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2 *Ecography*

3 **Seasonality, niche management, and vertical migration in landscapes**
4 **of relief**

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7

8 **Abstract**

9 Landscapes of vertical relief, such as mountains and continental slopes, intensify ecological and
10 climatological variation within narrow spatial windows. Seasonal vertical migrants exploit this
11 variation during their residence in, and movements between, vertically stratified seasonal
12 ranges. Animals in terrestrial, marine, and even human-ecological systems undergo similar
13 patterns of seasonal vertical movements. The diversity of arenas in which vertical migration
14 evolved lends insight to the factors promoting seasonal use of landscapes of relief. Because
15 animals must contend with both endogenous circannual rhythms and exogenous environmental
16 seasonality, vertical migrants may be sensitive to inconsistent change across stratified seasonal
17 ranges under climate change. To better understand how ongoing and future climatic and
18 environmental changes are likely to impact vertical migrants, we examine vertical migration in
19 the context of niche tracking and niche switching. Whereas niche trackers minimize variation in
20 realized environmental conditions throughout their seasonal movements, niche switchers
21 undergo seasonal transitions in realized niche space. These strategies mediate the relationship
22 between migrants and their changing environment, and can be used to forecast impacts of

23 future change and effectively conserve systems of vertical migration. Niche tracking may be
24 hindered by inconsistent or unpredictable environmental change along a single niche axis
25 across strata, while niche switching may be sensitive to incongruous spatiotemporal change
26 across factors. We suggest that climate change will affect seasonal patterns in vertical
27 environments discontinuously across time, space, and strata, and that vertical migrants are
28 likely to face additional anthropogenic threats that interact with environmental seasonality.
29 Conservation of vertical migrants should prioritize the availability of, and facilitate movement
30 between, stratified seasonal ranges.

31 **Keywords**

32 altitudinal, bathymetric, migration, seasonality, climate change, niche breadth

biotopic space. Axes of positions in a physical environment, such as elevation or depth.
fundamental niche. The range of biotic and abiotic conditions over which an organism can, theoretically, survive and produce viable offspring.
geographic migration. Long-distance seasonal movements, such as latitudinal migration.
landscape. Any environment with a substrate, including terrestrial and marine settings.
niche space. Axes of positions in an n-dimensional hypervolume of conditions that define an organism's suitable environment.
realized niche. The range of biotic and abiotic conditions over which an organism actually survives and produces viable offspring. This is a sub-set of the organism's fundamental niche constrained by realized environmental conditions at a specific location and time and by positive and negative interactions with other organisms.
season. An intra-annual subset of any annual cycle, such as tropical precipitation patterns, Arctic sea ice formation and depletion, or endogenous rhythms in hormone release.
strata. Two or more positions separated by vertical biotopic space, as in the case of movement along topographic or bathymetric slopes.
vertical migration. Short-distance seasonal movements across strata such as altitudinal or bathymetric migrations.

Glossary

33

34 **Introduction**

35 Migration is a taxonomically and geographically widespread adaptation to temporal
36 variation in the environment. Migration is characterized by movements between spatially
37 isolated ranges, on a much greater scale than typical day-to-day movements (Dingle and Drake
38 2007). Seasonal ranges may be separated by hundreds or thousands of kilometers across
39 geography, as is the case in long-distance migrations, or just a few hundred meters across
40 topography or bathymetry, as is the case in vertical migrations. Whereas long-distance migrants
41 face significant energetic demands and risks of mortality during their journeys, vertical migrants
42 minimize costs of travel while still realizing considerable ecoclimatic variation. Amphibians,
43 birds, and crustaceans are but a few of the taxa represented among Earth's vertical migrants
44 (Aguzzi et al. 2013, Boyle 2017, Hsiung et al. 2018). These climbing creatures seasonally
45 traverse mountain sides and marine canyons alike, in pursuit of resources, shelter, and mating
46 opportunities.

47 Mountains cover a quarter of Earth's land surface, and shallow seas comprise over a
48 tenth of the total global seabed area (Costello et al. 2010, Karagulle et al. 2017). While vast
49 distances must be traveled to realize significant changes in ecology or climatology across
50 latitudinal gradients, comparable variation can be experienced by traversing only a few
51 hundred meters up or down a mountainside (Körner 2007, Klinges and Scheffers 2020).
52 Similarly, physical properties of the water column generate abiotic and biotic gradients over
53 small distances in depth, compared to the thousands of kilometers that would be required for
54 the same apparent change across latitude (Sprintall and Cronin 2001). Together, elevation and

55 depth constitute a continuous, 1-dimensional biotopic space which vertical migrants can
56 navigate in order to realize or mitigate seasonal change in their environment (Figure 1).

