

1 **Highlights**

2 Population declines have been greater among migratory species because of their vulnerability  
3 to climate change and human pressure. Growing concerns for migratory animals necessitate  
4 new assessments of the outcome of environmental changes for species that rely on long-  
5 distance migration to the North.

6 A growing body of evidence suggests that North temperate and Arctic animals are currently  
7 experiencing lower food supply and availability, higher pathogen and parasite pressure, as  
8 well as increased predation rates, compared with previous decades.

9 We hypothesize that the natural advantages of migration to northern latitudes are being  
10 eroded. Understanding the underlying mechanisms of ecological impacts will allow better  
11 forecasting and mitigation, as well as insights into consequences for population dynamics of  
12 migratory animals.

13 **Animal Migration to Northern Latitudes: Environmental Changes and Increasing**  
14 **Threats**

15 Vojtěch Kubelka<sup>1-4\*</sup>, Brett K. Sandercock<sup>5</sup>, Tamás Székely<sup>2,3</sup> and Robert P. Freckleton<sup>1\*</sup>

16 <sup>1</sup>Department of Animal and Plant Sciences, University of Sheffield, Alfred Denny Building,  
17 Western Bank, Sheffield S10 2TN, UK

18 <sup>2</sup>Milner Centre for Evolution, University of Bath, Claverton Down, Bath BA2 7AY, UK.

19 <sup>3</sup>Department of Evolutionary Zoology and Human Biology, Faculty of Science, University of  
20 Debrecen, Egyetem tér 1, Debrecen, Hungary

21 <sup>4</sup>Department of Biodiversity Research, Global Change Research Institute, Czech Academy of  
22 Sciences, Bělidla 986/4a, Brno, 603 00, Czech Republic

23 <sup>5</sup>Department of Terrestrial Ecology, Norwegian Institute for Nature Research, Høgskoleringen  
24 9, Trondheim, 7485, Norway

25 \*Corresponding authors: Kubelka, V. (kubelkav@gmail.com), Freckleton, R. P.  
26 (r.freckleton@sheffield.ac.uk)

27 **Key words:** climate change, food supply, nest predation, parasites, population dynamics,  
28 trophic interactions

29 **Abstract**

30 Every year, many wild animals undertake long-distance migration to breed in the North,  
31 taking advantage of seasonally high pulses in food supply, fewer parasites and lower  
32 predation pressure in comparison with equatorial latitudes. Growing evidence suggests that  
33 climate change-induced phenological mismatches have reduced food availability.  
34 Furthermore, novel pathogens and parasites are spreading Northwards, and nest or offspring  
35 predation has increased at many Arctic and North temperate locations. Altered trophic  
36 interactions have decreased the reproductive success and survival of migratory animals.  
37 Reduced advantages for long-distance migration have potentially serious consequences for  
38 community structure and ecosystem function. Changes in the benefits of migration need to be  
39 integrated into projections of population and ecosystem dynamics and targeted by innovative  
40 conservation actions.

## 41 **The Northward Migration**

42 Each year numerous wild animals migrate to Arctic or North temperate **breeding grounds**  
43 (see Glossary). Migratory taxa include mammals, insects and, notably, many birds (e.g. [1,2],  
44 Figure 1, Figure S1), but the evolution and ecological implications of migratory behaviour are  
45 still not fully understood [1,3,4]. For such extensive, costly and dangerous behaviour to  
46 evolve, the benefits must be considerable [5,6]. Breeding at higher latitudes is assumed to  
47 have several advantages that outweigh the physiological costs and mortality risk connected  
48 with **migration**. Major benefits include (i) seasonal pulses of food supplies and long days for  
49 foraging [1,7], (ii) low prevalence of pathogens and parasites [8,9], and (iii) reduced predation  
50 pressure in comparison with southern areas [10,11].

51 Currently, many populations of terrestrial animals undertaking **long-distance migration** are  
52 threatened, declining in numbers, and performing worse than their resident counterparts [12–  
53 15]. Changes along **migratory routes**, especially habitat loss or deterioration, or disturbance  
54 and persecution on non-breeding areas, **stopover and staging sites** have already received  
55 considerable attention and are now well-recognized drivers of population declines through  
56 reductions in adult survival [12,16,17]. But here, we suggest that the recent declines are also  
57 driven, in part, by deterioration in the ecological quality of North temperate and Arctic  
58 breeding grounds. Breeding grounds have received less attention because they are often  
59 remote or inaccessible areas with less anthropogenic activity. Here, we highlight recently-  
60 documented impacts on ecological outcomes following long-distance migration in a range of  
61 terrestrial animals (Figures 2–3, Table 1).

62 In forecasting the impacts of future threats the spatial and temporal extents of possible  
63 changes are critical factors (Figure 3). For example, disturbances such as storms or temporary  
64 mismatch with food resources may be individually relatively brief **acute stressors (pulses)**

65 leading to short impacts and fast recovery (Figure 3A). In contrast, long-term **chronic**  
66 **stressors (presses)**, such as changes in ambient CO<sub>2</sub> and temperature are slower directional  
67 changes occurring over periods of decades ([18], Figure 3B). Acute and chronic stressors  
68 create a continuum: for instance, repeated pulses due to increasing frequency of storms or  
69 predation pressure can also act as presses. Moreover, once chronic stressors exceed a  
70 threshold, the state of an entire system can change, leading to population extinction or regime  
71 shift (Figure 3B). Similarly, characterizing the spatial extent of impacts is important (Figure  
72 3C), because the scale over which changes occur determines not just the extent of impacts but  
73 also the feasibility of conservation efforts.

74 Here, we review and discuss factors that appear to be driving reduced profitability of  
75 northward breeding grounds for different groups of migrating animals (Figures 2–3, Table 1).  
76 We argue that various on-going environmental changes are resulting in large-scale chronic  
77 stressors degrading habitats, emphasizing needs for targeted conservation actions.

## 78 **Evidence for Reduced Benefits of Animal Migration**

### 79 ***Food Supply***

80 Recent climate change has affected food supplies and their seasonal availability in Northern  
81 latitudes [19,20]. To successfully exploit short-term peaks of food abundance, reproduction of  
82 higher trophic levels needs to be synchronized with relevant periods of plant phenology or  
83 insect emergence [21]. However, mismatches are well recognized, and have been documented  
84 for various taxa [19,22]. The phenomenon is termed **trophic/phenological mismatch** and can  
85 occur at various scales (Figure 2, Table 1).

86 Phenological mismatch alone may not necessarily lead to detrimental fitness consequences. If  
87 minimum food requirements are still met, young can grow and survive, even if food

88 abundance is not at its peak [23,24]. Global warming usually advances the plant-growing  
89 season and the peak of arthropod abundance (Table 1), but the number of days with an  
90 adequate food supply may be unchanged [7,25]. Local and temporary mismatches represent  
91 acute stressors, whereas continuous imbalance between reproduction timing and food  
92 availability may result in long-term and large-scale chronic stressors affecting the entire  
93 ecosystem (Figure 3).

