction of 4 y. falls within the 95% c.i.), 180 cm (the shortest stature at which age 9 falls within the 95% c.i. of age estimates), and 215 cm (largest size at which age 9 falls within the 95% c.i.). Indicated below the axis in Fig. S2A for comparison are shoulder height estimates for male and female Asian elephants at ages 2, 4 and 9, calculated from von Bertalanffy growth curves modeled on wild-caught Sumatran animals (whose ages at the time of capture were estimated by head mahouts and a veterinarian; ref. 23). (Similarly for Asian elephants in Ceylon, McKay (24) identified a 'juvenile' category ranging 120-180 cm in height or approximately 3-12 years old for females, and 120-200 cm or 3-15 years for males.)

The body masses for African elephants were estimated from a least-squares regression on log-transformed values of 55 masses and shoulder heights of juvenile elephants in the range 71 – 230 cm (Figure S3). Least-squares estimates are appropriate here for reasons outlined by Smith (25). The model sample consisted of 32 heights and masses of *Elephas maximus* (27 provided by Benedict (26), 3 from Flower (27), 2 from Christiansen (28); the largest and smallest individuals in the sample were of this species) and 23 from *Loxodonta africana* (obtained from growth curves of six individuals, two or three time points each, reported by Lang (29), at heights ranging 90 – 190 cm). Average % prediction error was 13.5%; %SEE=19.4.

Figure S2B shows log (body mass) estimates and their 95% prediction intervals (hashmarks extending beyond the color blocks) for shoulder heights of 115, 140, 180, and 215 cm. Colors and boundaries are those used in Fig. 2 of the main text. Body masses of captive Indian elephants at ages 2, 4, and 9 as estimated from formulas provided by Sukumar (30) are also noted, below the axis.

Similarities among species in body sizes of juveniles: As indicated above, young individuals of modern *Loxodonta africana africana* and *Elephas maximus* appear on the whole to be similar in size at corresponding ages. Size estimates for young woolly mammoths show some geographic variability, but are also similar. Lister (31:458) reported that European *Mammuthus primigenius* was “roughly the same body size as living African elephants”. From dental evidence he estimated the age of young juvenile specimens from a Late-glacial (14.5-14 ka BP) assemblage in Shropshire to be in the range of 3-6 years. The diaphysis lengths of juvenile femora, tibiae, humeri, and ulnae in the same assemblage fall in the same ranges as those reported for modern elephants at similar dental stages (32), and during the juvenile period the two modern genera of elephants are similar to one another in their relationships between dental stages and age (cf ref. 33, Table 4 and ref. 34, Table 2 & Fig.7). Maschenko (18) inferred that as neonates, *M. primigenius* from Sevsk (Russia) overlapped the lower end of height distributions for modern elephant neonates, but experienced rapid growth in their first year and reached similar or slightly smaller sizes at 1 year of age. In general, with regard to body size, Haynes (ref. 17:24) presented evidence that “the sizes of cranial and postcranial elements in the skeletons of mastodons and mammoths indicate that in regard to stature, girth, and body length, they may not have been extraordinarily different from modern elephants” but that midshaft diameters of limb bones were greater. Body mass estimates based on (1) shoulder heights, using modern African and Asian elephant height-mass relationships, and (2) lengths or circumferences of limb bones, using interspecific relationships between these measurements and body masses of a wide size range of mammalian taxa, suggest that the Columbian mammoth, *Mammuthus columbi*, sometimes reached body masses exceeding those typically reported for modern elephants (35).

Christiansen (28) derived estimates of body mass for a variety of proboscidean fossils and suggested that as adults they attained sizes substantially larger than living forms. Estimates for *Mammuthus primigenius* from skeletons ranged 3897-10,917 kg; from single elements, 2421-10932 kg; for *M. columbi*, 4980-7859 kg from single bones (and for *M. imperator*, which is commonly synonymized with it, 5215-9143 kg from skeletons, 3466-5045 kg from single bones); for *Mammuthus americanum*, 3600-8953 kg from skeletons; 3267-7672 kg from single bones; and for *Elephas antiquus* 4313-13122 kg from single bones. Some of the limb bone elements from adult mammoths and mastodons are longer and/or wider than those typical for modern elephants, which suggests that these animals did attain larger sizes than the living species. However, caution is warranted in accepting some of the largest estimates, for several reasons: (1) the regression parameters employed were fit to best approximate the relationships between osteological measurements and body masses of seven (4 African, 3 Asian)

modern elephants whose masses ranged 850 – 6434 kg, yet—as the widely differing estimates derived even for single individuals suggest—bone shapes and body proportions of mammoths and mastodons differ from those of modern elephants; (2) often, inevitably, the estimates required extrapolating relationships beyond the size range of the original sample; (3) equations were based on median axis regressions, which yield steeper slopes and higher predictions at the upper ends of the relationships than least squares. For estimating body masses of extinct animals, extrapolating relationships beyond the ranges and outside the body proportions represented by living forms may be unavoidable. For our purposes, however, it is worth noting that the greatest size disparities in these species, whose adults show high sexual size dimorphism, are for adult males, and juveniles of both sexes and both living species tend to be much more similar in size than estimates of maximum adult size might suggest (see Figs. A & B and, *e.g.*, ref. 22 and sources cited therein). On the basis of current evidence we suggest that, while some populations may have been shifted towards the higher ends of the ranges, the size distributions for young mammoths, *Elephas antiquus*, and mastodons at the most vulnerable ages would have overlapped the ranges presented here.