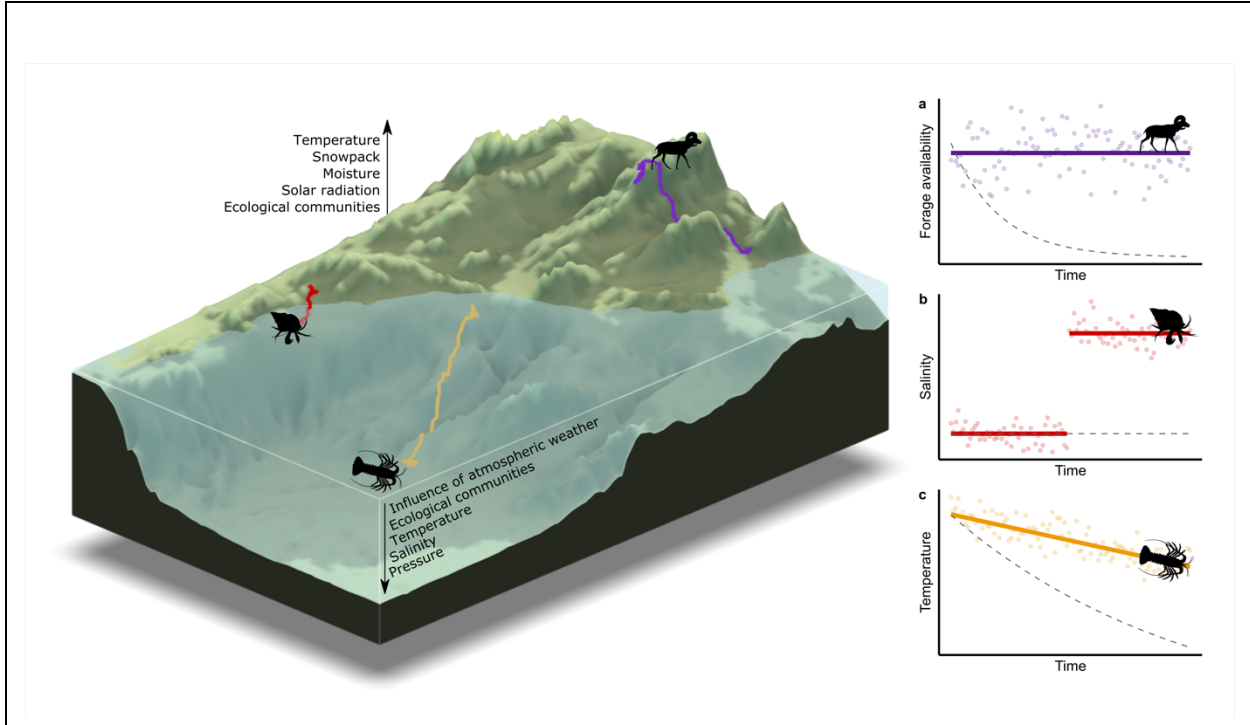


Figure 1. Landscapes of relief introduce multiple axes of environmental variation over short distances, which can be exploited by mobile organisms. In this conceptual illustration, exemplar species are shown at the destination end of their seasonal movement path in both the perspective landscape plot and niche panels (a-c). Environmental factors that vary vertically are indicated along arrows in the perspective plot. In a-c, one axis of niche space is plotted against time, with environmental conditions realized by seasonal migration shown in color, and environmental conditions realized by non-migration shown by a dashed line. Bighorn sheep track springtime plant growth as vegetation green-up progresses uphill (“niche tracking”, a). Soldier hermit crabs face a massive shift in realized salinity when they move from their terrestrial non-reproductive environment to their marine reproductive and developmental environment (“niche switching”, b). Other decapod crustaceans, such as some lobsters, dampen wintertime shifts in temperature by migrating to deeper waters (“dampened niche tracking”, c). Data are for illustrative purposes only; terrain data from ETOPO1 (Amante and Eakins 2009), movement paths generated using directed random steps, and niche space generated using normal distributions with shifting center across time (R version 3.6.1). Animal silhouettes adapted from [phylopic](#); illustrated by Scott Harmon, Ekaterina Kopeykina, and Joanna Wolf; and under public domain and creative commons licenses [CC0 1.0](#) and [CC 3.0](#).

58 The magnitude of seasonal rhythms with which migrants contend often varies across
59 strata. For example, snowpack is deeper and the snow season is longer in temperate alpine
60 regions than in the foothills below. Similarly, deep seafloors are more protected from seasonal
61 storms than shallow coastal waters. Two strategies are available to migrants for coping with
62 spatially structured seasonal environmental variation: they can track spatiotemporal variation
63 along niche axes (“niche tracking,” Figure 1a,c) or undergo seasonal transitions in the niche
64 space they occupy (“niche switching,” Figure 1b) over the course of their migratory journey
65 (Martínez-Meyer et al. 2004, Gómez et al. 2016). Whereas niche trackers maintain constancy or
66 dampen seasonal oscillations in their realized niche (e.g. Somveille et al. 2019, Bay et al. 2021),
67 niche switchers abandon niche space at the turn of the season and exacerbate variation along
68 some niche axes (e.g. Ponti et al. 2020). If seasonal ranges undergo inconsistent environmental
69 change, past strategies of niche tracking and niche switching may fail to accommodate novel
70 conditions.

71 Prevailing patterns of seasonal change across stratified environments may be
72 threatened by changes in the climate regime. Inconsistent change in the timing and magnitude
73 of seasons across strata, as well as long-term trends in bioclimatic and species distributions,
74 modify the pattern of emergence of seasons across landscapes of relief. If phenological shifts in
75 exogenous seasonal factors are inconsistent across elevation or depth (e.g. Inouye et al. 2000),
76 the ability of both niche trackers and niche switchers to cope with forecasted environmental
77 change may be compromised. Inconsistent trends among axes of fundamental niche space may
78 lead to the loss of suitable niche space within a range. Simultaneously, direct human impacts
79 such as land use change and fencing infrastructure limit migrants’ historical access to seasonal

80 ranges. Effective conservation management plans for vertical migrants will account for the
81 spatiotemporal complexities of landscapes of relief.

82 In this review, we ask the following questions: 1) *Why migrate vertically rather than*
83 *geographically?* 2) *In what vertical migratory systems do niche tracking and switching emerge?*
84 and 3) *How will anthropogenic change affect systems of vertical migration?* To answer these
85 questions, we identify how landscapes of relief modify seasonal variation in limiting factors
86 across vertical space, and examine the means by which vertical migrants cope with seasonal
87 variation in the environment. We then explore how climate shapes the progression of seasons
88 across strata, and discuss the mechanisms through which ongoing and future change are likely
89 to affect vertical migrants. We conclude by outlining conservation priorities that will help
90 protect vertical migrants in the face of forecasted change.

91 **Environmental variation in landscapes of relief**

92 Seasonal movements such as migration arise in response to temporal variation in
93 endogenous or exogenous factors. Relative to the scale of seasonal migration, some
94 environmental factors that vary spatially appear to be constant through time, with a similar
95 spatial pattern of variation persisting across years. Migrants that experience significant
96 seasonal endogenous variation (such as breeding or molting) may exploit temporally “static”
97 variation by moving between disparate ranges as the need for - or challenges posed by -
98 different conditions arises. Conversely, other environmental factors vary both spatially and
99 seasonally due to Earth’s axial tilt and position along its revolution around the Sun. Such

100 dynamic exogenous variation may promote migratory movements regardless of the migrant's
101 endogenous state if the relative favorability of seasonal ranges varies across seasons.

102 *Static variation: space*

103 Elevation and depth generate environmental variation over short distances through
104 static influences on abiotic environmental factors. In general, solar radiation increases while air
105 density and temperature decrease with increasing altitude (Körner 2007). Other factors such as
106 precipitation and wind may vary across altitude but the pattern of their variation is locally or
107 regionally idiosyncratic. For example, temperate latitudes tend to experience increasing annual
108 precipitation at higher elevations, while in polar and equatorial latitudes the opposite trend is
109 observed. Within regions, relationships among precipitation and elevation are nonlinear, and
110 vary across seasons (Körner 2007). Water depth similarly imposes stratified abiotic variation:
111 incoming light and solar radiation dramatically decrease through the photic zone, while
112 buffering from atmospheric weather is increased. Conversely, temperature, oxygen
113 concentration, and salinity vary with depth, but their profiles can be nonlinear and locally
114 idiosyncratic (Paulmier and Ruiz-Pino 2009, Shadwick et al. 2015).

