

1 **Herbivores, resources and risks: alternating regulation along primary**
2 **environmental gradients in savannas.**

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21

22 **Abstract**

23 Herbivores are regulated by predation under certain environmental conditions, whereas in
24 others they are limited by forage abundance and nutritional quality. Whether top-down or
25 bottom-up regulation prevails depends both on abiotic constraints on forage availability
26 and body size, because size simultaneously affects herbivores' risk of predation and their
27 nutritional demands. Consequently, ecosystems composed of similar species can have
28 different dynamics if they differ in resource supply. We use large herbivore assemblages
29 in African savanna ecosystems to develop a framework that connects environmental
30 gradients and disturbance patterns with body size and trophic structure. This framework
31 provides a model for understanding the functioning and diversity of ecosystems in
32 general, and unifies how top-down and bottom-up mechanisms are dependent on
33 common underlying environmental gradients.

34

35 **Herbivore regulation and the implications of body size**

36 The global decline of large herbivores, due to human-induced landland use changes,
37 raises concerns for the long-term conservation of species whose ranges are being reduced
38 to a handful of protected areas [1]. The local extirpation of large herbivores has
39 consequences for entire ecosystems, because of their role in maintaining the diversity of
40 predators and primary producers [2]. Understanding herbivore regulation across resource
41 gradients, such as rainfall, is important for the long-term management and conservation
42 of ecosystems, especially if shifts in global climate result in a mismatch between the
43 location of protected areas and a species' preferred niche. Here, we investigate how

44 resource gradients simultaneously influence top-down and bottom-up processes in
45 ecosystems, using the large herbivore community of African savannas as a generalized
46 example. The model could also prove useful in understanding the relation between
47 disturbance, resource gradients and trophic structure in other ecosystems.

48

49 Classic food chains represent relationships between trophic levels as linear bottom-up or
50 top-down processes: abiotic factors such as rain determine primary production, which is
51 consumed by herbivores, which are, in turn, consumed by carnivores. The abundance of
52 herbivores can therefore be controlled through top-down mechanisms, such as predation
53 [3-5], or through bottom-up constraints on primary production, such as soil fertility
54 (Figure 1) [6-9].

55

56 Trophic cascades in linear models of herbivore regulation (Figure 1,) involve the knock-
57 on effects of predation expressed at alternate trophic levels [10]. In the classic example,
58 predators limit the abundance of herbivores, which releases grazing pressure on plants
59 (the ‘green world’ hypothesis) [11]. In this hypothesis, the abundance of vegetation is
60 determined largely by the availability of abiotic resources as herbivores are regulated by
61 predators [5, 12].

62

63 Previous research has elucidated the complexity of trophic interactions by breaking each
64 trophic level into more fundamental components (Figure 1). Specifically, the role of
65 abiotic factors, disturbances, quality and quantity of primary production, and the effect of
66 body size have each been shown to influence independently the distribution and

67 abundance of herbivores. Here, we show how common underlying environmental
68 gradients influence both top-down and bottom-up regulation simultaneously. Differences
69 in the relative accessibility of limiting resources can cause ecosystems with similar
70 species to have different regulatory mechanisms. In addition, the body sizes of herbivores
71 determine both their susceptibility to predators and their resource requirements. We
72 show how this generalized model accounts for observed differences in the trophic
73 functioning of the large herbivore community in savanna sites across Africa. Although
74 humans evolved in African savannas and historically affected herbivores through hunting
75 and fire, substantial landuse changes and increasing human populations have put
76 unnatural demands on these systems, which calls for a better understanding of ecosystem
77 dynamics.

78

79 **Predation: not all herbivores are affected equally**

80 The simple food-chain view of predator-prey interactions (Figure 1) ignores the fact that
81 not all carnivores can consume all herbivores, and not all herbivores are equally
82 susceptible to all carnivores. Large prey, such as buffalo (*Syncerus caffer*), are difficult to
83 capture and are only consumed by the largest predators, such as lion. Whereas small
84 predators can only consume small prey, large predators might consume both large and
85 small prey (Figure 2a) [13, 14]. Recent work proposes that predation has a greater impact
86 on regulating a population of small body-sized herbivores (e.g. oribi, *Ourebia ourebi*)
87 when the prey base of small predators is nested within that of large predators as this
88 exposes smaller herbivores to more enemies (Figure 2a) [15]. Conversely, if predators
89 specialize on particular size classes of prey (i.e. they partition the prey base), predation

90 pressure is expected to be relatively even across all body sizes, until the prey become too
91 large (Figure 2b).

92

93 Therefore, the degree of regulation of herbivore populations by predation depends on
94 both the relative body size of predators and prey, and the way in which the available prey
95 base is partitioned by carnivores of different sizes (Figure 3a). Furthermore, the largest
96 predators (lion) are also the most dominant and actively kill smaller predators (cheetah,
97 hyena), or chase them away from their prey (Figure 3a) [16, 17].

98

99 Recent studies provide evidence that both size-nested and size-partitioned predation
100 occurs (Figure 3b). In the Serengeti system (Tanzania) large prey, such as buffalo
101 (450kg), generally escape predation except from cooperatively hunting lions, while oribi
102 (18 kg) are eaten by many species [3, 18] illustrating size-nested predation (Figure 3b).
103 Studies in Kruger Park, Hluhluwe-iMfolozi, and Phinda (South Africa) provide evidence
104 for size-partitioned predation where large predators, such as lion, concentrate on prey
105 centered around the predators' body mass [4, 19] (but see Ref. [15]). Only prey species
106 with a body mass >1000kg, such as hippo (*Hippopotamus amphibius*) and white rhino
107 (*Ceratotherium simum*), tend to escape predator regulation in Kruger, whereas prey
108 species heavier than 150kg, such as buffalo, escape predator regulation in Serengeti. The
109 largest herbivores, such as elephant (*Loxodonta africana*), are too big to be captured
110 irrespective of whether predation is size-nested or size-partitioned [20, 21] (with the
111 exception of juveniles).