Similarities among species in timing of social independence: Behaviorally the two modern species are on similar schedules: for Asian elephants in Ceylon, McKay (2) distinguished 'infants', which suckle frequently, are small enough to walk under their mothers, and maintain continuous proximity to an adult (ref. 2:9), from the older category of 'juvenile' (age estimates in the range 3-12 y. for females and 3-15 y. for males) and indicated that while young elephants tend to remain in groups with their mothers, "Whenever the group of females is feeding in a relatively stationary position, older infants frequently stray from them and indulge in fairly extensive play behavior" (ref. 2:69) and that "juveniles tend to remain together and form play-groups when a herd is feeding in one spot (ref. 2:9)". For the same species, Eisenberg *et al.* (ref. 36:219) noted that "Young males apparently begin to wander farther from the cow herd from the age of about 6 years on".

Even so, the body masses of adults and timing of sexual maturity can vary widely intraspecifically, and are heavily influenced by resource availability. Laws *et al.* (15), for example, reported ages of attainment of sexual maturity that ranged between 10.77 and 17.2 years for males and 11.73 – 22.8 years for females in five different populations in Uganda, Kenya, and Tanzania.) Resource availability also affects weaning schedules. Laws *et al.* (15) reported mean calving intervals ranging across populations from 2.9 to 9.1 years, roughly in parallel with population density and in inverse correspondence with habitat resources. In Amboseli, mean interbirth intervals for cows conceiving two successive surviving calves ranged from 3.5 y., when conception took place in wet years (and food availability was high), and 5.6 y. during years of low rainfall (16). Calves typically continued suckling until birth of the next calf, but some over 4.5 y.o. were weaned without the birth of another calf and double-suckling continued in the case of a few individuals for up to 12 mo. (16).

By all indications, calving intervals were also similar in extinct elephants (including mammoths) and mastodons. For fossil forms, *Mammuthus americanum* and *Mammuthus primigenius* have been the sources of the most data. Close analysis of annual growth increments in the tusks of female mastodons show a cyclic pattern that Fisher *et al.* (37) (see also, e.g., refs. 38-39) have persuasively interpreted as calving intervals that average 3-4 years. First conception in an individual with “no signs of interrupted growth or any long-term nutritional stress” (Fisher *et al.* ref. 37:461) was estimated to have occurred at an age of ~9-10 years. Growth increments in the tusks of a male mastodon showed an aseasonal pattern that underwent a transition between ages 4 and 5 years to a more regular, seasonal pattern that suggested weaning occurred at that age (40). A drop in the growth rates of tusks occurring at age 10 in male mastodons from seven localities was interpreted as reflecting nutritional stress in adapting to an independent life after their expulsion from the matriarchal family group at sexual maturity (38).

A long-term shift in stable carbon and nitrogen isotope composition in tusks of a 5.5-6 y.o. *M. primigenius* from Wrangel Island (Russia) suggested to Rountrey *et al.* (41) a declining dependence on its mother’s milk and pointed to a lower limit of 5 years of age for weaning, which they compare to ages of weaning of African elephants in high-stress environments. From additional stable isotope analyses on woolly mammoth tusks from Yukon (Canada), Metcalfe *et al.* (42) suggested that juveniles began consuming plants at 2-3 years of age and continued to consume milk at least until age 3. Tooth wear, beginning at an estimated age of 6-7 months in woolly mammoths from Sevsk (Russia) was interpreted by Maschenko (18) to indicate that the juveniles had begun consuming plants at that age, and that they relied upon a diet of vegetation exclusively by an age of 1.5 years. He inferred that the animals reached maturity at 8-10 years and that growth decelerated by 15-17 years with pregnancy and nursing. Recognizing similarities in this timing with *E. maximus*, Maschenko (18) also emphasized the variation among populations living in different environments.

§ 7) Stable Isotope Analyses of Pleistocene Carnivore Diets

A survey of published stable isotope ratio studies of Pleistocene Old and New World hypercarnivores finds relatively little evidence of any single predator species showing a specialization on mammoth or any other megaherbivore species. There are occasional individuals that seem to have favored mammoth or woolly rhino (lion, ref. 43; gray wolf, ref. 44), but most individuals of large prehistoric hypercarnivores were generalists, consuming a mix of large bovids and horses (43-50). The late Pleistocene New World sabertooth cat, *Smilodon*, has been the subject of two stable isotope ratio studies, one in the western United States (46) and the other in southern Chile (49). Both found that *Smilodon* ate a variety of large prey that included various ungulates as well as giant ground sloths, *Mylodon darwini* in Patagonia and *Paramylodon harlani* in California, and there was no evidence of a preference for proboscideans.