94 The number and intensity of summer storms is increasing [26]. Arthropod availability for  
95 insectivorous migrants is reduced during inclement weather events [21,25], and may reduce  
96 the availability of food during critical windows during the offspring rearing period, increasing  
97 the probabilities of abandonment and mortality of young [21]. In contrast, for herbivorous  
98 migrants such as caribou and geese, climate warming may increase available plant biomass  
99 during the brood-rearing period in summer [17,27]. However, it is possible that increasing  
100 plant biomass may not be sufficient to negate the consequences of phenological mismatch  
101 [27]. Predatory long-distance migrants, such as skuas (*Stercorarius* spp.) can be negatively  
102 affected by long-term chronic stressors at breeding grounds due to ongoing shortages in the  
103 abundance of prey species, as well as increased competition with other predators [28].

#### 104 ***Pathogens and Parasites***

105 The prevalence of disease agents was historically low in boreal and Arctic regions, because  
106 the pathogens are typically unable to complete their life cycle in harsh environments, as well  
107 as because of a limited number of suitable vectors [8,9]. There is now evidence that a variety  
108 of pathogens, parasites and their vectors have shown poleward shifts in their distributions.  
109 Emerging diseases are consistent with earlier projections based on impacts of global warming  
110 [29,30] and novel pathogens represent an increasing threat for wildlife at high latitudes [31–  
111 33] (Figures 2–3, Table 1). Examples include acute stressors, such as avian cholera outbreaks

112 in the Canadian Arctic leading to mortality of common eiders *Somateria mollissima* [34] or an  
113 extensive and rapid mass mortality event at calving grounds of saiga antelopes *Saiga tatarica*  
114 in Central Kazakhstan caused by hemorrhagic septicemia following unusually high  
115 temperatures and humidity in the region [35,36]. New pathogens and parasites invading  
116 northern latitudes including helminths of mammals and birds represent chronic stressors for  
117 migrating animals (Figures 2–3, Table 1). Migratory birds or bats are also important because  
118 they transport non-native pathogens. For example, the *Plasmodium* causing avian malaria is  
119 now able to complete the transmission cycle in the Arctic [37].

## 120 **Predation**

121 In general, predation pressure appears to be increasing for Arctic and North temperate wildlife  
122 (Figure 2, Table 1). The impacts range from acute to chronic stressors, both at various spatial  
123 scales (Figure 3), creating novel predator-prey interactions [38]. Historically, predation  
124 pressure has been thought to decline from the tropics towards the poles [10,11,39]. However,  
125 climatically-induced rapid alterations in boreal and the Arctic ecosystems, including changes  
126 in predator numbers and predator guild composition, have been predicted to induce increased  
127 predation pressure on breeding birds [21,40,41] and such changes have been recently detected  
128 (Figure 2, Table 1).

129 In some Arctic regions, climate change-induced damping of the population cycles and  
130 abundance of lemmings and voles [42–44] may have influenced the behaviour of predators  
131 that consume nests and chicks of birds as alternative prey [21,45]. For example, loss of  
132 lemming cycles may be a factor limiting breeding productivity and population size of brant  
133 geese *Branta bernicla* [46]. Elevated nest predation rates have also been reported in temperate  
134 Europe [45,47], together with changes in cyclicity and lower abundances of voles [48,49], and  
135 similar ecological mechanisms may occur in both North temperate and Arctic regions [50].

136 However, interactions of predators with rodents and bird nests as an alternative prey can be  
137 highly dynamic and some studies have found only weak relationships between rodent  
138 abundance and population trends of other animals [51].

139 The behaviour of predators may have changed, altering their distributions and increasing their  
140 impacts during the breeding season. For example, changing sea ice dynamics (a chronic  
141 stressor) have led to stranding of polar bears *Ursus maritimus* in coastal areas across the  
142 Arctic where they can now prey on breeding colonies of geese, ducks, gulls and auks [52,53].  
143 The geographic ranges of some generalist predators have also increased Northward, including  
144 the red fox *Vulpes vulpes* [19,21]. Generalist avian predators such gulls and corvids have  
145 increased their numbers and spread, supported by human activities [54]. Moreover, sites with  
146 increased primary productivity (greening, another chronic stressor for High Arctic wildlife) in  
147 a warming Arctic experienced higher predation rates on artificial nests, suggesting an elevated  
148 risk of nest predation in tundra ecosystems [55]. Increased predation pressure may not be  
149 restricted to migratory species or birds, for example Arctic ground squirrels *Urocitellus*  
150 *parryii* inhabiting Canadian boreal forest were nearly extirpated by increased predation [56].

## 151 **Responses of Migrating Animals to Changing Environmental Conditions**

152 Migratory animals can modify their behaviour, life-history or physiology through phenotypic  
153 plasticity or adaptation to account for changes in the profitability of migration. Such changes  
154 may ameliorate the consequences of the aforementioned disruptions to migration benefits,  
155 particularly chronic stressors. Migratory schedules commonly change, specifically earlier  
156 arrival on the breeding grounds, matching phenological advances [19,20,22,57]. However,  
157 species can only adjust phenology within certain limits. For example, in migratory birds,  
158 flexibility is limited because of the need to build up energetic reserves prior to migration  
159 [19,23]. Similarly, caribou can change timing of migration [58], however when tracking



160 frozen water bodies which enhance landscape connectivity, rising temperatures and thinner  
161 ice impede caribou migration [17]. Migratory animals can change wintering grounds as well,  
162 tracking the altered environmental conditions, where older individuals with more experience  
163 can be critical for developing new migration behaviours, as shown for cranes [59]. Migratory  
164 routes or the timing of migration may also change in response to predation pressure [60].  
165 Several species of migratory birds have shown long-term reductions in wing length, possibly  
166 as an adaptation to improve aerial agility in response to increased predation pressure  
167 following recovery of falcon populations [61].

168 Due to the rapid pace and complexity of recent changes at breeding grounds (Table 1, Figure  
169 3), migratory animals may not have developed suitable responses to all novel threats. Current  
170 migratory behaviour might become less advantageous or even maladaptive (Figure S2). In the  
171 worst-case scenario, breeding locations in the Arctic tundra, as well as in boreal and North  
172 temperate zones, could now represent **ecological traps** with lower profitability than  
173 alternative locations [62,63] or **degraded environments** with no better alternatives in the  
174 surrounding landscape for migrating animals (Figure S3).