112

113 In summary, top-down processes are modified by the way in which carnivores partition
114 their food niches and the degree to which larger carnivores dominate smaller carnivores.
115 Explanations as to why there are differences in the mechanism of niche partitioning of
116 predators in otherwise similar food chains can be found in the type of vegetation that
117 supports herbivores and the disturbance regime, which we explore below.

118

119 **Forage quality and abundance: not all that is green is edible**

120 Geographic processes involving erosion of parent material, and rainfall, determine key
121 environmental gradients, such as soil fertility and water availability [22, 23], which
122 influence vegetation structure [7, 24, 25]. Plant structure, in turn, determines the quality
123 and quantity of digestible material available to herbivores (Figure 4a) [9, 26-30]. Primary
124 production varies along environmental gradients [31] and regulates herbivore populations
125 through classic bottom-up processes of resource limitation (Figure 4a) [8, 32-34]. The
126 quantity of primary production increases with rainfall and soil fertility, such that in the
127 absence of herbivory or fire, the largest standing biomass is found in fertile areas with
128 unlimited moisture while declining when either water or soil nutrients become limiting
129 (Figure 4b) [7, 29, 35]. Under high rainfall conditions, plants invest more resources in
130 structural support and protection against herbivory (e.g. stems, lignified tissues,
131 secondary compounds and mechanical defenses [35, 36]). As a result, the digestible
132 quality of primary production is inversely related to rainfall (Figure 4b), such that the
133 amount of energy and nutrients per unit biomass that is extractable by herbivores declines
134 as conditions become wetter [37].

135

136 Herbivory provides the most direct access to consumable energy and has evolved
137 multiple times in many unrelated taxa (e.g. molluscs, birds, mammals, insects, reptiles,
138 fish, and marsupials) from both carnivorous and detritivorous ancestors [38]. To digest
139 cellulose, ungulates use a symbiotic fermentation process in the rumen or caecum that is
140 relatively time-consuming and requires a specialized gastrointestinal tract. Small
141 ungulates like oribi and Thomson's gazelle (*Gazella thomsoni*) have smaller
142 gastrointestinal systems, and therefore shorter ingesta retention times [20, 21, 39], which
143 means they cannot process coarse vegetation. Furthermore, small endotherms have a
144 higher energy expenditure per unit mass. These two factors mean that small herbivores
145 have to select the most nutritious, highest energy forage (Figure 4c) [26, 28, 40, 41].
146 Larger herbivores are relatively unconstrained by the size of their gastrointestinal tract,
147 they have longer retention times, and so can extract sufficient energy from poorer quality
148 food, providing there is sufficient quantity [40, 42] (Figure 4c).

149

150 In summary, plant quality and biomass are determined by both environmental gradients
151 and plant growth form (Figure 4a). These affect small and large herbivores differently
152 due to differing metabolic constraints. The result is that smaller herbivore populations are
153 nutritionally limited by the quality of forage, whereas populations of larger grazers are
154 limited by the quantity of food (Figure 4d).

155

156 **The role of disturbances, facilitation and ecosystem engineers**

157 Sudden shifts in primary production caused by abiotic disturbances such as fire, or
158 marked changes in consumption rates (either herbivory, predation or infection)

159 potentially rearrange the dynamics of an ecosystem either temporarily or semi-
160 permanently into a new state [43, 44]. Given that local densities of herbivores can be
161 regulated by both predation and attributes of the plant community (e.g. structure, quality
162 and quantity), disturbances are factors that can change the primary mechanism of
163 herbivore regulation and lead to nonlinear responses in abundance (Figure 5a) [22, 45-
164 47].

165

166 Reciprocal effects occur between large herbivores and primary producers (represented
167 with double arrows in Figure 5a) that can lead to grazing facilitation between species [25,
168 48, 49]. Mega-herbivores, such as white rhino or hippo, create and maintain low-biomass
169 grass swards composed of nutritious grazing-tolerant grasses, which subsequently support
170 other smaller grazers [50, 51]. The repetitive grazing of specific patches by multiple
171 species where more dung and urine are deposited might have similar consequences,
172 resulting in fertile hotspot locations where grazers consistently occur over time [52, 53].

173

174 Reciprocal interactions alter the probability of a disturbance occurring in an ecosystem,
175 whereas interaction modifiers (as per Ref. [14]) alter the severity of a disturbance. For
176 example, the relative proportion of trees and grasses in a savanna influences its potential
177 flammability [54] because grasses (which senesce seasonally) contribute more to the
178 fuel load than do trees. Once grasses dominate the plant community, owing to
179 disturbances such as herbivory (e.g. Ref. [55]), a positive feedback between grass
180 abundance and fire frequency can arise (double arrow in Figure 5a). In addition, the
181 accumulation of grass biomass also alters the intensity of a fire (dotted line in Figure 5a),

182 which when combined with the positive feedbacks between fire frequency and grass
183 abundance, maintains an open grassland landscape, preventing tree invasion [54, 56-58].
184 As a result, fires can prevent invasion of grasslands by trees, which is engineered, in part,
185 by the grazing intensity of herbivores [59, 60].

186

187 Disturbances such as grazing and fire can act additively in savanna systems by changing
188 the competitive balance between grazing tolerant and intolerant grasses (Figure 5b) [46]
189 and influencing the nutritional quality of the forage supporting herbivores (Figure 5c)
190 [61, 62]. So where some savannas have sufficient rainfall to support closed forests, they
191 persist as mixed grasslands owing primarily to disturbances [46, 63]. Thus, two systems
192 with similar rainfall and nutrient regimes could have different woodland-grassland
193 structures because disturbances push systems between multiple states [55, 58] and this, in
194 turn, affects the abundance of herbivores.