Based on the stable isotope ratio data, it would seem that the large hypercarnivores of the Pleistocene Old and New World rarely killed or even scavenged the largest of the megaherbivores, mammoths and rhinoceroses. However, it is important to note that the numbers of individual predators that have been sampled for stable isotopes is relatively small in most cases. With the exception of cave bears and gray wolves, each of which are represented by 70 or more individuals, other hypercarnivorous species (*Panthera atrox*, *P. leo spelaea*, *Homotherium spp.*, *C. crocuta spelaeus*) are represented by fewer than 20 individuals that span the past fifty thousand years. Given this limited sampling, the fact that even occasional individuals stand out as being specialized on megaherbivores suggests that some populations at various times may have been regular hunters of mammoths or woolly rhinos. This is supported by a different line of evidence from the fossil record, carnivore-produced bone accumulations, as discussed in the text.

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TABLE S1. Extant and extinct guilds of large carnivores with body masses >21 kg. References used for body mass estimates are shown, as well as dietary categorization.

Old World late Pleistocene -Scladina Cave, Belgium (80-110 kyr)				
Species	Family	Est Mean Body mass (kg)	Reference	Diet
<i>Crocota crocuta spelaea</i>	Hyaenidae	165	51	Large Prey
<i>Canis lupus</i>	Canidae	26	51	Large Prey
<i>Cuon priscus</i>	Canidae	21	51	Large Prey
<i>Homotherium latidens</i>	Felidae	249	51	Large Prey
<i>Panthera leo spelaea</i>	Felidae	249	51	Large Prey
<i>Panthera pardus</i>	Felidae	198	51	Large Prey
<i>Ursus arctos</i>	Ursidae	68	51	Omnivore
<i>Ursus spelaeus</i>	Ursidae	178	51	Omnivore-Herbivore
OW-Early Pleistocene - Venta Micena, Spain, Upper Villafranchian, circa 1 MA				
<i>Canis falconeri</i>	Canidae	28	51	Large Prey
<i>Canis etruscus</i>	Canidae	21	51	Large Prey
<i>Lynx aff. issiodorensis</i>	Felidae	13	51	Large Prey
<i>Megantereon whitei</i>	Felidae	54	51	Large Prey
<i>Homotherium latidens</i>	Felidae	249	51	Large Prey
<i>Pachycrocuta brevirostris</i>	Hyaenidae	127	51	Large Prey
<i>Ursus etruscus</i>	Ursidae	160	51	Omnivore
NW LATE PLEISTOCENE- Rancho La Brea, California, USA 50,000-10,000 ybp				
<i>Canis lupus</i>	Canidae	35	51	Large Prey
<i>Canis dirus</i>	Canidae	50	51-52	Large Prey
<i>Puma concolor</i>	Felidae	54	51	Large Prey
<i>Panthera onca</i>	Felidae	85	51	Large Prey
<i>Homotherium serus</i>	Felidae	190	51-52	Large Prey
<i>Panthera atrox</i>	Felidae	430	51-52	Large Prey
<i>Smilodon fatalis</i>	Felidae	230	51-52, 54	Large Prey
<i>Ursus americanus</i>	Ursidae	111	51	Omnivore
<i>U. arctos</i>	Ursidae	196	51	Omnivore
<i>Arctodus simus</i>	Ursidae	650	51-53	Omnivore
NW EARLY PLEISTOCENE -Leisey Shell Pit, Florida, USA, Late E. Irvingtonian, circa 1.3 MA				
<i>Canis armbrusteri</i>	Canidae	37	51	Large Prey
<i>Canis edwardii</i>	Canidae	25	51	Large Prey
<i>Xenosmilus sp.</i>	Felidae	328	51	Large Prey
<i>Smilodon gracilis</i>	Felidae	100	51	Large Prey
<i>Miracinonyx inexpectatus</i>	Felidae	57	51	Large Prey
<i>Arctodus pristinus</i>	Ursidae	133	51	Omnivore
Yellowstone National Park, U.S.A.				

<i>Puma concolor</i>	Felidae	60	56	Large Prey
<i>Canis lupus</i>	Canidae	45	3, 57	Large Prey
<i>Canis latrans</i>	Canidae	13	3	Small-Med Prey
<i>Gulo gulo</i>	Mustelidae	14	3, 58	Omnivore
<i>Taxidea taxus</i>	Mustelidae	8.5	3	Omnivore
<i>Ursus americanus</i>	Ursidae	150	3	Omnivore
<i>Ursus arctos</i>	Ursidae	263	3	Omnivore
Royal Chitawan National Park, Nepal				
<i>Panthera tigris</i>	Felidae	162	3, 59	Large Prey
<i>Panthera pardus</i>	Felidae	45	3, 60	Large Prey
<i>Neofelis nebulosa</i>	Felidae	17	3, 60	Large Prey
<i>Cuon alpinus</i>	Canidae	15	3, 61	Large Prey
<i>Melursus ursinus</i>	Ursidae	95	3	Insectivore/omnivore
Etosha National Park, Namibia				
<i>Panthera leo</i>	Felidae	162	3, 62	Large Prey
<i>Panthera pardus</i>	Felidae	45	3,60	Large Prey
<i>Acinonyx jubatus</i>	Felidae	38	3, 63	Large Prey
<i>Lycaon pictus</i>	Canidae	22	3	Large Prey
<i>Crocuta crocuta</i>	Hyaenidae	52	3	Large Prey
<i>Hyaena brunnea</i>	Hyaenidae	41	55, 64	Large Prey/omnivore

Table S2. Recorded kills of megaherbivores with number killed, estimated number of predators involved, location, time period of observation, and reference.