115 Stratified variation in abiotic factors drives vertical zonation in ecological communities,
116 composed of species that are simultaneously adapted to a window of conditions and
117 interspecific interactions. Physiological limits of animals and plants determine the vertical range
118 of habitat available to them on both mountains (Janzen 1967) and in oceans (Carney 2005). The
119 process through which relief generates community stratification is perhaps most famously
120 illustrated by the upper limit of tree growth on mountainsides (Körner and Paulsen 2004). In

121 the ocean, primary production is largely limited to the shallowest reaches, where sunlight is
122 able to penetrate and the rate of photosynthesis is greater than that of respiration (Dennison
123 1987). Consequently, a steep decline in forage availability emerges for primary consumers with
124 increasing ocean depth. Similarly, a balance of pressure tolerance, temperature, and oxygen
125 availability dictates the bathymetric range of many marine animals (e.g. Brown and Thatje
126 2011, Brown et al. 2017).

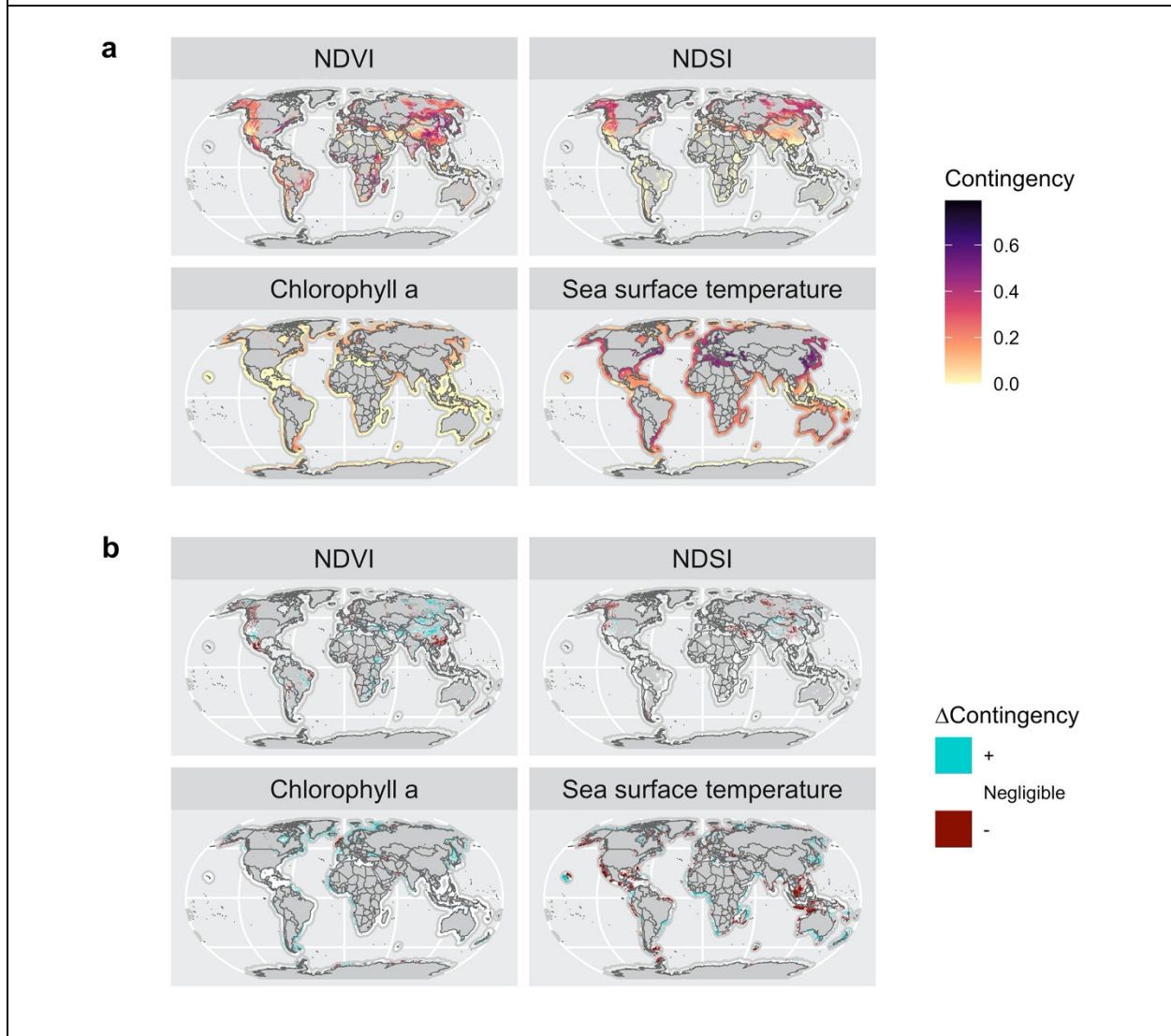
127 *Dynamic variation: space-time*

128 Seasonal variation in abiotic factors drives seasonal variation in ecological communities
129 (Post 2019). Just as accumulated temperature and precipitation regulate plant growth in many
130 terrestrial systems (Cleland et al. 2007), so too do these factors impact the timing, magnitude,
131 and species composition of marine phytoplankton blooms (Thompson et al. 2015).
132 Temperature, precipitation, and primary productivity cycles underlie the adaptive benefits of
133 dormancy and emergence by many terrestrial animals. Seasonal pulsed reproduction by plants
134 and animals alike are linked to cascading effects of abiotic seasons (e.g. madwort flowering:
135 Gómez 1993, caribou and muskox parturition: Kerby and Post 2013, bivalve spawning:
136 Philippart et al. 2014). When seasonal variation follows predictable cycles (e.g. Box 1), migrants
137 can rely on temporal cues such as photoperiod to coordinate movements with anticipated
138 change at the destination.

Box 1. Seasonal variation is not universally predictable in landscapes of relief.