195

196 **Emerging properties: top-down and bottom-up processes are not mutually**
197 **exclusive**

198 The separate roles of predation (Figure 3a), primary production (Figure 4a), or
199 disturbance (Figure 5a) in the regulation of herbivore populations have different
200 consequences when they are combined as opposed to when considered separately.
201 Underlying environmental and landscape gradients affect top-down and bottom-up
202 processes simultaneously [2] by influencing the forage quality and quantity available to
203 herbivores while changing their exposure to predation (Figure 6a). Thus, in Kruger
204 rainfall simultaneously affects predation and primary production [64]. Moreover, the

205 topography of the landscape (e.g. hills, catenas and rivers) affects the quality and
206 abundance of primary production available to herbivores [7, 22, 45], while at the same
207 time determines the amount of cover for a predator, and increases their hunting success
208 by concealing them from their prey (Figure 6b, c) [65-69]. These indirect (Figure 4a),
209 additive (Figure 5a), reciprocal (Figure 5a), and interaction-modifying (Figure 5a)
210 relationships strengthen the interdependencies between primary production, disturbances
211 and predation in regulating herbivores (Figure 6a).

212

213 **The scale of heterogeneity determines the mode of regulation**

214 Top-down and bottom-up processes are modified by the scale of both spatial and
215 temporal heterogeneity [70]. Suitable habitats can occur heterogeneously at a coarse scale
216 where large patches are separated by long distances, or as a fine-scale mosaic [71, 72].
217 When heterogeneity is coarse, herbivores migrate long distances between suitable
218 patches. When habitats are locally heterogeneous animals move frequently between small
219 patches but do not move far. On a temporal scale, strong seasonality causes animals to
220 move between patches as phenological conditions change (such as the seasonal drying of
221 grass), whereas weak seasonality enables animals to remain in local areas. The
222 consequences of these different scales are seen in the long-distance seasonal migrations
223 of wildebeest (*Connochaetes taurinus*) in Serengeti, Coke's hartebeest (*Alcelaphus*
224 *buselaphus*) on the Athi plains of Kenya, or white-eared kob (*Kobus kob leucotis*), topi
225 (*Damaliscus lunatus*) and Mongalla gazelle (*Gazella thomsoni albonotata*) in Sudan [1].
226 Such movements reduce the impact of predation as predators cannot follow the
227 herbivores over these long distances [73, 74] and result in bottom-up regulation [34] (Box

228 1). By contrast, fine-scale spatial and temporal heterogeneity keeps herbivores within the
229 territories of predators and results in top-down regulation. This distinction is seen by
230 comparing resident wildebeest in Pilanesberg, Hluhluwe-iMfolozi, Kruger (South
231 Africa), Ngorongoro Crater (Tanzania), and Etosha (Namibia) [4, 75-78] which are
232 predator-regulated, to migrant wildebeest in Serengeti [34] which are food-regulated
233 (Box 1). Therefore, abiotic gradients influencing the regulation of herbivores at large
234 scales through predation and nutrition are replicated at finer scales and determine how
235 animals are distributed over a landscape [79].

236

237 **Abiotic gradients determine the direction of regulation**

238 The availability of abiotic factors, such as soil nutrients and rainfall, determines the mode
239 of population regulation of herbivores [2, 80]. High nutrient supply, such as in volcanic
240 or riparian soils, leads to higher quality plant forage as seen in eastern Kruger, southern
241 Serengeti and Samburu (Kenya) [71, 81]. Plants with high nutrient content and low
242 amounts of fiber can support animals of small body size that are top-down regulated. By
243 contrast, sandy soils of granitic origin are low in nutrients, and result in fibrous plants that
244 are less digestible. Such plants are eaten by large herbivores, such as elephants and
245 buffalo, which are bottom-up regulated [29, 42, 82]. This gradation forms the basis for
246 our interpretation of the different savanna systems in Africa, which we describe below.

247

248 **Synthesis: Shifting regulation across gradients of productivity in African savannas**

249 The integration of allometric differences in susceptibility to top-down and bottom-up
250 regulation along gradients of productivity enables an interpretation of the various
251 herbivore dynamics across African savannas (Figure 7). The relative importance of
252 predation, forage quality and forage abundance in regulating grazers should change under
253 different rainfall and soil fertility regimes. Figure 7 proposes how different areas in
254 Africa, often with functionally similar species, could be regulated by different
255 mechanisms along gradients of rainfall and soil fertility [27-29].

256

257 The highest vegetative biomass occurs with high water availability (rainfall or flooding)
258 on fertile soils usually of volcanic or fluvial origin seen best in the Ruwenzori grasslands
259 (Uganda), the Nile floodplains and the flooded Boma grasslands of southwest Sudan.
260 (Figures 3b, 6a) [61, 83-85]. Under these moist fertile conditions, the quantity of food is
261 effectively unlimited and, therefore, regulates only the largest herbivores (dashed blue
262 line, Figure 7a). However, high rainfall causes the grass to have a large proportion of
263 poor quality stems, making digestion more difficult and reducing the overall nutritional
264 quality (dashed green line, Figure 7a). The high standing biomass under high rainfall
265 conditions also conceals predators, making small grazers more susceptible to top-down
266 effects (solid red line, Figure 7a). As a result, under high rainfall and fertile conditions,
267 small herbivores become regulated by predation, as in the Ruwenzori system [86],
268 whereas medium and large herbivores are regulated by the quality of the available forage
269 since the quantity is effectively limitless.