Predator	Prey	Prey Age (yrs)	Predator Group Size	# Killed	Time period	Location	Reference
Lion	African elephant	4-11	>27	4	Oct. 4-25, 2005	Chobe NP, Botswana	65
		4-9 mostly	18	74	1993-1996	Chobe NP, Botswana	20
		young adult	unknown	10	1998-2003	Samburu NP, Kenya	66
		< 7, one 10-yr old bull	8	9	1981-1984	Central African Republic	67
		<8	1-10	44	1998-2004	Hwange NP, Zimbabwe	68
	Black Rhinoceros	3-4	4	3	June-Sept. 1995	Etosha NP, Namibia	69
		old bull	2	1	1960's	Kenya	70
		8 mos.	unknown	1*	2008	Hluhluwe-Mfolozi Park,SA	71
		Almost 2	1 male	1*	1987	Mfolozi Game Reserve	72
Spotted Hyena	Elephant	4 newborn, one 5-yr old	7-10	5	1999	Hwange NP, Zimbabwe	73
	Black Rhinoceros	< 1	unknown	3*	1991-1993	Namibia	74

* inferred but not observed

Table S3. Historical Data on Predator Group Size.

Species	Location	Year	Quote	Citation
Lion, African Wild Dog	Rift Valley, Kenya	1901-1931	"In the days when such well-known personalities as Lord Delamere habitually encountered prides of up to 36 lions on their ranches in Kenya, when hunters all too frequently stumbled on prides of between 17 and 40 on a kill or when the lone horseman found himself surrounded by packs of between 25 and 40 Cape hunting dogs , these animals were sufficiently numerous to keep elephant calf mortality at a high level."	75:253-254
Lion	Etosha NP, Namibia	1989-1997	Lion ..."Pride sizes decreased from 6.3 adult females in 1989 to 2.8 lionesses in 1997. . . 82 % of all known lion mortalities were caused by humans."	76:345
Lion	Ruvana Plains, Tanzania	1903-1926	"During my expedition to Uganda, I met an American who had killed 26 lions in but two weeks" ...On the Ruvana Plain I saw the biggest troop of lions I have ever seen. I counted 26 of them."	77:4.
Lion	Tanzania	1903-1926	"There were still about fifteen lions hidden in the high grass,..."	77:13.
Lion	Ruvana Plain, Tanzania	1903-1926	"...I noticed a number of animals moving and through my field-glass I made them out to be lions. One was a big, heavily maned beast; there was another smaller one and nine lionesses."	77:28.
Lion	Free State, South Africa	1835	Reverend James Archbell reported seeing a group of 18 lions	78: 102.
Lion	Free State, South Africa	1843	Reverend John Bennie reported seeing as many as 14 together.	78: 108.
Lion	South Africa	1902-1946	". . . prides may number as many as thirty individuals, and even more. In fact, one of thirty-five has been recorded in the Kruger National Park."	79:150
Lion	Kenya	1894	"At the end of the last century, Sir Frederick Jackson watched a pride of 23 near Machakos. . ."	80:64

Lion	Kenya	1911	"In 1911, a pride of no less than 40 lions was seen on the Kapiti Plains."	80:64
Lion	Kenya	1951-1952	". . . a pride of 31 was seen in the Marsabit Reserve, and a year later 32 were reported from the Amboseli Reserve."	80:64
Lion	southern Africa	~1950s	"In the border area between northern Rhodesia and Angola, on the Rivers Mashu and Kwandu, prides of 40 or more specimens are by no means rare."	80:64
Lion	South Africa	1880s	"Twelve is the largest number I have seen, though I have heard of as many as fifteen in a troop."	81: Loc. 9780 Kindle edition
Lions	Southern Africa	1887	"In the interior of South Africa, one more commonly meets with four or five lions consorting together, than with single animals, parties of ten or twelve are not uncommon."	82
Lions	Orange River, South Africa	1830s	"Not long since he had see fifteen prowling at the foot of a neighbouring mountain."	83: Loc.1644
Lions	East Africa	1900s	". . . but lions; he counted recent-three or twenty-five of them all around him." "	84: Loc. 685 Kindle Edition
Lions	East Africa	1900s	"Mr H. R. M'Clure of the Government Service is, I believe, responsible for this story. The total I heard was 43..."	84: Loc. 3651
Lions	East Africa	1900s	"The late Mr H. A. F. Currie is said to have shot a lion out of a minimum of 18"	84: Loc. 3659
African wild dogs	Free State, South Africa	1850	J. Leyland saw a pack of 20 wild dogs and claimed packs of 100 were observed.	78: 140 & 141.
African wild dogs	Free State, South Africa	1853	British officer, William St John hunted a pack of 60 wild dogs on 19 th of October 1852, and on the 26 th of January 1853 he came across a pack of 40 or 50.	78: 141.
African Wild Dog	Uganda, Tanzania	1903-1926	"I have seen them hunt in packs of from five to fifty ...I never saw more than fifty in a pack, but hunters declare that packs of a hundred are not rare."	77:259.