### 175 **Implications for Population Dynamics**

176 Migratory behaviour presumably evolved as an adaptive strategy to maximize fitness as a  
177 trade-off between reproductive success and adult mortality in seasonal environments [5,6].  
178 However, conditions on breeding grounds are changing, with potential to reduce reproductive  
179 success, lowering the profitability of migration. Negative consequences are likely for  
180 individual fitness, population trends and recovery from perturbations (Figure 3, Table 1). The  
181 breeding ranges of migratory birds may track the distributions of predators and alternative  
182 prey species [4,64], suggesting that some species might avoid breeding grounds with high nest  
183 and chick predation if more suitable alternatives are available. Similar predator avoidance at

184 larger scales would be more difficult for ground-travelling mammals, for example caribou  
185 which prefer remote calving grounds with lower predation pressure [17]. Conversely, regions  
186 with high predation pressure, or sites with lack of alternative prey, could experience local  
187 extinction of migratory animals [4,64] and thereby initiate significant alterations in predator-  
188 prey interactions, changing trophic food webs with cascading effects for the ecosystem.

189 Responses vary across different species or populations. For example, invertivorous shorebirds  
190 in North America are generally declining whereas herbivorous geese are increasing [65,66].  
191 In these cases, the complex drivers are quite different: multiple factors are responsible for  
192 declines in shorebirds whereas increases in numbers of geese are mainly driven by changes in  
193 agriculture practices and improvements in winter habitat quality [67], regardless of possible  
194 changes in the profitability of migration.

195 Many temperate species have shifted ranges northward following global warming. However,  
196 High Arctic migratory animals usually cannot extend their northward distribution owing to  
197 the geographic barrier of the Arctic Ocean [40]. Migratory species often have inflexible life-  
198 history strategies and hence are particularly at risk from any environmental changes [19,40].  
199 The life cycle of migratory animals typically consists of distinct stages in different  
200 environments, each with different limiting factors. Serious disruption at any stage of the life-  
201 cycle could lead to a steep decline of the whole population [16,68]. Traditional harvest of  
202 migratory ungulates and birds remains important for many groups of indigenous people  
203 [17,69], consequently reduced populations of migratory animals could also socioeconomically  
204 impact human communities.

## 205 **Implications for Conservation**

206 It will be challenging to directly mitigate the large-scale impacts of climate change for  
207 migratory species that are dependent on multiple environments distributed across several  
208 regions of the globe. Conservation efforts at all spatial and temporal scales are important,  
209 starting from local direct nest protection to regional habitat management. However, large-  
210 scale conservation projects are essential to secure future for migratory animals (Figure S3),  
211 including the development or expansion of international networks of protected areas [70,71].

212 Climate change is most pronounced in Arctic regions where suitable habitats are changing  
213 rapidly [19,40]. Environmental protections in the Arctic require cooperation among  
214 governments and indigenous peoples [69] in the face of economic incentives for development  
215 of mining and oil drilling and to manage exploitation of natural resources and wildlife.  
216 Growing Arctic settlements need proper waste disposal systems to avoid supplemental  
217 feeding of generalist predators [54]. New trading routes, currently opening across the more  
218 ice-free Arctic Ocean [72], need to be carefully planned and well-controlled from the start to  
219 minimize their impact on the Arctic ecosystems, especially when many Arctic regions still  
220 remain largely unprotected [73]. Issues encountered by migrating animals at the North  
221 temperate breeding grounds are more complex, involving climate change, habitat degradation  
222 due to intensification of agriculture and forestry production or urban areas spreading, direct  
223 persecution, disturbance or increased predation pressure [45,47,74], requiring coordinated  
224 conservation activities across large scales (Figure S3). Moreover, it is important to maintain  
225 landscape connectivity by reducing obstacles in traffic corridors such as telecommunication  
226 towers, wind turbines, and powerlines for migrating bats and birds or gas pumpjacks and  
227 fence-lines for migrating ungulates [74].

228 We suggest that Arctic and North temperate breeding grounds need substantial conservation  
229 attention, in addition to well-recognized problems at stopover sites and wintering areas of

230 migratory species [12,16,74]. Targeting only one or two stages may be not enough [45],  
231 therefore integrated conservation measures based on international cooperation will be  
232 essential to cover the entire life-cycle, and the critical areas used by migratory animals  
233 throughout the year.

#### 234 **Concluding Remarks and Future Perspectives**

235 Ecological conditions at Arctic and North temperate breeding grounds may be deteriorating  
236 for many migrating animals owing to recent changes in the availability of food resources,  
237 prevalence of pathogens and parasites, and increased predation rates. Animals adapted for  
238 migration to the Arctic and North temperate regions may face dual threats from low breeding  
239 productivity at breeding grounds and deteriorating adult survival during their migratory  
240 movements [45]. This double jeopardy for long-distance migrants could further intensify the  
241 negative population trends of migratory species.

242 When mitigating impacts of chronic stressors at larger scales, it is vital to rank habitats by  
243 their quality, although challenging to be able to recognize: (i) habitats supporting sufficient  
244 reproductive output and likely maintaining good source populations of migratory species; (ii)  
245 disturbed habitats, which are still advantageous for migrating animals, only less than they  
246 used to be; (iii) ecological traps or degraded environments with negative consequences for  
247 reproductive output and subsequent population trends (see Figures S2–S3 for details). The  
248 distinction will be essential for effective targeting of conservation measures, mitigating the  
249 impacts of current human pressure and climate change induced pulse or press stressing events.  
250 More extensive and well-connected networks of protected areas, building on previous efforts  
251 such as Ramsar wetlands or Natura 2000 sites across breeding, migratory and wintering areas,  
252 as well as population-specific protective measures, will be essential (see Outstanding  
253 Questions). Last, recent developments in tracking technologies facilitating effective tracking

254 of complete journeys and life cycles of individuals, represent a breakthrough for studies of  
255 migratory connectivity and population dynamics.

256 Author contributions

257 All authors conceived the study; V.K. compiled the data with help from B.K.S. and R.P.F.

258 V.K. wrote the manuscript with inputs from all co-authors.

259 Acknowledgements

260 We thank Andrea E. A. Stephens and three anonymous reviewers for constructive comments

261 and suggestions. V.K. was supported by the Ministry of Education, Youth and Sports of the

262 Czech Republic (CZ.02.2.69/0.0/0.0/19\_074/0014459). V.K. and T.S. were supported by

263 ÉLVONAL-KKP 126949 of the Hungarian government. B.K.S. was supported by Norwegian