270

271 In contrast, forage quality is highest under low rainfall conditions on fertile soils, such as
272 on the open treeless Serengeti plains and the saline soils of Etosha. Here, the low grass
273 biomass exposes predators making them visible to herbivores, and results in minimal top-
274 down regulation. Instead, the quantity of forage regulates the number of medium and
275 large grazers, with predation accounting for most of the mortality in only the smallest size
276 classes (Figure 7b) [41]. The quality of the grass has little impact on regulating
277 populations, because nutritious forage is abundant and readily available.

278

279 The nutritional quality of grass is lowest in high rainfall areas with infertile soils and,
280 therefore, becomes more important in regulating herbivore populations (Figure 7c). Such
281 areas can be found in the broad-leaved miombo woodlands of the south and central
282 African plateau, such as Selous (Tanzania), Hwange (Zimbabwe), Chobe (Botswana) and
283 Kafue (Zambia), the coastal sand dune savannas of St. Lucia (South Africa) and the
284 Maputu Elephant Reserve (Mozambique), and the moist savannas of Garamba (DRC) and
285 Mole (Ghana) [29, 87-89]. In these communities, there is a high biomass of woody
286 vegetation but also of tall grasses of lower nutritional value, which support fewer
287 herbivores and predators than areas with fertile soils. Predation is expected to be high
288 because the high grass biomass conceals predators, making them more effective [66, 68].
289 The requirements of small herbivores for high forage quality combined with their
290 susceptibility to predation limit their abundance in these areas (Figure 7c). The dominant
291 grazers are elephants [29, 89]. Even mega-herbivores might not be able to escape the
292 limitations of quality because the methane production associated with ingesting a lot of
293 low quality food reduces their capacity to absorb nutrients [39].

294

295 Semi-arid systems with infertile soils, such as Tsavo and Samburu (Kenya), and the
296 Kgalagadi Transfrontier (South Africa and Botswana), tend to have sparse and
297 nutritionally poor plants, supporting lower herbivore densities and, as a result, fewer
298 predators [29, 90, 91]. These conditions are more extreme in desert environments, such
299 as the stony plains of the Jiddah (Oman) home of the Arabian oryx (*Oryx leucoryx*) [92],
300 and the Ouadi Achim Faunal Reserve (Chad) where the scimitar-horned oryx (*O.*
301 *dammah*) used to live [93], and the Skeleton Coast (Namibia). Under these conditions,
302 predation does not limit herbivore abundance. Herbivores are regulated by forage quality
303 (especially for smaller herbivores) and forage quantity (for larger animals) (Figure 7d).

304

305 **Conclusions and future directions**

306 Predation and competition for resources interact synergistically rather than operate
307 independently [94]. Reciprocal, indirect, additive and interaction modifying relationships
308 shape this synergism to explain functional differences between ecosystems. In essence,
309 the interplay between: (i) the availability of limited abiotic resources (such as nutrients
310 and rainfall) that determine the quality and quantity of primary production; (ii) the
311 evolutionary tradeoffs related to body size (including predation sensitivity, digestive
312 capacity and metabolic requirements); (iii) adaptive behaviors (such as migration or
313 group vigilance), which enable primary consumers to escape regulation; and (iv) the
314 extent and frequency of disturbances (such as fires, storms, extreme temperatures, salinity
315 shifts, scouring, etc.) are processes affecting how predation and competition collectively
316 structure communities. This conceptual structure yields testable predictions for how

317 global environmental changes might affect the distribution of different sized herbivores
318 and potential regime shifts in ecosystem dynamics [95]. For example, changes in rainfall
319 due to global warming could shift the importance of food and predation in regulating
320 herbivore populations, such that decreasing rainfall would push an ecosystem along the x-
321 axis in Figure 7 from panel a to b, or c to d. The evolutionary role of early hunter-
322 gatherer humans in regulating herbivores as predators and as agents of disturbance fits the
323 framework of Figure 6. However, modern humans have escaped from factors regulating
324 their population density, which destabilizes framework.

325

326 Future research should test the predictions of Figure 7. More data are required to resolve
327 the consequences of predation. Specifically, the analysis of herbivore carcasses suggests
328 small prey are prone to many predators (i.e. size-nested predation, Figure 2a) [3], but this
329 is not supported by data on carnivore diets which suggest predation is size-partitioned
330 (Figure 2b) [4]. In addition, an evaluation of mortality in juvenile age classes might show
331 that predation by a single predator, with low capture rates, could still impose strong
332 population regulation, especially for larger species.

333

334 In summary, we propose that abiotic factors determine the importance of predation,
335 forage quality and forage abundance in regulating herbivores of different sizes (Figure 7)
336 and this alters the relative strength of the connections between biotic and abiotic
337 components in ecosystems (Figure 6a). The availability of key environmental resources
338 has profound consequences for herbivore regulation and ecosystem dynamics by
339 simultaneously affecting multiple top-down and bottom-up processes. The different

340 herbivore dynamics of the many savanna systems of Africa can be understood in the
341 context of this framework. These concepts could help our understanding of other
342 ecosystems where strong abiotic gradients influence the shape of the community (such as
343 salinity and desiccation in intertidal ecosystems, dissolved oxygen and opacity in aquatic
344 ecosystems, or body mass and predation risk in avian communities). The strength of this
345 framework is that it captures how environmental gradients can switch top-down and
346 bottom-up processes that regulate animal abundance.

347

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- 574
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577 Box 1. Adaptive responses to regulation: migrations, crypsis and vigilance

578

579 Large herbivores are occasionally found in areas of low plant biomass while small
580 herbivores are sometimes found in areas of high plant biomass, which is contrary to the
581 expectations of Figure 7 (main text). The realized niche of many herbivores differs from
582 expected because they escape regulation through behavioral and physiological
583 adaptations selected through evolutionary time [26].