African Wild Dog	Tanzania	1903-1926	"I came upon this large pack of thirty-five to forty dogs on the Ngare Dowash one late afternoon during one of my bird-collecting trips."	77:260.
African Wild Dog	Kruger NP, South Africa	1902-1946	"Consequently, they had greatly increased and multiplied and roamed about in packs of fifty or more, ..."	85:55
African wild dogs	East Africa	1900s	"I have seen hundreds of wild dogs, and always in packs - rarely less than five together, and more often from ten to forty."	84: Loc. 832 Kindle Edition
African wild dogs	Ethiopia	1900s	"In Abyssinia I once followed a pack about fifty strong"	84: Loc. 489 Kindle Edition
African wild dogs	East Africa	1900s	"I have stood with thirty or more of them in lines in front of me ..."	84: Loc. 489 Kindle Edition

Table S4. Percent of total number of teeth broken in life for Pleistocene carnivores and the mean value for 36 species of extant carnivores. Data for Pleistocene New World and extant carnivores are from Van Valkenburgh (2009). Data for Zoolithen Cave lions and hyenas were collected by BVV in The Museum fur Naturkunde, Berlin, Germany.

SPECIES	LOCALITY	Total # teeth	% broken
NEW WORLD PLEISTOCENE			
<i>Canis dirus</i>	Rancho La Brea, USA	949	8
<i>Canis latrans</i>	Rancho La Brea, USA	1280	5
<i>Smilodon fatalis</i>	Rancho La Brea, USA	1775	8
<i>Panthera atrox</i>	Rancho La Brea, USA	275	11
<i>Canis dirus</i>	San Josecito Cave, MX	212	4
<i>Canis dirus</i>	Talara Tar seeps, Peru	329	5
<i>Canis lupus</i>	Alaskan permafrost, USA	373	10
<i>Panthera atrox</i>	Alaskan permafrost, USA	129	25
OLD WORLD PLEISTOCENE			
<i>Panthera leo spelaea</i>	Zoolithen Cave, Germany	77	19
<i>Crocota crocuta spelaea</i>	Zoolithen Cave, Germany	161	11
AVG FOR 36 EXTANT SPECIES			2
AVG FOR 11 EXTANT SPECIES >21 KG			2.3 +/- 1.3

Table S5. Estimated body mass equations used to predict extinct species dietary parameters of accessible and largest prey depending upon hunting strategy. See Fig. S1.

Dietary component	Hunting strategy	Equation
Smallest accessible prey	Solitary	$y = 15.74\ln(x) - 33.749$
	Group	$y = 0.6869x + 2.4044$
Largest accessible prey	Solitary	$y = 2.2425x - 19.49$
	Group	$y = 204.78\ln(x) - 279.59$
Largest prey	Solitary	$y = 504.74\ln(x) - 1166.6$
	Group	$y = 22.781x - 92.089$

Table S6. Data from the Absolut Mammal database used for the analysis of the diversity of megaherbivores and hypercarnivores >21 kg in modern mammal communities.

LOCALITY NAME	COUNTRY	# MEGAHERBIVORE S	# HYPERCARNIVOR ES>21 kg
Lunda Norte	Angola	1	4
Huíla Plateau	Angola	3	4
Sevan Lake	Armenia	1	4
Talysh Mountains, Steppe	Azerbaijan	1	2
Okavango Delta	Botswana	4	6
Sangmelima	Cameroon	1	1
Northern Savannah Cameroon	Cameroon	4	5
La Maboké	Central African Republic	2	1
Northern Tibetan Plateau	China	1	2
Xishuangbanna	China	1	3
Mayombe, Kouilou Basin	Congo	1	1
Český les	Czechoslovakia	1	1
Asmara	Eritrea	2	5
Backo, Wollega	Ethiopia	2	2
Southern Lake Tana	Ethiopia	2	2
Dire Dawa	Ethiopia	2	5
Arba Minch	Ethiopia	3	3
Lake Ziway	Ethiopia	3	3
Awash National Park	Ethiopia	3	4
Gondaraba	Ethiopia	4	4
Dohonta	Ethiopia	4	4
Normandie-Maine	France	1	1
Nord-Pas-de-Calais	France	1	1
Colmar	France	1	1
Gamba	Gabon	2	1
Makokou	Gabon	2	2

Greater Caucasus Mountains	Georgia and Azerbaijan	1	3
Kiskunság National Park	Hungary	1	1
Hortobágy National Park	Hungary	1	1
Taï National Park	Ivory Coast	1	1
Lamto	Ivory Coast	2	3
South Turkana National Reserve	Kenya	2	3
Amboseli National Reserve	Kenya	4	5
Lengwe Nature Park/Mwabvi Game Reserve	Malawi	3	4
Lake Malombe	Malawi	3	4
Kasungu National Park	Malawi	3	5
Nyika National Park	Malawi	3	5
Middle Gunung Benom	Malaya	1	3
Low Gunung Benom	Malaya	3	3
LowlandSabah	Malaysia	1	1
Malaysian Lowland Rain Forest	Malaysia	3	3
Maputo Elephant Reserve	Mozambique	3	4
Zinave National Park	Mozambique	3	4
Northern Tete District	Mozambique	3	4
Gilé Wildlife Reserve	Mozambique	3	4
Kaokoveld Desert	Namibia	4	6
Etosha National Park	Namibia	4	6
Langtang National Park	Nepal	1	4
Royal Chitwan National Park	Nepal	3	2
Benin City	Nigeria	1	1
Cross River National Park	Nigeria	2	1
Kainji Lake National Park	Nigeria	2	4
Yankari National Park	Nigeria	3	5
Lake Chad Game Reserve	Nigeria	4	2
Bieszczady Mountains	Poland	1	0
Pila Region	Poland	1	1
Trzebnickie Hills	Poland	1	1