264 Institute for Nature Research. T.S was funded by the Royal Society (Wolfson Merit Award

265 WM170050, APEX APX\R1\191045).

266 Supplemental information

267 Supplemental information associated with this article can be found at doi:XXXXXXX'

268 **References**

- 269 1 Lack, D. (1968) Bird migration and natural selection. *Oikos* 19, 1–9
- 270 2 Fryxell, J.M. *et al.* (1988) Why are migratory ungulates so abundant? *Am. Nat.* 131,  
271 781–798
- 272 3 Newton, I. (2008) *The Migration Ecology of Birds*, Academic Press.
- 273 4 Gilg, O. and Yoccoz, N.G. (2010) Explaining Bird Migration. *Science* 327, 276–277
- 274 5 Stearns, S.C. (1976) Life-history tactics: A review of the ideas. *Q. Rev. Biol.* 51, 3–47
- 275 6 Alerstam, T. *et al.* (2003) Long-distance migration: evolution and determinants. *Oikos*  
276 103, 247–260
- 277 7 Tulp, I. and Schekkerman, H. (2008) Has prey availability for arctic birds advanced  
278 with climate change? Hindcasting the abundance of tundra arthropods using weather  
279 and seasonal variation. *Arctic* 61, 48–60
- 280 8 Laird, M. (1961) A lack of avian and mammalian Haematozoa in the Antarctic and  
281 Canadian Arctic. *Can. J. Zool.* 39, 209–213
- 282 9 Piersma, T. (1997) Do global patterns of habitat use and migration strategies co-evolve  
283 with relative investments in immunocompetence due to spatial variation in parasite  
284 pressure? *Oikos* 80, 623–631
- 285 10 Pienkowski, M.W. (1984) Breeding biology and population dynamics of Ringed  
286 plovers *Charadrius hiaticula* in Britain and Greenland: nest- predation as a possible  
287 factor limiting distribution and timing of breeding. *J. Zool.* 202, 83–114
- 288 11 McKinnon, L. *et al.* (2010) Lower predation risk for migratory birds at high latitudes.  
289 *Science* 327, 326–327
- 290 12 Bairlein, F. (2016) Migratory birds under threat. *Science* 354, 10–12

- 291 13 Billerman, S.M. *et al.*, eds. (2021) *Birds of the World*, Cornell Laboratory of  
292 Ornithology (<https://birdsoftheworld.org/>)
- 293 14 Koleček, J. *et al.* (2021) Global population trends in shorebirds: migratory behaviour  
294 makes species at risk. *Sci. Nat.* 108, 9(1–8)
- 295 15 Rushing, C.S. *et al.* (2020) Migratory behavior and winter geography drive differential  
296 range shifts of eastern birds in response to recent climate change. *Proc. Nat. Acad. Sci.*  
297 117, 12897–12903
- 298 16 Studds, C.E. *et al.* (2017) Rapid population decline in migratory shorebirds relying on  
299 Yellow Sea tidal mudflats as stopover sites. *Nat. Commun.* 8: 14895, 1–7
- 300 17 Mallory, C.D. and Boyce, M.S. (2018) Observed and predicted effects of climate  
301 change on Arctic caribou and reindeer. *Environ. Rev.* 26, 13–25
- 302 18 Jentsch, A. and White, P. (2019) A theory of pulse dynamics and disturbance in  
303 ecology. *Ecology* 100, e02734 (1-15)
- 304 19 Gilg, O. *et al.* (2012) Climate change and the ecology and evolution of Arctic  
305 vertebrates. *Ann. N. Y. Acad. Sci.* 1249, 166–190
- 306 20 Parmesan, C. (2006) Ecological and Evolutionary Responses to Recent Climate  
307 Change. *Annu. Rev. Ecol. Evol. Syst.* 37, 637–669
- 308 21 Møltofte, H. *et al.* (2007) *Effects of climate variation on the breeding ecology of Arctic*  
309 *shorebirds – Meddelelser om Grønland Bioscience 59*, Danish Polar Center
- 310 22 Scheffers, B.R. *et al.* (2016) The broad footprint of climate change from genes to  
311 biomes to people. *Science* 354, 719 (aaf7671)
- 312 23 Saalfeld, S.T. *et al.* (2019) Phenological mismatch in Arctic-breeding shorebirds:  
313 Impact of snowmelt and unpredictable weather conditions on food availability and  
314 chick growth. *Ecol. Evol.* 9, 6693–6707

- 315 24 Kwon, E. *et al.* (2019) Geographic variation in the intensity of warming and  
316 phenological mismatch between Arctic shorebirds and invertebrates. *Ecol. Monogr.*  
317 89, e01383
- 318 25 Shaftel, R. *et al.* (2021) Predictors of invertebrate biomass and rate of advancement of  
319 invertebrate phenology across eight sites in the North American Arctic. *Polar Biol.*  
320 DOI: 10.1007/s00300-020-02781-5 (<https://www.springer.com/journal/300>)
- 321 26 Sepp, M. and Jaagus, J. (2011) Changes in the activity and tracks of Arctic cyclones.  
322 *Clim. Change* 105, 577–595
- 323 27 Lameris, T.K. *et al.* (2018) Arctic geese tune migration to a warming climate but still  
324 suffer from a phenological mismatch. *Curr. Biol.* 28, 2467–2473
- 325 28 Perkins, A. *et al.* (2018) Combined bottom-up and top-down pressures drive  
326 catastrophic population declines of Arctic skuas in Scotland. *J. Anim. Ecol.* 87, 1573–  
327 1586
- 328 29 Harvell, C.D. *et al.* (2002) Climate warming and disease risks for terrestrial and marine  
329 biota. *Science* 296, 2158–2162
- 330 30 Altizer, S. *et al.* (2013) Climate change and infectious diseases: from evidence to a  
331 predictive framework. *Science* 341, 514–519
- 332 31 Colella, J.P. *et al.* (2020) Conservation genomics in a changing Arctic. *Trends Ecol.*  
333 *Evol.* 35, 149–162
- 334 32 Davidson, R. *et al.* (2011) Arctic parasitology: why should we care? *Trends Parasitol.*  
335 27, 238–244
- 336 33 Galaktionov, K. V (2017) Patterns and processes influencing helminth parasites of  
337 Arctic coastal communities during climate change. *J. Helminthol.* 91, 387–408
- 338 34 Iverson, S.A. *et al.* (2016) Avian Cholera emergence in Arctic-nesting northern



339 Common Eiders: using community-based, participatory surveillance to delineate  
340 disease outbreak patterns and predict transmission risk. *Ecol. Soc.* 21, 12 (1–24)

341 35 Kock, R.A. *et al.* (2018) Saigas on the brink: Multidisciplinary analysis of the factors  
342 influencing mass mortality events. *Sci. Adv.* 4, eaao2314(1-10)

343 36 Fereidouni, S. *et al.* (2019) Mass die-off of saiga antelopes, Kazakhstan, 2015. *Emerg.*  
344 *Infect. Dis.* 25, 1169–1176

345 37 Loiseau, C. *et al.* (2012) First evidence and predictions of *Plasmodium* transmission in  
346 Alaskan bird populations. *PLoS ONE* 7, e44729 (1-5)