584

585 For example, migrations enable a population to escape the limitations of both low forage
586 biomass and predation. By sequentially moving between the best available food patches
587 migrants are in essence increasing the total available biomass of highest quality food,
588 without exhausting the overall food supply. Small resident populations of wildebeest in
589 Pilanesberg, Kruger and Serengeti are regulated by predation [64, 77], whereas the large
590 migratory wildebeest population in the Serengeti (~1.2 million) are regulated by the
591 availability of sufficient forage especially during the dry season when food is most
592 limited [34]. Other resident herbivore species in Kruger (e.g. roan, *Hippotragus equinus*)
593 and Serengeti become more predator regulated in areas they share with migrant
594 populations, because the seasonal influx of migratory prey supports more predators that
595 tend to supplement their diet with resident prey [80]. As a result, the abundances of
596 migrants versus residents can differ by orders of magnitude [73, 74, 76]. In
597 circumstances where migratory routes are blocked the population declines rapidly [1] and
598 becomes predator regulated. Migrations explain how very large herbivores can persist in
599 low biomass areas, such as the paradox of mammoths which occupied the low biomass

600 habitats of the tundra. Other behavioral adaptations to predation include group dilution
601 and group vigilance where individuals decrease their probability of being caught [96].
602
603 Avoiding predators through habitat selection or crypsis requires physiological
604 adaptations. For instance, addax (*Addax nasomaculatus*), gemsbok (*Oryx gazelle*) and
605 Arabian oryx avoid predators by living in very arid habitats, however they require
606 complex adaptations for water retention and cooling [97]. Digestive adaptations are
607 required especially by small energetically constrained herbivores relying on crypsis in
608 thick low-quality vegetation [98]. In both circumstances, populations escape predation
609 but become regulated at much lower densities by forage quantity and quality,
610 respectively.

611 Figure 1. Several factors are responsible for regulating herbivore abundances across
612 ecosystems. In this classic food chain approach, the abundance of herbivores is
613 regulated by top-down processes, such as predation, and by bottom-up processes
614 through primary production (arrows indicate the direction of influence).
615 Herbivores (like carnivores) are considered as a single unit, despite showing strong
616 functional divergences based on body size. Primary production in this framework
617 is a general term that fails to distinguish the differences between the quality,
618 quantity, and structure of a plant community.

620 Figure 2. The degree of herbivore mortality due to predation depends on the predators'
621 diet selection. (a) If large predators are opportunists and consume prey of all sizes,
622 while small predators only kill small prey, then the prey base of small predators is
623 nested within that of large predators (size-nested predation). The cumulative
624 mortality on small prey is much greater than large prey because they are exposed to
625 more predators. (b) If predators are selective, and only consume prey of a specific
626 size classes (size-partitioned predation), then large predators do not supplement
627 their diet with small prey. When predation is size-partitioned as opposed to size-
628 nested, the cumulative mortality due to predation on small prey is much less,
629 whereas large prey are killed more often.

631 Figure 3. The relative importance of predation in regulating herbivore populations
632 depends on the body size of predators and their degree of specialization for certain
633 prey. (a) If large carnivores only eat large prey, and small carnivores only eat small
634 prey (solid vertical lines), the prey base is partitioned. If large carnivores eat both
635 large and small prey (solid and dashed vertical lines), the prey base of smaller
636 carnivores is nested within that of larger carnivores. Large carnivores dominate
637 small carnivores and reduce their efficiency (solid horizontal line). (b) When
638 predation is nested, small prey are exposed to more predator species and become
639 increasingly predator regulated, as in the Serengeti example (solid line). When
640 predation is partitioned, large prey suffer greater predation than do small prey
641 because large predators do not supplement their diet with small prey, as in the
642 Kruger example (dashed line). Data for Serengeti and Kruger reproduced with
643 permission from Refs. [3, 4, 64]. (Abbreviations: B = African Buffalo, E =
644 Elephant, G = Giraffe, H = Hippo, I = Impala, O = Oribi, R = Black Rhino, T =
645 Topi, W = resident Wildebeest, Z = resident Zebra)

648 Figure 4. The quality versus quantity of primary production regulates large herbivores
649 differently from small herbivores. (a) Physical and environmental gradients have
650 direct and indirect affects (i.e. between non-adjacent levels) on the plant
651 community structure and on the quality and quantity of primary production. (b) For
652 example, the quantity of plant biomass is positively related to increasing rainfall
653 and soil fertility, whereas the digestible quality of the plant declines with increasing
654 rainfall. (c) Large herbivores consume greater quantities of lower quality food,
655 whereas small herbivores consume less food of higher quality because they are
656 constrained by their high metabolism and limited digestive capacity. (d) Therefore,

657 large herbivores, such as elephant, tend to be regulated by food abundance (dashed
658 blue line), whereas smaller herbivores, such as wildebeest, are regulated by food
659 quality (dotted green line). The smallest herbivores, such as oribi, are mainly
660 predator regulated (solid red line). Reproduced with permission from Refs. [35] (b)
661 and [21] (c).
662
663