Swietokrzyski National Park	Poland	1	1
Suwalki Region	Poland	1	1
Białowieża National Park	Poland	1	1
Dagestan Caucasus	Russia	1	2
Ciscaucasian Subdistrict	Russia	1	3
Nyungwe National Park	Rwanda	1	2
Akagera National Park	Rwanda	3	4
Kelabit Plateau, Sarawak	Sarawak, Malaysia	1	0
Nord Ferlo	Senegal	2	4
Thabazimbi	South Africa	1	5
Pietersburg	South Africa	2	3
Wakkerstroom	South Africa	2	4
Potchefstroom	South Africa	3	4
Zeerust	South Africa	3	5
Pretoria	South Africa	3	5
Swartwater, Transvaal	South Africa	3	5
Messina	South Africa	3	6
Punda Milia	South Africa	3	6
Germiston	South Africa	4	4
Nelspruit	South Africa	4	5
Komatipoort	South Africa	5	6
Jebel Marra	Sudan	1	4
Udzungwa Mountains	Tanzania	2	3
Lake Rukwa Valley	Tanzania	4	5
Serengeti National Park	Tanzania	4	6
Central Vietnam	Vietnam	2	2
South-Central Vietnam	Vietnam	3	3
Southwestern Kivu Lake	Zaire	1	1
Kibara Plateau, Upemba National Park	Zaire	1	5
Ituri Forest	Zaire	2	1
Lake Upemba, Upemba National	Zaire	2	3

Park			
Rwindi–Rutshuru Plain, Virunga National Park	Zaire	2	4
Bagbele	Zaire	4	2
Garamba National Park	Zaire	4	4
Kafue National Park	Zambia	2	5

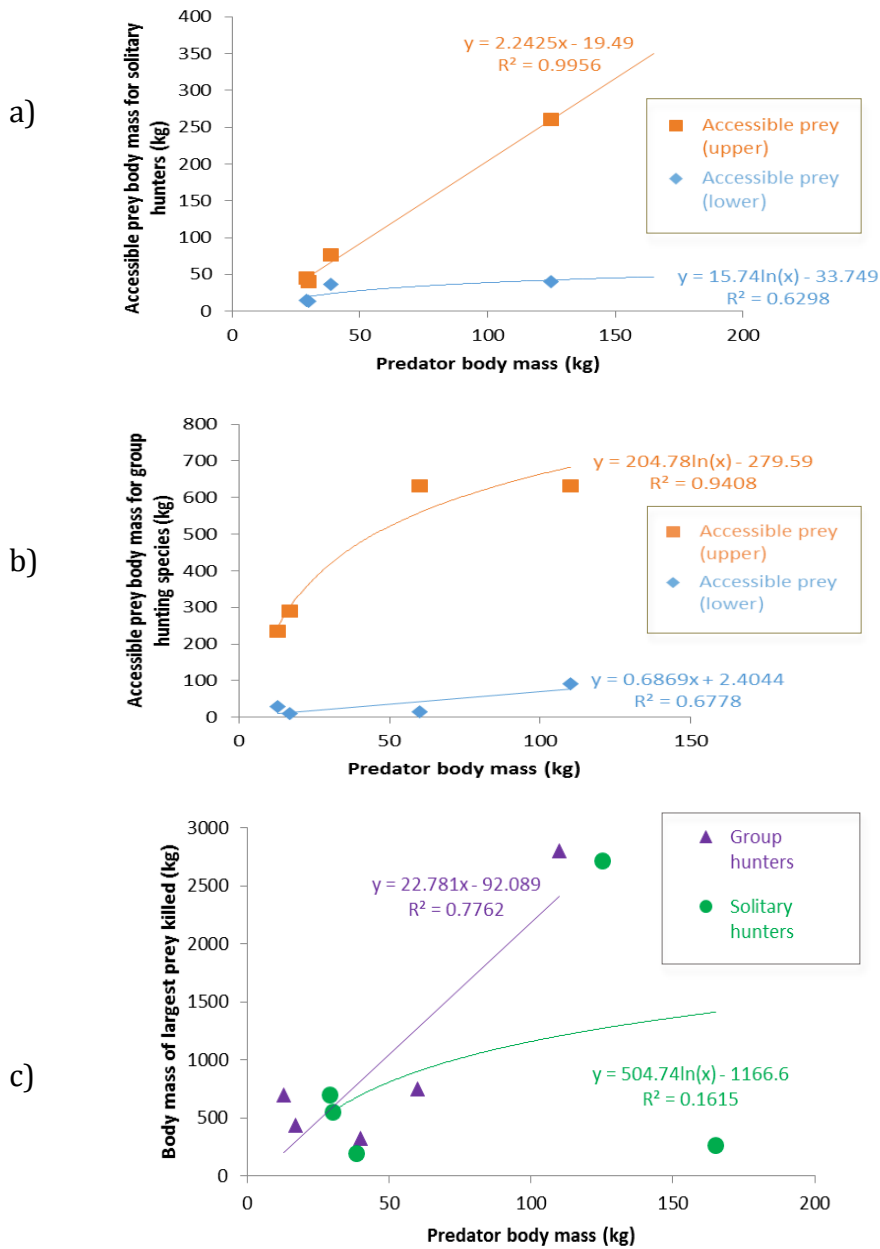


Figure S1. Relationships based on extant predators that were used to predict the upper and lower accessible prey of a) solitary and b) group hunting predators, and the largest (c) prey killed by extinct Pleistocene predators.