347 38 Guiden, P.W. *et al.* (2019) Predator – prey interactions in the Anthropocene:  
348 Reconciling multiple aspects of novelty. *Trends Ecol. Evol.* 34, 616–627

349 39 Skutch, A.F. (1949) Do tropical birds rear as many young as they can nourish? *Ibis* 91,  
350 430–455

351 40 Wauchope, H.S. *et al.* (2017) Rapid climate-driven loss of breeding habitat for Arctic  
352 migratory birds. *Global Change Biol.* 23, 1085–1094

353 41 Zöckler, C. *et al.* (2003) Wader populations are declining – how will we elucidate the  
354 reasons? *Wader Study Group Bull.* 100, 202–211

355 42 Ims, R.A. *et al.* (2008) Collapsing population cycles. *Trends Ecol. Evol.* 23, 79–86

356 43 Kausrud, K.L. *et al.* (2008) Linking climate change to lemming cycles. *Nature* 456,  
357 93–98

358 44 Aharon-Rotman, Y. *et al.* (2014) Loss of periodicity in breeding success of waders  
359 links to changes in lemming cycles in Arctic ecosystems. *Oikos* 124, 861–870

360 45 Kubelka, V. *et al.* (2018) Global pattern of nest predation is disrupted by climate  
361 change in shorebirds. *Science* 362, 680–683

- 362 46 Nolet, B.A. *et al.* (2013) Faltering lemming cycles reduce productivity and population  
363 size of a migratory Arctic goose species. *J. Anim. Ecol.* 82, 804–813
- 364 47 Roodbergen, M. *et al.* (2012) Revealing the contributions of reproduction and survival  
365 to the Europe-wide decline in meadow birds: review and meta-analysis. *J. Ornithol.*  
366 153, 53–74
- 367 48 Cornulier, T. *et al.* (2013) Europe-wide dampening of population cycles in keystone  
368 herbivores. *Science* 340, 63–66
- 369 49 Bierman, S.M. *et al.* (2006) Changes over time in the spatiotemporal dynamics of  
370 cyclic populations of field voles (*Microtus agrestis* L.). *Ame. Nat.* 167, 583–590
- 371 50 Kubelka, V. *et al.* (2019) Pilot study on nest predation in shorebirds breeding at the  
372 Caspian steppe lakes. *Wader Study* 126, 142–150
- 373 51 Doyle, S. *et al.* (2020) Anthropogenic impacts on the demographics of Arctic-breeding  
374 birds. *Polar Biol.* 43, 1903–1945
- 375 52 Prop, J. *et al.* (2015) Climate change and the increasing impact of polar bears on bird  
376 populations. *Front. Ecol. Evol.* 3, 33 (1–12)
- 377 53 Iverson, S.A. *et al.* (2014) Longer ice-free seasons increase the risk of nest depredation  
378 by polar bears for colonial breeding birds in the Canadian Arctic. *Proc. Royal Soc. B*  
379 281, 20133128 (1–9)
- 380 54 Weiser, E.L. and Powell, A.N. (2010) Does garbage in the diet improve reproductive  
381 output of Glaucous Gull? *Condor* 112, 530–538
- 382 55 Ims, R.A. *et al.* (2019) Arctic greening and bird nest predation risk across tundra  
383 ecotones. *Nat. Clim. Change* 9, 607–610
- 384 56 Donker, S.A. and Krebs, C.J. (2012) Evidence for source–sink dynamics in a regional  
385 population of arctic ground squirrels (*Urocitellus parryii plesius*). *Wild. Res.* 39, 163–

- 386 170
- 387 57 Gill, J.A. *et al.* (2014) Why is timing of bird migration advancing when individuals are  
388 not? *Proc. Royal Soc. B* 281, 20132161 (1–6)
- 389 58 Kauffman, M.J. *et al.* (2021) Mapping out a future for ungulate migrations. *Science*  
390 372, 566–569
- 391 59 Teitelbaum, C.S. *et al.* (2016) Experience drives innovation of new migration patterns  
392 of whooping cranes in response to global change. *Nat. Com.* 7, 12793(1–7)
- 393 60 Lank, D.B. *et al.* (2003) Effects of predation danger on migration strategies of  
394 sandpipers. *Oikos* 103, 303–319
- 395 61 Lank, D.B. *et al.* (2017) Long-term continental changes in wing length, but not bill  
396 length, of a long-distance migratory shorebird. *Ecol. Evol.* 7, 3243–3256
- 397 62 Schlaepfer, M.A. *et al.* (2002) Ecological and evolutionary traps. *Trends Ecol. Evol.*  
398 17, 474–480
- 399 63 Székely, T. (1992) Reproduction of Kentish Plover *Charadrius alexandrinus* in  
400 grasslands and fish-ponds: the habitat mal-assessment hypothesis. *Aquila* 99, 59–67
- 401 64 Léandri-Breton, D.J. and Bêty, J. (2020) Vulnerability to predation may affect species  
402 distribution: plovers with broader arctic breeding range nest in safer habitat. *Sci. Rep.*  
403 10, 5032 (1–8)
- 404 65 Munro, B.Y.M. (2017) What’s killing the world’s shorebirds? *Nature* 541, 16–20
- 405 66 Smith, P.A. *et al.* (2020) Status and trends of tundra birds across the circumpolar  
406 Arctic. *Ambio* 49, 732–748
- 407 67 Abraham, K.F. *et al.* (2005) The dynamics of landscape change and snow geese in mid-  
408 continent North America. *Global Change Biol.* 11, 841–855

- 409 68 Moores, N. *et al.* (2008) *The 2006-2008 Saemangeum shorebird monitoring program*  
410 *report*, Birds Korea.
- 411 69 Barry, T. *et al.* (2020) How does the Arctic Council support conservation of Arctic  
412 biodiversity? *Sustainability* 12, 5042 (1–18)
- 413 70 Amano, T. *et al.* (2018) Successful conservation of global waterbird populations  
414 depends on effective governance. *Nature* 553, 199–202
- 415 71 Davidson, S.C. *et al.* (2020) Ecological insights from three decades of animal  
416 movement tracking across a changing Arctic. *Science* 370, 712–715
- 417 72 Smith, L.C. and Stephenson, S.R. (2013) New Trans-Arctic shipping routes navigable  
418 by midcentury. *Proc. Nat. Acad. Sci.* 110, E1191–E1195
- 419 73 Saura, S. *et al.* (2017) Protected areas in the world’s ecoregions: How well connected  
420 are they? *Ecol. Indic.* 76, 144–158
- 421 74 Wilcove, D.S. and Wikelski, M. (2008) Going, going, gone: is animal migration  
422 disappearing? *PLoS Biol.* 6, e188
- 423 75 Kutz, S.J. *et al.* (2013) Invasion, establishment, and range expansion of two parasitic  
424 nematodes in the Canadian Arctic. *Global Change Biol.* 19, 3254–3262
- 425 76 Both, C. *et al.* (2010) Avian population consequences of climate change are most  
426 severe for long-distance migrants in seasonal habitats. *Proc. Royal Soc. B* 277, 1259–  
427 1266
- 428 77 van Gils, J.A. *et al.* (2016) Body shrinkage due to Arctic warming reduces red knot  
429 fitness in tropical wintering range. *Science* 352, 819–821
- 430 78 Fossøy, F. *et al.* (2014) Reproductive success is strongly related to local and regional  
431 climate in the Arctic snow bunting (*Plectrophenax nivalis*). *Polar Biol.* 38, 393–400