664 Figure 5. Abiotic and biotic disturbances, such as fires or intense grazing, alter the
665 primary production and the plant community structure of a landscape which, in
666 turn, affects the abundance of different sized herbivores. (a) Disturbances can have
667 reciprocal effects (double arrows) on primary production. For example, herbivores
668 can reduce the biomass of grass, which reduces the probability of fire; but,
669 conversely, fires removes grass necromass which stimulates re-growth of high
670 quality shoots that are preferred by herbivores. Interaction-modifying relationships
671 (dotted arrow) alter the effects of a disturbance, such as large amounts of
672 flammable biomass altering the severity of a fire. Positive feed-back loops have
673 additive effects, such as grass biomass increasing the probability of a fire, which
674 removes trees, and provides grass with a competitive advantage. (b) Biotic
675 disturbances, such as grazing lawns created by white rhino, modify the quality and
676 quantity of vegetation over time by altering the competitive balance between
677 grazing tolerant and grazing intolerant grass species (described by Refs. [46, 50,
678 51]). (c) Abiotic disturbances, such as fires, alter the short-term abundance and
679 nutritional quality of the grasses available to herbivores by removing senescent
680 vegetation and stimulating nutrient-rich re-growth (described by Refs [56, 61, 99]).
681 Long-term species succession, could reverse this trend.
682
683

684 Figure 6. An ecosystem template of the macroecological processes determining the
685 abundance and distribution of herbivores. (a) Underlying environmental gradients
686 simultaneously affect both the quality and quantity of forage available to herbivores
687 as well as the efficiency of predators at capturing prey. Thus, bottom-up and top-
688 down processes are not independent. For example, (b) lions select areas with
689 denser vegetation, and (c) areas that are closer to rivers for hunting more often than
690 expected, based on the availability of these resources across the landscape.
691 Therefore, the plant community structure (such as % tree cover) and topographic
692 features (such as rivers) contribute to the predation risk for herbivores, while
693 simultaneously influencing the quality and quantity of forage available to them
694 (data from Ref. [66]).
695
696

697 Figure 7. Predictions of the relative importance of predation (solid red line), food quality
698 (dotted green line) and food abundance (dashed blue line) in regulating herbivores
699 of increasing body mass across rainfall and soil fertility gradients, assuming all else
700 is equal (e.g. availability of drinking water, size and isolation of protected area,
701 etc.). We consider herbivores between 10kg to 1000kg, with a major portion of
702 grass in their diet [33]. (a) High rainfall and soil nutrients. Food abundance

703 regulates large herbivores, food quality regulates medium sized herbivores and
704 predation regulates small herbivores. (b) Low rainfall, high soil nutrients. Food
705 quality does not regulate. Food abundance regulates large and medium sized
706 herbivores, predation regulates small ones. (c) High rainfall, low soil nutrients.
707 Food quality regulates all herbivores, with predation acting synergistically at small
708 size, and food abundance at very large size. (d) Low rainfall and soil nutrients.
709 Predation is not regulating, food quality acts at small size and food abundance at
710 medium and large sizes. For more details see main text. If rainfall patterns change
711 due to global warming, herbivore regulation within a savanna ecosystem is
712 predicted to shift from panel a to b, or c to d, or visa versa.

713

714

715

Figure 1

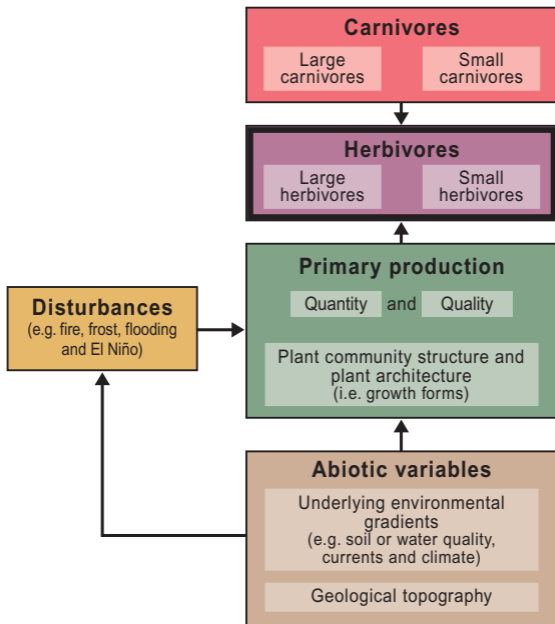
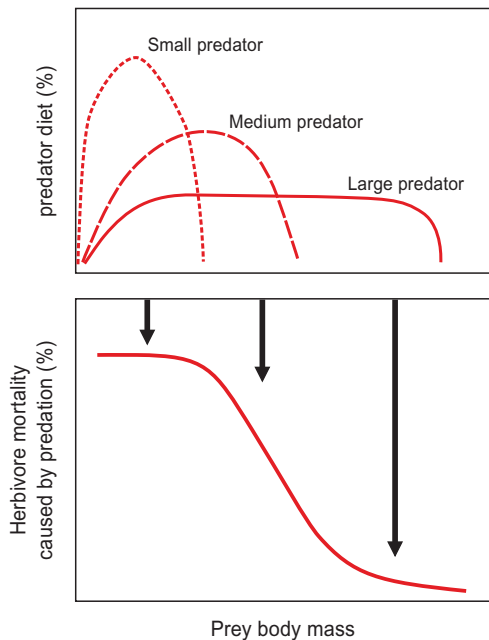