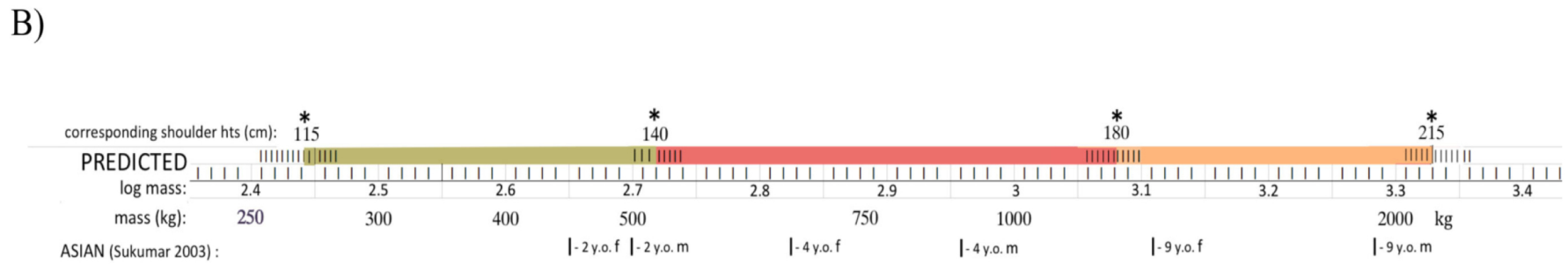
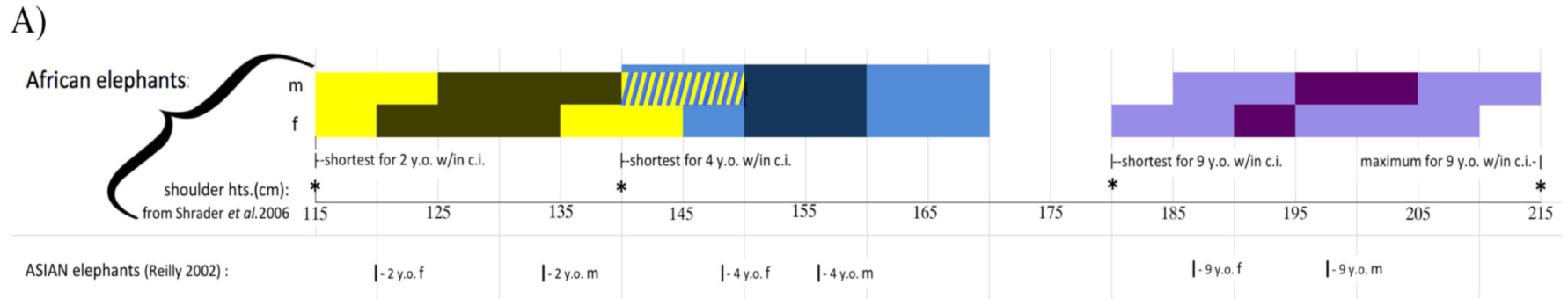


Figure S2: A) shows intervals of shoulder height (in cm) at which male (m) or female (f) African elephants are predicted by ref. (1) to be two (yellow), four (blue), and nine (purple) years of age (y.o.), respectively; paler shades represent 95% confidence intervals (c.i.) for those age estimates. (Note that c.i.s for two and four y.o. males overlap; those for two and four y.o. females do not. The c.i.s for predicted ages greater than nine y.o. broaden greatly and show considerable overlap with one another.) For the intervals shown in Fig. 2 (main text) we estimated body mass for the following shoulder heights: 115 cm (shortest stature at which an age prediction of two y. falls within the 95% c.i.), 140 cm (shortest stature at which an age prediction of four y. falls within the 95% c.i.), 180 cm (the shortest stature at which age nine falls within the 95% c.i. of age estimates), and 215 cm (largest size at which age 9 falls within the 95% c.i.). Indicated below the axis in Fig. S2A for comparison are shoulder height estimates for male and female Asian elephants at ages two, four and nine, calculated from von Bertalanffy growth curves modeled on wild-caught Sumatran animals (whose ages at the time of capture were estimated by head mahouts and a veterinarian; ref. 1). (Similarly for Asian elephants in Ceylon, McKay (2) identified a 'juvenile' category ranging 120-180 cm in height or approximately 3-12 years old for females, and 120-200 cm or 3-15 years for males.) **B)** shows log (body mass) estimates and their 95% prediction intervals (hashmarks extending beyond the color blocks) for shoulder heights of 115, 140, 180, and 215 cm. Colors and boundaries are those used in Fig. 2 of the main text. Body masses of captive Indian elephants at ages 2, 4, and 9 as estimated from formulas provided by Sukumar (3) are also noted, below the axis.