Migration is an adaptive life history strategy when seasonal variation follows a predictable schedule (high “contingency”; Colwell 1974, Shaw and Couzin 2013, Riotte-Lambert and Matthiopoulos 2020). In mountain and coastal regions, contingency of a given factor may vary across space (a). For example, although snow cover is highly contingent in the northern Canadian Rocky Mountains, reliable seasonal cycles are diminished toward the south (NDSI, a). However, seasonal variation may become more or less predictable over time (b) as factors individually respond to cascading effects of climate change. For

example, sea surface temperature variation in numerous areas throughout the Malay Archipelago became less predictable in the 2010's compared to the 2000's, but the same pattern was not observed for chlorophyll concentration. In (a), contingency was calculated following Colwell (1974) for MODIS Terra NDVI and NDSI, and MODIS Aqua L3SMI Chlorophyll a and Sea surface temperature products aggregated to 10km pixel resolution. In (b), contingency of each factor during 2010-2019 was compared against contingency during 2000-2009 to identify change, with $|\Delta\text{Contingency}| < 0.025$ considered "Negligible."



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Because variation in abiotic and biotic factors is neither linear across strata nor uniform

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among factors, temporal environmental variation is spatially structured in landscapes of relief.

142

Landscapes of relief modify the spatial structure of seasonal environmental variation in three

143

forms of "seasonal transformation": phase shifts, phase differences and phase products (Figure

144 2). In phase shifts (Figure 2a), the timing of variation in a factor is shifted with respect to strata,
145 but the magnitude, and baseline condition of variation in that factor, remain unchanged. For
146 example, plant growth may be delayed with respect to elevation due to the recession of the
147 snowline and variation in accumulated temperature along an elevational gradient. In phase
148 differences (Figure 2b), the baseline condition of a factor changes with respect to strata, but
149 the timing and magnitude of variation in that factor remain unchanged. For example, the timing
150 and magnitude of seasonal temperature variation may be consistent across a mountainside,
151 while high elevations experience predictably colder temperatures than lower elevations. In
152 phase products (Figure 2c), the magnitude of variation in a factor is adjusted with respect to
153 strata, while the timing and baseline condition of the factor remain unchanged. For example,
154 disturbance from seasonal atmospheric storms is greater in shallow waters than in protected
155 waters deep below. Finally, landscapes of relief may introduce a combination of these phase
156 modifications (Figure 2d), where two or more forms of spatiotemporal variation emerge. For
157 example, the duration and magnitude of seasonal snow cover may be much greater in an alpine
158 environment than the foothills below. Similarly, photosynthetic activity in the euphotic zone
159 generates a primary productivity seasonality regime in shallow waters, while primary
160 production in deeper waters may be negligible.

161 *Geography and the relative role of relief*

162 Migration may facilitate the pursuit of favorable conditions, or escape from unfavorable
163 conditions (Winger et al. 2019). Across broad geographic distances, spatiotemporal variation in
164 environmental conditions is generally reliable. For example, movement between a savanna and
165 tropical rainforest ensures directional change in moisture and ecosystem structure (Whittaker

166 1970). This is also the case to a certain extent across topography or bathymetry (e.g. Whittaker
167 and Niering 1968); however, regional variation in climate may predominate in spite of
168 considerable local variation in some factors. For example, persistent snow is common at high
169 elevations in the Arctic, while seasonal snow cover characterizes variation across elevations
170 (Hammond et al. 2018). Vertical migration would thus be insufficient as a means of complete
171 escape from snow in the Arctic; geographical movements are the only available option if that
172 region is to be utilized by migrants that cannot cope with snow. At more temperate and tropical
173 latitudes, however, persistent snow is rare, and seasonal snow at high elevations is not
174 matched at low elevations (Hammond et al. 2018). Altitudinal movements would facilitate a
175 response to snow at these latitudes.

176 Conversely, vertical retreat from unfavorable winter conditions may be adaptive in
177 marine settings at higher latitudes, where sea surface temperatures are most dramatically
178 seasonal in shallow waters (Box 1). This combined phase difference and amplification at high
179 latitudes introduces a strong seasonal signal in the depth gradient in water temperature, of
180 particular importance for ectotherms. Tropical waters face comparatively little seasonal
181 variation in temperature, but strong seasonality in sea surface salinity due to winds and fresh
182 water inputs by precipitation and runoff (Delcroix and Hénin 1991). While vertical migration in
183 these waters may satisfy seasonal endogenous requirements, so too may it allow migrants to
184 cope with temporally dynamic exogenous factors.

185 Terrain structure further complicates general patterns of environmental variation along
186 elevation or depth. Slope and aspect are particularly impactful factors in terrestrial temperate
187 and Arctic settings, where increased direct solar irradiance on slopes facing the equator leads

188 to increased temperature, reduced moisture, and advanced phenology (Jackson 1966, Griffiths
189 et al. 2009, Geroy et al. 2011). In marine settings, topography underlies the mixing and
190 movement of water masses (Huthnance 1995, Shapiro et al. 2003), impacting temperature,
191 deposition of organic matter, availability of oxygen and nutrients, and community diversity
192 (Robertson et al. 2020). Coastal upwelling and its drivers add additional complexities to
193 seasonal variation in environmental factors along depth, but simultaneously introduce cues
194 that reliably predict imminent environmental change (Largier 2020). Upwelling in California
195 (USA), for example, leads to a seasonal influx of nutrients from deeper waters, and
196 consequently increases in production of phytoplankton and ultimately krill. Thus ensures
197 seasonal arrival by geographically migratory blue whales (*Balaenoptera musculus*) when
198 availability of their preferred food source is high there (Croll et al. 2005). Further from the
199 coast, organic carbon deposition reliably follows upwelling events, leading to seasonal influxes
200 of consumable detritus and visitation by mobile benthic species on the deep sea-floor (Smith et
201 al. 2013). Thus, in spite of coarse regional patterns that underlie some unavoidable seasonal
202 environmental, landscapes of relief introduce additional dimensions of local environmental
203 variation along which migratory animals can track existing conditions, or seek out novel
204 environments.