Figure 2

(a) Size-nested predation



(b) Size-partitioned predation

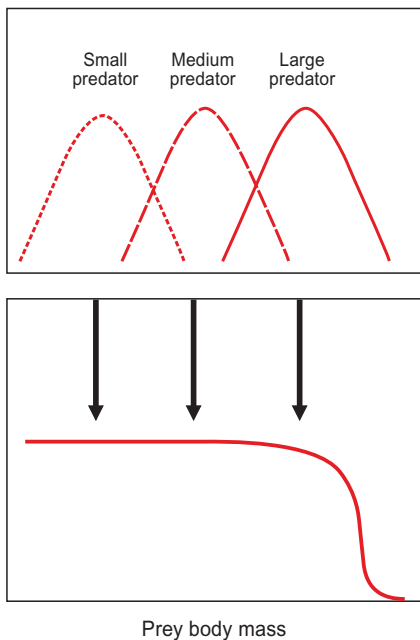
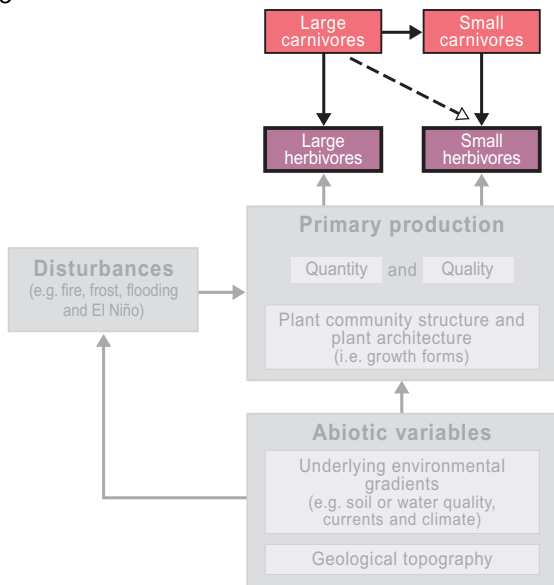


Figure 3

(a)



(b)

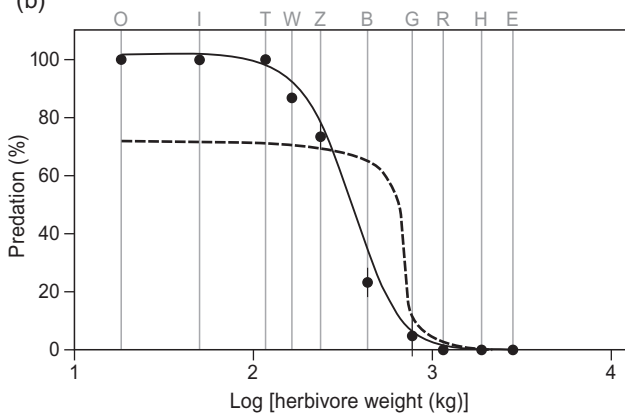
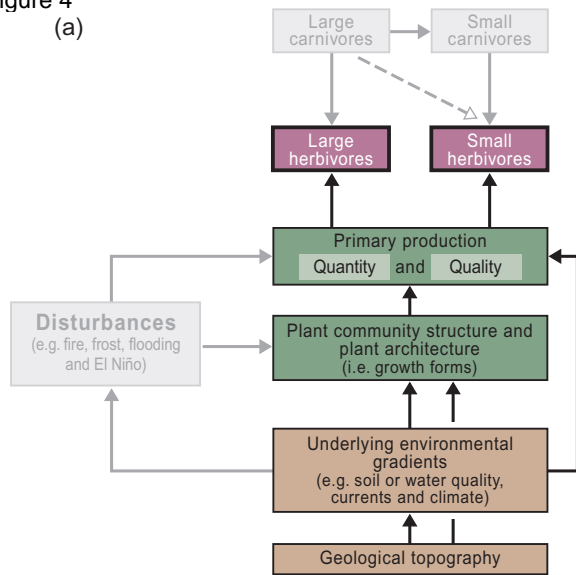
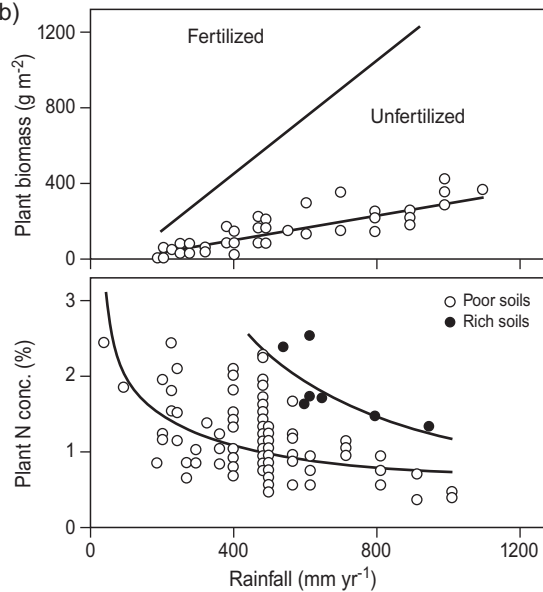


Figure 4

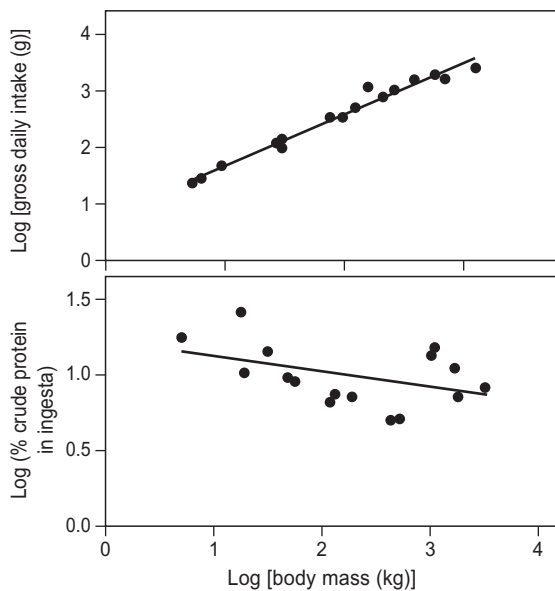
(a)



(b)



(c)



(d)