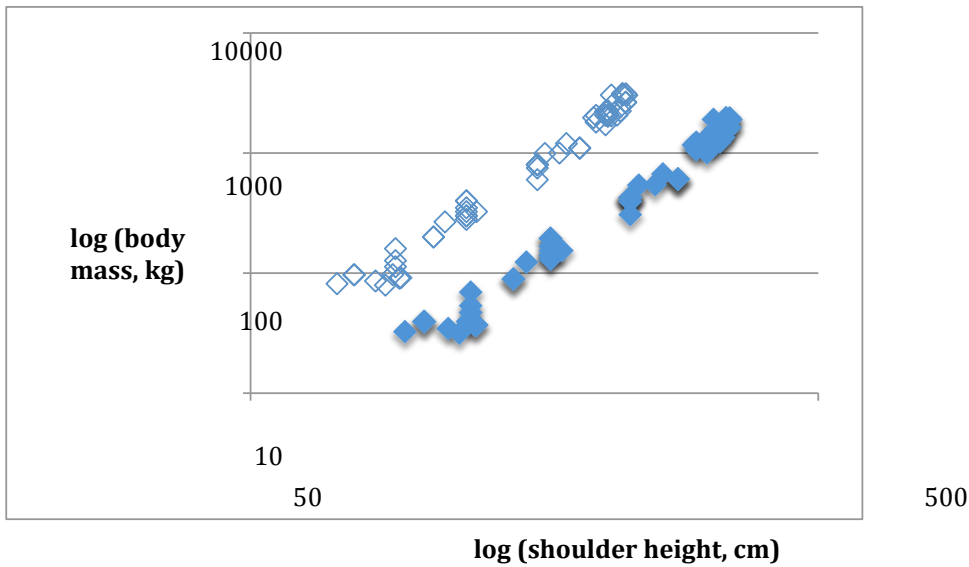


Figure S3. Least-squares regression on log-transformed values of 55 masses and shoulder heights of juvenile elephants in the range 71 – 230 cm. Equation: $\log \text{ mass} = -4.3283 + (3.2848 * \log \text{ ht})$.

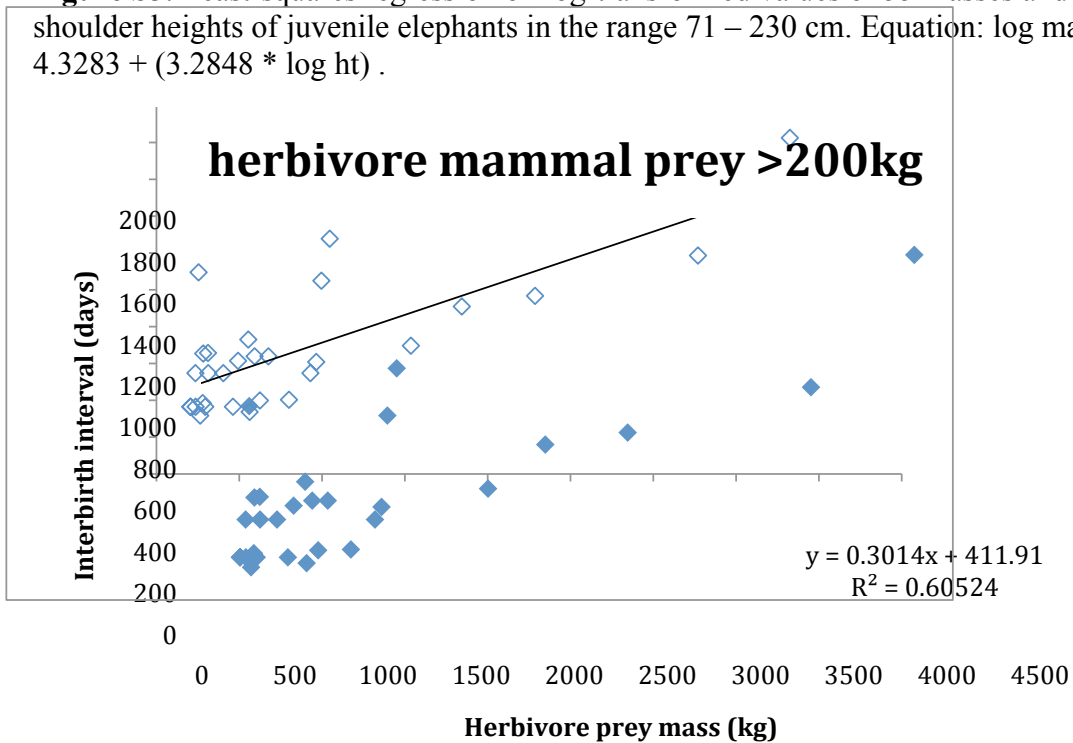


Figure S4. Least-squares regression of interbirth interval in days against body mass for 29 species of herbivores with masses greater than 200 kg. Data derived from PanTHERIA database for extant mammals (<http://esapubs.org/archive/ecol/e090/184/metadata.htm>).