- 432 79 Hallworth, M.T. *et al.* (2018) Tracking dragons: stable isotopes reveal the annual cycle  
433 of a long-distance migratory insect. *Biol. Lett.* 14, 20180741(1–5)
- 434 80 Altizer, S. *et al.* (2015) Do healthy Monarchs migrate farther? Tracking natal origins of  
435 parasitized vs. uninfected Monarch butterflies overwintering in Mexico. *PLoS ONE* 10,  
436 e0141371 (1-14)
- 437 81 Huebschman, J.J. *et al.* (2019) Detection of *Pseudogymnoascus destructans* during  
438 Summer on Wisconsin Bats. *J. Wild. Dis.* 55, 673–677
- 439 82 Stefanescu, C.O. *et al.* (2012) Parasitism and migration in southern Palearctic  
440 populations of the painted lady butterfly, *Vanessa cardui* (Lepidoptera: Nymphalidae).  
441 *Eur. J. Entomol.* 109, 85–94
- 442 83 Morgan, E.R. *et al.* (2007) Parasite transmission in a migratory multiple host system.  
443 *Ecol. Modell.* 200, 511–520
- 444 84 Nyholm, N.E. (2019) Return rates of nest box breeding Pied Flycatchers *Ficedula*  
445 *hypoleuca* to their breeding site in subalpine birch forest in Swedish Lapland, during  
446 1965-2018. *Ornis Svecica* 29, 53–68
- 447 85 Madsen, J. *et al.* (2019) A gloomy future for light-bellied brent geese in Tusenøyane,  
448 Svalbard, under a changing predator regime. *Polar Res.* 38, 3393 (1–6)
- 449 86 Eldøy, I.H. (2019) Variation in predation and nest survival rates in the Arctic breeding  
450 Common Eider (*Somateria mollissima*). Master thesis. Norwegian University of  
451 Science and Technology ([https://ntnuopen.ntnu.no/ntnu-](https://ntnuopen.ntnu.no/ntnu-xmlui/handle/11250/2618100?locale-attribute=en)  
452 [xmlui/handle/11250/2618100?locale-attribute=en](https://ntnuopen.ntnu.no/ntnu-xmlui/handle/11250/2618100?locale-attribute=en))
- 453 87 Flemming, S.A. *et al.* (2019) Spatio-temporal responses of predators to hyperabundant  
454 geese affect risk of predation for sympatric-nesting species. *PLoS ONE* 14(8),  
455 e0221727 (1-16)

456 **Outstanding Questions**

457 Which levels of predation, pathogen prevalence or reduced food supply (acute or chronic  
458 stressors) at breeding grounds represent ecological traps, or degraded environments, with  
459 declining populations? Which sites remain advantageous for migration but are less profitable  
460 than before? Which limiting factors are most influential in driving global population trends?  
461 Are these effects independent or do they have synergistic interactions? Are there common  
462 patterns within a community, or are the ecological drivers different for each species or  
463 population?

464 Which species are best able to cope with novel conditions at breeding grounds and which are  
465 more likely to be vulnerable? How do life-history traits and social behaviour influence the  
466 species adaptability to novel environmental changes? How are the challenged advantages for  
467 the long-distance animal migration in the North relevant to: (i) short-distance or partial  
468 migrants, nomadic and resident species; (ii) other geographical regions such as mammals in  
469 sub-Saharan Africa; (iii) to non-terrestrial taxa such as fish or cetaceans?

470 Is there currently more or less intense competition among migrating animals in the Northern  
471 latitudes in comparison with earlier decades? Given the declining numbers of many migrating  
472 animals nowadays, density-dependent competition could be reduced if resources are  
473 unchanged. However, sources and habitat carrying capacity have probably changed, and  
474 interspecific competition with new species spreading poleward could offset any additional  
475 advantages for long-distance migration to the North.

476 Developing effective conservation strategies for populations of migratory animals will be a  
477 crucial task for coming decades. But what are the most efficient protective measures for  
478 migratory species with complex life histories? Inevitably, there will be need for prioritisation,

479 and we need to understand well the main drivers of global population trends in migratory  
480 species, and we need to apply correctly large-scale conservation measures as well as species-  
481 specific rescue action plans.



483

484 **Figure 1. Examples of Migratory Terrestrial Animals with Recently Reduced Long-**

485 **distance Migration Benefits.** Clockwise from top left: Arctic tern *Sterna paradisaea*, chicks

486 and eggs of American golden plover *Pluvialis dominica*, semipalmated plover *Charadrius*

487 *semipalmatus* in distraction display, Arctic skua *Stercorarius parasiticus* incubating a clutch,

488 saiga antelope *Saiga tatarica* family (photo by Navinder Singh), monarch butterfly *Danaus*

489 *plexippus*, resting caribou *Rangifer tarandus* (photo by Robert McCaw), flying common green

490 darner *Anax junius* (photo by Peter Chen, Wikimedia Commons) and hunting Eastern red bat

491 *Lasiurus borealis* (centre, photo by Michael Durham). All other photos by Vojtěch Kubelka.