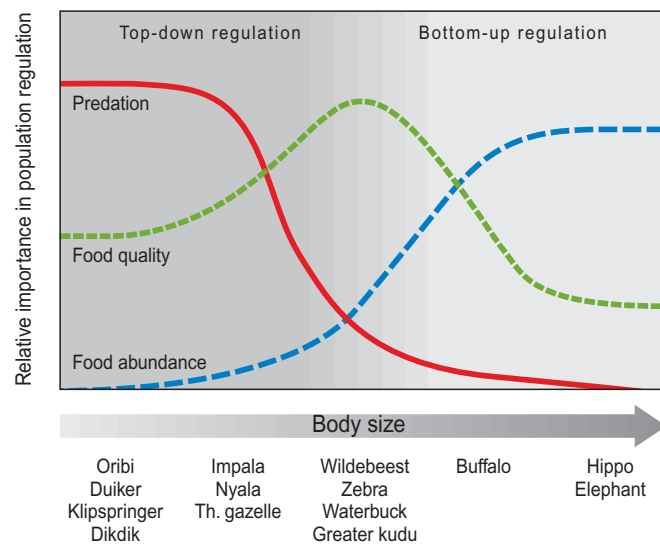
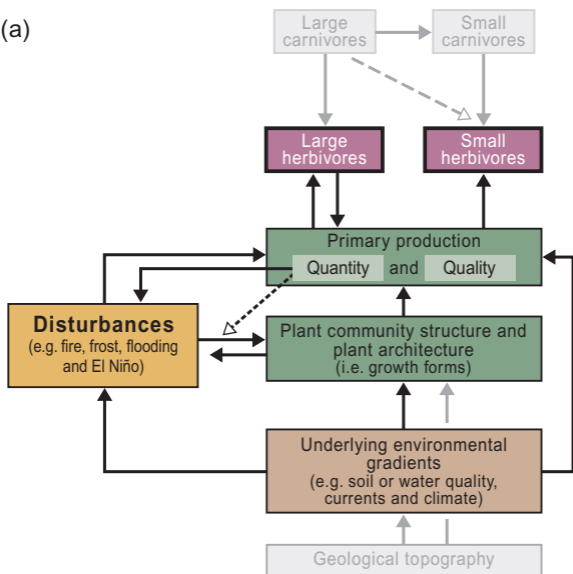
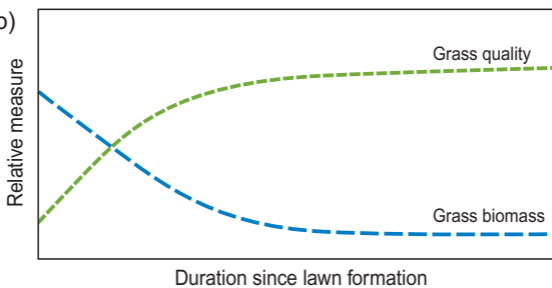


Figure 5

(a)



(b)



(c)

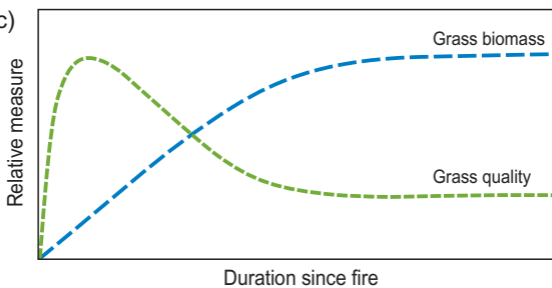


Figure 6

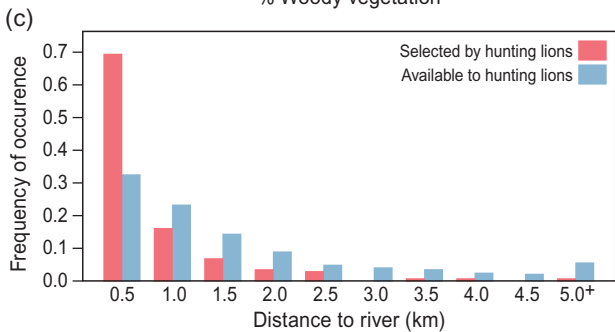
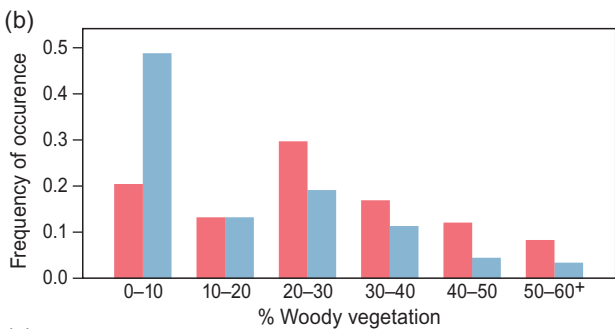
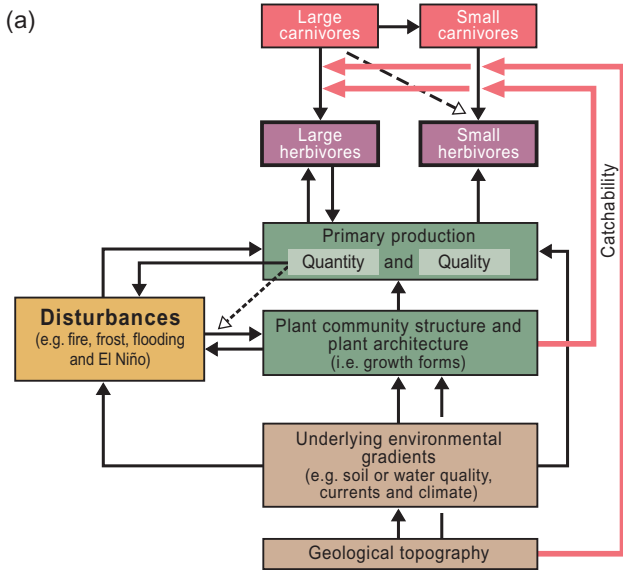


Figure 7