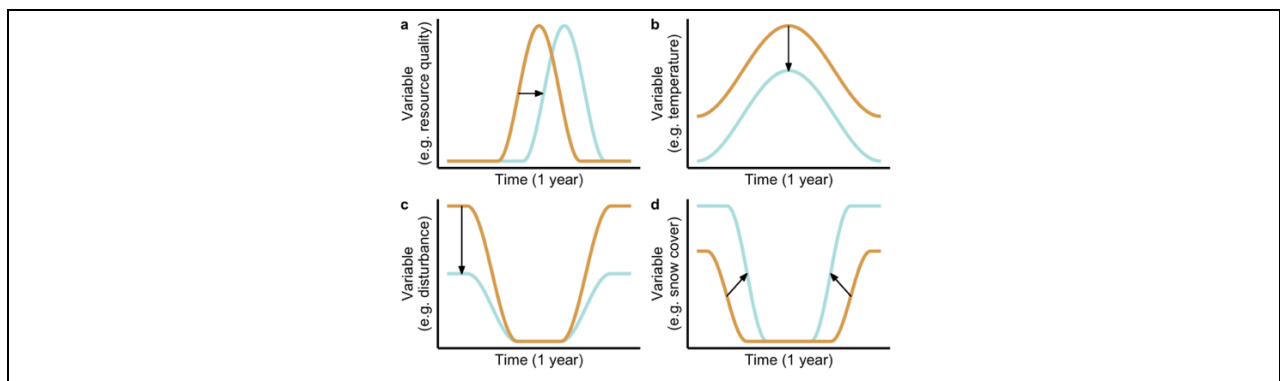


Figure 2. Landscapes of relief shape the spatiotemporal environmental variation through seasonal transformations across strata (tan and blue lines correspond to two environments of different elevation or depth). Phase *shifts* (a) involve a temporal advance or delay in the seasonal cycle without change in the intercept or magnitude of the cycle. Phase *differences* (b) arise when the cycle's intercept changes but without changing the amplitude of the cycle. Phase *products* occur where the same baseline conditions exist across space, but the magnitude of the cycle varies in the form of amplification or attenuation (c). Further spatiotemporal complexities may arise through a combination of these phase changes (d). The primary axes of phase variability are illustrated with black arrows.

205 **Niche management in multi-season space**

206 The environmental variation introduced by landscapes of relief can be exploited without
207 moving across vast distances, as would be required of long-distance migrants. This reduces the
208 endogenous cost of migration, such as energy usage and storage limitations, as well as external
209 pressures such as unfavorable conditions faced during long-distance migrations (Alerstam et al.
210 2003, Boyle 2017). Migrants must navigate a gamut of exogenous seasons while simultaneously
211 facing seasonal variation in endogenous factors such as hormone release, body condition, and
212 reproductive status. Animals may therefore use seasonal migration as a tactic to navigate
213 biotopic space while managing their use of niche space. For a given niche axis, migrants may
214 maintain access to consistent conditions (“niche tracking”) or seek out completely different
215 conditions during a subset of year (“niche switching”, Martínez-Meyer et al. 2004). Whereas
216 niche tracking involves the maintenance of niche space during migration, niche switching
217 involves a fundamental shift in the realized environmental factors during or as a result of
218 migration. Vertical migrants may employ either or both of these strategies to cope with the
219 numerous seasons of change they face.

220 *Niche tracking*

221 Niche tracking vertical migrants follow constant environmental conditions across strata
222 in landscapes of relief. Because the phenology of seasonal factors such as forage availability and
223 accumulated temperature may vary across strata (e.g. Hopkins 1920), niche trackers can
224 synchronize their vertical movements with spatially predictable change in their environment
225 (Riotte-Lambert and Matthiopoulos 2020). Migrants using this strategy can minimize
226 physiological or behavioral adjustments that would be required for major transitions in realized
227 environmental factors.

228 Many migrating ungulates follow the phase shift in spring plant growth upslope during
229 spring in a form of niche tracking called “surfing the green wave” (Albon and Langvatn 1992,
230 Mysterud et al. 2001, Merkle et al. 2016). Because newly emergent plant material is nutrient-
231 rich and easily digested, these migrants can extend the window of high-quality forage access
232 and increase fat reserves by tracking green-up through space (Middleton et al. 2018). Recent
233 evidence indicates that ungulates are able to more effectively track resource phenology when
234 vegetation growth is rapid and progresses sequentially across the landscape, as is the case in
235 many mountain settings (Aikens et al. 2020). Evidence of resource tracking in landscapes of
236 relief has also been reported in tortoises (Blake et al. 2013), hares (Rehnus and Bollmann 2020),
237 birds (Loiselle and Blake 1991), and human-ecological systems (Beck 1991).

238 Niche tracking vertical migrations are also useful for coping with seasonal variation in
239 temperature. For example, sea surface temperature off the coast of Maine (USA) is highly
240 seasonal, generating an attenuation in temperature variation toward deeper, more insulated

241 water. Seasonal bathymetric movements have been detected through mark-recapture of deep-
242 sea lobsters (*Homarus americanus*) in Maine, where they use shallower waters during the
243 summer before retreating to the continental slope for winter (Cooper and Uzmann 1971).
244 Migratory lobsters were able to maximize their growth rate, in part because they maintained a
245 warm narrow range of temperature across the year, while sympatric lobsters that remained
246 resident in shallow waters year-round experienced colder temperatures that were more
247 variable (Cooper and Uzmann 1971). In the Mediterranean Sea, similar seasonal variation in
248 bathymetric distributions of benthopelagic fish and crustaceans appears to coincide with
249 temperature and salinity variability introduced by seasonal subsurface water currents (Aguzzi et
250 al. 2013).

251 *Niche switching*

252 Niche switching vertical migrants face major shifts in realized niche space during or as a
253 result of their migration. Migrations associated with endogenous schedules, such as dormancy,
254 molting, and reproduction frequently include some element of niche switching. Because
255 seasonality in physiology and behavior may cyclically require different habitat parameters,
256 many niche switching vertical migrants exploit stratified environmental variation that is not
257 necessarily seasonal. By partitioning niche space over the course of the year, niche switchers
258 can adjust habitat use to accommodate seasonal life history requirements.

259 Seasonal dormancy allows organisms to cope with periodically harsh conditions, and is
260 frequently associated with vertical migratory movements. In Costa Rican dry forests, seasonal
261 variation in rainfall generates seasonality in vegetation growth. Skipper butterfly (*Aguna*

262 *asander*) caterpillars rely on leaves of *Bauhinia unguolata*, a shrub that produces leaves only
263 during the rainy season (Assunção et al. 2014). In the absence of *Bauhinia* leaves, the insects
264 migrate uphill to colder strata where they estivate in trees and among rocks. Once the seasonal
265 rains resume, *A. asander* return to the rejuvenated lowland forests to feed and reproduce
266 (Janzen 2004). Interestingly, a sympatric predatory paper wasp (*Polistes variabilis*) undergoes
267 similar vertical movements to reach dry season dormancy habitat (Hunt et al. 1999). Bears and
268 bats also undergo seasonal vertical movements to access hibernacula, which are separated
269 from habitat used during their waking life through amplified snowpack and stratified
270 geomorphology (Grachev 1976, Neubaum et al. 2006).