492 See also Online Supplemental Information Figure S1. Note that the pool of long-distance

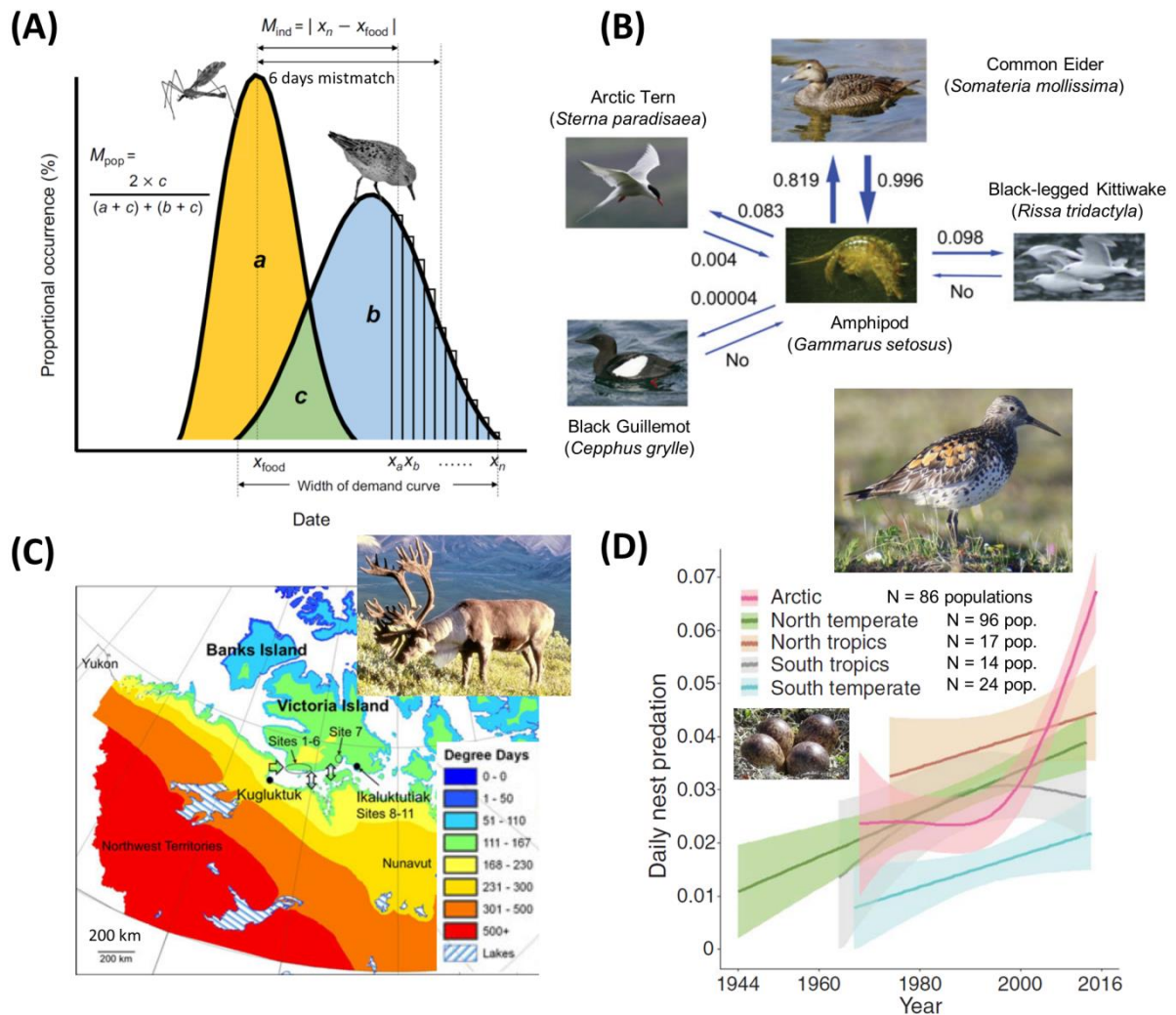
493 terrestrial migrants travelling more than 1,000 km to North temperate and the Arctic breeding

494 grounds includes numerous birds (800+ long-distance migrants, mostly insectivorous or

495 herbivorous and some predatory species), some well-studied species of insects (ca. 12+ spp.,

496 dragonflies, butterflies and moths) and a few species of mammals (ca. 5+ spp., bats, caribou





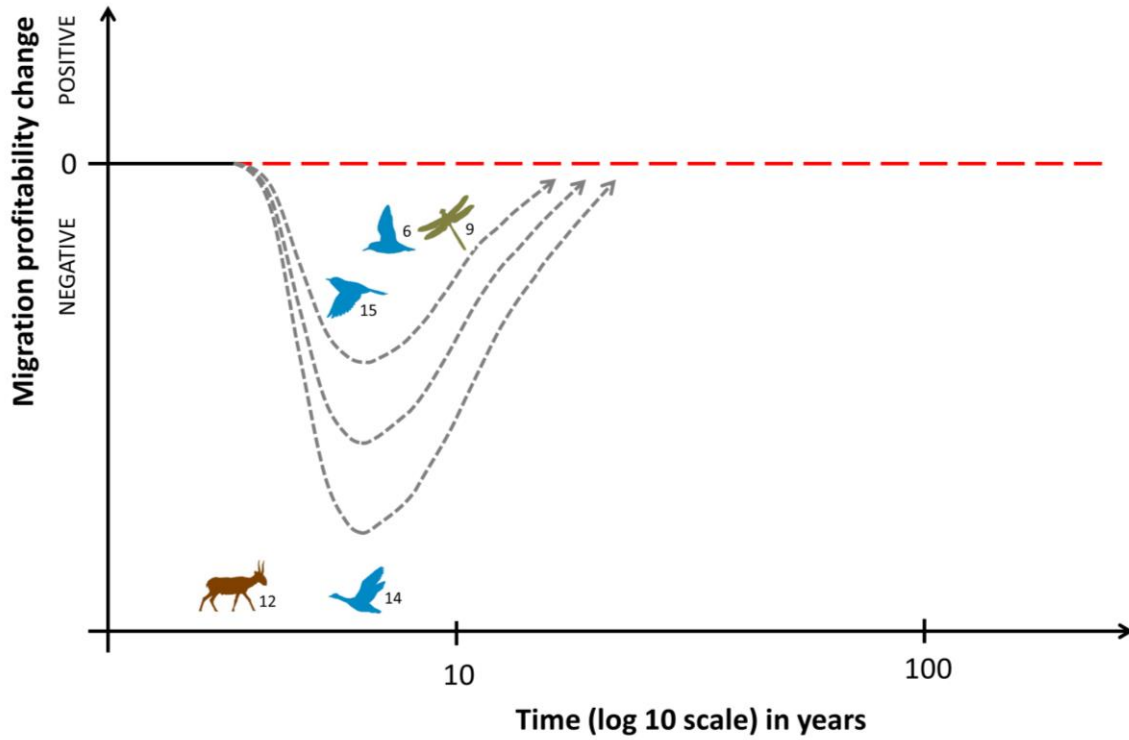
498

499 **Figure 2. Novel Emerging Threats for Migratory Animals at Northern Latitudes.**

500 Selected examples from Table 1. (A) **Food supply and trophic mismatch.** Mean values (+6  
 501 days on average from a range -20 to +22 days range) of individual-nest-level mismatch  
 502 between food (insect) abundance peak and chick food demand peak for six shorebird species  
 503 across the North American Arctic. Modified from Kwon *et al.* (2019) [24]. (B) **Novel host-  
 504 parasite interactions.** Circulation and transmission rates of the acanthocephalan  
 505 *Polymorphus phippsi* and the main host, common eider *Somateria mollissima*. The host-  
 506 parasite network has recently expanded and the parasite has colonised new avian host species  
 507 of seabirds, starting with low infection rates. Modified from Galaktionov (2017) [33].