271 Niche switching may also facilitate reproductive tasks. A perhaps extreme example of
272 this occurs in a transit between terrestrial and marine environments. The ecotone between
273 land and sea is inherently stratified and delineates an impressive array of environmental
274 conditions. Soldier crabs (*Coenobita clypeatus*) of Mona Island (Puerto Rico, USA) undergo
275 seaward migrations during an annual summer spawning event called the *cobada* (Nieves-Rivera
276 and Williams 2003). Adult *C. clypeatus* spend most of their time in terrestrial habitat, but
277 migrate to the ocean where eggs are deposited, larvae develop, and juveniles claim shells.
278 Seaward migrations by land hermit crabs are timed to coincide with seasonal peaks in water
279 temperature, but also attuned to the lunar cycle for favorable tide and lighting conditions (Doi
280 et al. 2016, Nio et al. 2019). Niche switching reproductive migrations are also observed among
281 some birds, when seasonal courtship, foraging, and roosting habitat are vertically stratified
282 (Mussehl 1960, Crawford and Pelren 2001, Zwickel and Bendell 2003), or when reproduction is
283 seasonal and sensitive to stratified variation in predation (Boyle 2008).

284 *Navigating the fundamental niche: the tracking-switching continuum*

285 Across the entirety of a migrant's fundamental niche, both niche tracking and niche
286 switching are no doubt employed over the course of a year. Any environmental condition that
287 varies in space can be tracked or abandoned by a mobile organism, so long as relocation does
288 not require exceeding some physiological constraint (i.e. departing from fundamental niche
289 space). Because migrants face a crash of endogenous and exogenous seasons, they potentially
290 realize inconsistent variation among niche axes while moving across strata. For example,
291 ascending a marine canyon may increase access to forage, but simultaneously lead to changes
292 in light, salinity, and hydrostatic pressure. By navigating physical space, migrants can realize
293 considerable differentiation along one niche axis, but minimize variation along another.
294 Incomplete migratory patterns, such as partial and facultative migrations, lend insight into how
295 animals use vertical movements to track or switch among seasonal niche spaces.

296 Partial migration is common among vertically migratory species, wherein a subset of the
297 population migrates while another does not (e.g. Cooper and Uzmann 1971, Boyle 2017).
298 Whereas migration can facilitate tracking favorable environmental variation, remaining resident
299 in a particular range may limit the extent to which niche tracking is possible (Laube et al. 2015,
300 Gómez et al. 2016). For example, many Sierra Nevada bighorn sheep (*Ovis canadensis sierrae*)
301 in California (USA) move to low-elevation slopes during winter to avoid deep snow and seek
302 foraging opportunities, but they face increased predation pressure compared to those
303 remaining in the barren snowy alpine zone (Spitz et al. 2020). While individuals that remain
304 resident in the alpine zone may reduce interactions with predators, they must cope with
305 increased seasonal variation in temperature, wind speed, snow cover, and forage availability.

306 Other factors favoring partial migration include seasonal intraspecific competition and conflict,
307 intrapopulation niche differentiation, and thermal tolerance (Chapman et al. 2011). Coping
308 with, or evading, these factors, likely underlies how individuals manage niche space: evading
309 conditions that are sought after by conspecifics ensures a different realization of total annual
310 niche space occupied by the individual.

311 Facultative and sub-seasonal movements add flexibility to migrants' seasonal
312 distribution schedule. Niche tracking vertical migrants may be able to entrain their movements
313 with environmental conditions by making multiple vertical movements leading up to and during
314 the migratory season. The close geographic proximity of seasonal ranges afford vertical
315 migrants the opportunity to scout conditions and find refuge from unfavorable weather by
316 revisiting seasonal ranges (Horvath and Sullivan 1988, Hahn et al. 2004, Rice 2008, Boyle et al.
317 2010). Sub-seasonal vertical movements also arise among multiple-breeding animals, such as
318 some birds seeking to track high-quality conditions across reproductive attempts (Brambilla and
319 Rubolini 2009, Ceresa et al. 2020). Such fine-scale adjustments in vertical distribution may
320 enable some migrants to buffer climate change and maintain access to tracked niche space
321 throughout the season (Frey et al. 2016).

322 Tracking and switching lie on opposite ends of a continuum of strategies for managing
323 seasonal niche space. Migrants likely employ both approaches during vertical migration
324 because of the numerous factors that vary across strata in landscapes of relief. However, the
325 interaction between physiology, behavior, and environmental factors governs whether a
326 migrant generally tracks niche space across the year, or switches between seasonal subsets of
327 niche space. For example, ectotherms that face seasonal temperature variation without going

328 into dormancy track vertical zonation in temperature in order to maintain metabolic activity
329 (e.g. Cooper and Uzmann 1971, Crossin et al. 1998). Conversely, species that migrate to
330 hibernacula in order to escape unfavorable conditions undergo a massive shift in metabolism,
331 activity, and preferred habitat (e.g. Hunt et al. 1999, Haroldson et al. 2002, Neubaum et al.
332 2006). Niche switching is similarly dramatic for diadromous and semi-diadromous migrants as
333 they deposit larvae in ontogenic habitat (e.g. Elliott et al. 2007, Amorim et al. 2016). The niche
334 management strategy employed by vertical migrants, and the faculty to use alternative
335 approaches during migration, will likely determine their ability to persist under future climate
336 change.

337 **Climate shapes configuration of seasonal niche space**

338 Climate modifies stratified environmental variation in landscapes of relief by shaping
339 patterns within and relationships among niche axes across biotopic space (Colwell and Rangel
340 2009). Changes in temperature and precipitation, and the seasonality thereof, lead to ecological
341 response across strata and through time. Common ecological responses to warming include
342 advancing phenology due to more rapidly accumulated spring temperature, and range shifts to
343 accommodate physiological constraints (Parmesan 2006). However, the degree of phenological
344 change may vary across strata, as well as the propensity to shift ranges and the availability of
345 novel, suitable habitat. Climate-induced changes in migratory propensity may influence
346 reproductive exchange within partially migratory populations, ultimately impacting gene flow
347 and genetic diversity, potentially driving, or limiting, evolutionary change. Vertical migrants'
348 ability to “keep up” with climate change will depend on modifications to the spatiotemporal

349 arrangement of environmental variation in their historic, current, and potentially future habitat,
350 as well as change in the phenology and distribution of organisms with which migrants interact.

351 *Seasonal niche space*

352 Inconsistent alteration of stratified ranges by climate change will impact the relative
353 timing of transitions between seasons, and consequently the ability of some migrants to track
354 spatial variation along some niche axes (Figure 3, Niche 1). For example, in the European Alps,
355 rapid phenological advance at high elevations has led to more uniform green-up across
356 elevation (Vitasse et al. 2018). Earlier green-up and flowering by plants related to climatic
357 warming were not matched by parturition of roe deer (*Capreolus capreolus*) in the Alps (Rehnus
358 et al. 2020). Roe deer are income breeders and therefore depend on forage availability during
359 the energetically expensive parturition and fawn rearing season. Thus, exacerbation of
360 conditions that result in trophic mismatch can be consequential for offspring production and
361 survival (Kerby and Post 2013). However, due to the elevational gradient in green-up timing, *C.*
362 *capreolus* were able to buffer the expanding mismatch by adjusting their migratory schedule to
363 account for shifting spatiotemporal dynamics of forage quality (Rehnus et al. 2020). The
364 advantages of such a buffer may be stymied by more spatio-temporally uniform plant
365 phenological dynamics. A similar pattern is expected for bark beetles (*Ips typographus*), as the
366 timing of diapause termination and swarming is expected to shift more rapidly at high
367 elevations and on south-facing slopes in the Alps (Jakoby et al. 2019). Adjustments to the
368 historic elevational trend in insect swarming phenology may impact insectivorous altitudinal
369 migrants that live there (e.g. grey wagtails, Klemp 2003). Inconsistent shifts across stratified

370 ranges can also be expected in marine settings, where effects of climate change are not
371 constant between surface waters and deep ocean (Capotondi et al. 2012).

372 Inconsistent climatic change across space also modifies the historic overlap and
373 adjacency of stratified niche spaces between seasons. Disruption of the historic relative timing
374 of seasonal niche availability may impact the seasonal migration schedule of some niche
375 switching vertical migrants (Figure 3, Niches 2 and 3). For example, grizzly bears (*Ursus arctos*
376 *horribilis*) in the greater Yellowstone ecosystem undergo a partial downhill migration after
377 emerging from high-elevation dens in spring (Haroldson et al. 2002). They then follow the
378 elevational progression of snowmelt into high-elevation habitat in the summer, coincident with
379 the summertime aggregations of Army cutworm moths (*Euxoa auxiliaris*) that migrate to the
380 alpine from the Great Plains (Pruess 1967, Servheen 1983, French et al. 1994, White et al.
381 1998). This influx of forage facilitates the hyperphagia required of bears leading up to
382 hibernation (Nelson et al. 1983, French et al. 1994). Throughout the Rocky Mountains, the
383 predictability of seasonal variation in snow cover has changed during recent years (Box 1).
384 Warming winters and reduced snow accumulation are expected in that area for the coming
385 decades (Lapp et al. 2005). If emergence and departure timing of *E. auxiliaris* in the Great Plains
386 becomes decoupled from snowmelt and den emergence by *U. a. horribilis* in the Rocky
387 Mountains, the efficacy of migratory bears' switch from hibernating to foraging may be
388 compromised.

389 *Fundamental niche space*

390 Response to climate change through range shifts is additionally limited for vertical
391 migrants. Abiotic stressors covary with both altitude and depth, imposing novel pressures on
392 animals that undergo distributional shifts in response to increasing temperatures (Spence and
393 Tingley 2020). In mountains, a finite upper biotopic limit (mountain peaks and ridgelines)
394 prohibits indefinite upslope response. The available surface area of mountains is furthermore
395 not constant across strata (Körner 2004, Elsen and Tingley 2015) and presents an added
396 constraint on altitudinal migrants, particularly if density dependent factors like intraspecific
397 competition for resources underlies the elevational distribution of seasonal ranges. In marine
398 systems, range shifts toward deeper water are a common response to warming surface waters
399 (Pinsky et al. 2013). However, physical and chemical factors prevent an indefinite retreat to
400 deeper waters. The twilight zone introduces an absolute maximum depth for visual animals,
401 while the sea surface delimits a minimum depth available to marine obligates. Reduced oxygen
402 and increased hydrostatic pressure in deep water impose limits on metabolism, introducing a
403 barrier that can only be overcome through physiological adaptation (Brown and Thatje 2015).
404 Because hypoxia tolerance is strongly linked to temperature in some depth migrants (Bigford
405 1979, Deutsch et al. 2015), change in ocean temperature and oxygenation, and shifts in the
406 depth of the oxygen minimum zone, may be especially limiting for depth shifts by seasonal
407 vertical migrants in the ocean.

408

409

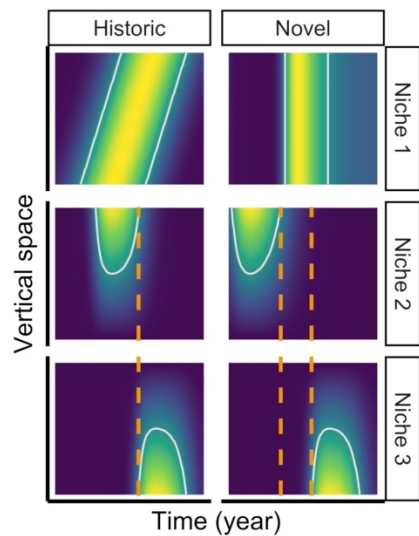


Figure 3. Inconsistent seasonal transformations across biotopic and niche axes precludes effective niche tracking and niche switching. Each panel illustrates the vertical gradient (y-axis) in niche value (color intensity) across the course of a year (x-axis). The left column of panels represents an historic climate regime, while the right column represents novel climatic conditions. Each row represents a different niche axis which may be tracked or differentially prioritized. Grey contour lines indicate an arbitrary threshold value within niche space that migrants seek out. For Niche 1, novel conditions lead to a more synchronous season across biotopic space. A tracking migrant may have historically migrated vertically to follow the spatiotemporal progression of Niche Variable 1, but under novel conditions more synchronous emergence of threshold niche access may reduce incentives to relocate in a coherent vertical pattern. For Niches 2 and 3, an advance in the timing of Niche 2 relative to Niche 3 leads to a temporal window where suitable space is not available along either niche axis under novel conditions (delineated with orange dashed lines). If a niche switcher depends on the historical synchrony of senescence in Niche Variable 2 and emergence of Niche Variable 3 during its migration, inconsistent change in the onset and termination of these factors will disrupt the pattern of environmental variation under which migration was adaptive.

410

411 **Conserving systems of vertical migration**

412 While the emergence of suitable niche space across vertical biotopic space is prone to
 413 adjustments by climate, migrants' access to and availability of seasonal ranges face additional
 414 threats. Direct processes such as infrastructure development and harvest may inhibit access by

415 animals to seasonal ranges, influence population dynamics, and drive microevolutionary change
416 in vertical migrants (Pecl et al. 2009, Pigeon et al. 2016, McInturff et al. 2020). Because
417 migrants have more expansive annual ranges than nonmigratory animals, more opportunities
418 arise for anthropogenic processes to disrupt their life history. Management of systems of
419 vertical migration should account for the multifaceted spatiotemporal requirements of
420 migrants.

421 Seasonal landscapes that actively face direct human-induced change such as
422 deforestation and destructive harvest are likely to be immediate concerns for vertical migrants.
423 For example, due to rapid and accelerating global deforestation (Hansen et al. 2013, Kim et al.
424 2015), habitat fragmentation is of particular concern for migrants that specialize on forests for
425 at least part of their life history. The Eastern Arc Mountains of Tanzania house a large
426 proportion of forest-dependent birds, but also face a high degree of deforestation (Buchanan et
427 al. 2011). Such loss of forest habitat is concerning for both the numerous endemic tree species
428 (Hall et al. 2009) and avian altitudinal migrants that live there (Burgess and Mlingwa 2000). One
429 such species is the Banded green sunbird (*Anthreptes rubritorques*), a forest-dependent
430 altitudinal migrant endemic to the Eastern Arc Mountains, which is now listed as a Threatened
431 species due to habitat loss through deforestation (BirdLife International 2017). Conservation
432 efforts must prioritize areas of rapidly vanishing habitat, especially where these intersect with
433 forecasts of climate change impacts on habitat availability, and particularly in instances where
434 highly endemic species face extinction (Post and Brodie 2015).

435 Parallel to habitat destruction, movement barriers limit the realized availability of
436 seasonal ranges to migratory animals. Human infrastructure, including fencing, roads, and

437 bridges, interrupt directed movement, thereby preventing migrants from accessing seasonal
438 ranges. Because anthropogenic delineations between land designations rarely fall along
439 ecological boundaries, fences bar passage across otherwise continuous niche space. Semi-
440 permeable solutions, such as wildlife-friendly fencing, increase connectivity between seasonal
441 ranges, while still fulfilling the original intention of the infrastructure (Paige 2008, Sawyer et al.
442 2013). Barrier permeability may be similarly important in marine settings. For example, many
443 crabs undergo seasonal movements between shallow estuarine and deeper ocean waters (e.g.
444 Bigford 1979). The construction of bridges connecting capes at the mouth of estuaries may
445 present temporary or permanent barriers to crab migration: project scheduling and design
446 permeability must be considerations if the conservation of marine-estuarine bathymetric
447 migrants is a priority.

448 Effectively managing for the future will rely on focused consideration of the seasonal
449 role humans play in the systems vertical migrants occupy, and the sensitivity of niche space to a
450 changing climate. One such example lies in a climate change hotspot, the East Australian
451 Current (EAC), which has faced changes in the strength and seasonality of its flow and poleward
452 shifts in the distribution of species that inhabit it (Johnson et al. 2011, Champion et al. 2018).
453 The southern rock lobster (*Jasus edwardsii*) is an economically important species that
454 undergoes seasonal bathymetric movements to shallow water for molting (MacDiarmid 1991).
455 *Jasus* sp. lobsters face seasonal fishery pressure in Tasmania, which is also highest in shallow
456 waters near the coast. Lobster recruitment is expected to fall in the coming decades as sea
457 temperature rises (Pecl et al. 2009), but they face the added threat of invading sea urchins that
458 arrived in response to overfishing (Johnson et al. 2011). Accounting for spatiotemporal

459 variation in lobster abundance, and determining whether effects of climate and harvest are
460 additive or synergistic, are crucial to the success of fishery management there.

461 While systems of vertical transhumant pastoralism may not fit neatly under some
462 contemporary definitions of “migration,” their seasonal structure of landscape use presents an
463 opportunity to forecast dynamics and examine policy through analogy: transhumant
464 pastoralism is the practice of leading livestock along seasonal routes to track pasture quality,
465 and mountain ranges across the world have served as systems for vertical pastoralism
466 throughout human history. A primary concern of shepherds is the condition of grazing and
467 wintering lands; decisions about grazing timelines dictate forage quality for livestock, and
468 frequently incite conflict (Beck 1991). Shifts in predictable forage phenology may be difficult to
469 match by pastoralists due to the often-rigid transit schedules imposed on them through
470 political avenues. Nepalese herders have reported earlier snowmelt and advanced plant
471 phenology in the Himalaya, where they undergo seasonal vertical movements to maintain yaks
472 and other ungulates (Aryal et al. 2016). Added to shifting social and cultural landscapes,
473 environmental change amplifies the expected decline of vertical transhumant pastoralism in
474 the coming decades (Hock et al. 2019). Protection of the routes that connect seasonal ranges,
475 and flexibility by officials to account for changes in ecosystem dynamics, will be central to the
476 conservation of this imperiled lifestyle.

477 **Conclusions**

478 The evolution of analogous patterns across ecosystems lends insight to both basic and
479 applied research questions (Burkepile et al. 2020). One such pattern is the seasonal movement

480 between vertically separated habitats: vertical migration. Seasonal vertical migrations have
481 evolved not only within marine and terrestrial habitats, but in some cases they involve transit
482 between these inherently stratified ecosystems. Vertical migrants can be found throughout the
483 animal kingdom, including among arthropods, birds, and even human-ecological systems. The
484 heightened environmental variation generated by landscapes of relief allows animals to exploit
485 or mediate the seasonality of their environment in order to facilitate growth and reproduction.
486 However, the historic pattern of variation may be sensitive to adjustment by climate and
487 accessibility by other anthropogenic impacts. Migrants' capacity to manage seasonal niche
488 space by navigating biotopic space may underlie their ability to persist under climate and land
489 use change, but well-planned conservation action and policy can help ensure the longevity of
490 vertical migrants.

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496

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