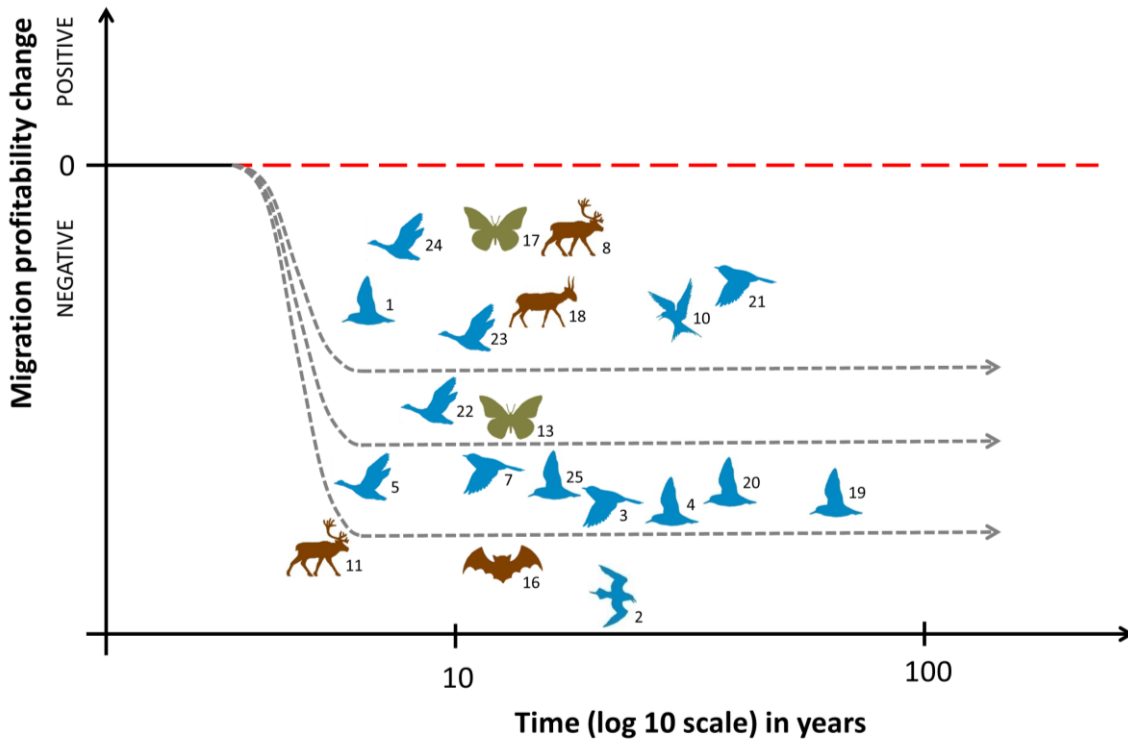
508 **Geographic expansion of parasites.** Continuous degree-day surface map showing  
509 accumulation of degree-days for the development of protostrongylid lungworm  
510 *Umingakstrongylus pallikuukensis* from first larvae (L1) to infective third (L3) stage. From  
511 2000 to 2006, development from L1 to L3 (167 degree-days accumulated) could occur in a  
512 single summer on Southwestern Victoria Island whereas previously conditions were  
513 unsuitable. Protostrongylids parasitize caribou *Rangifer tarandus* (depicted, photo by Dean  
514 Biggins, Wikimedia Commons). Modified from Kutz *et al.* (2013) [75]. (D) **Temporal**  
515 **increase in nest predation.** Nest predation rates for 237 populations of 111 shorebird species  
516 worldwide, divided according to five latitudinal areas. Generalized additive model fits with  
517 95% confidence intervals. Adult and eggs of great knot *Calidris tenuirostris* are depicted  
518 (photos by Vojtěch Kubelka). Modified from Kubelka *et al.* (2018) [45].

**(A) ACUTE STRESSORS (PULSES)**



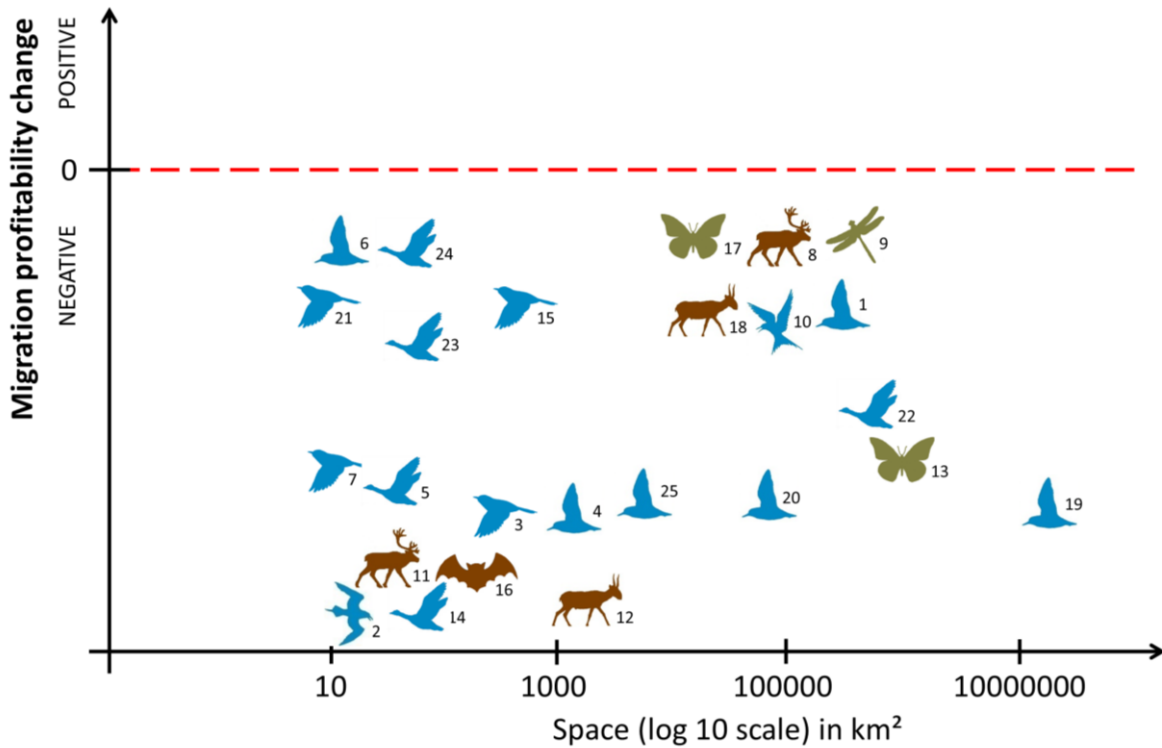
519

**(B) CHRONIC STRESSORS (PRESSES)**



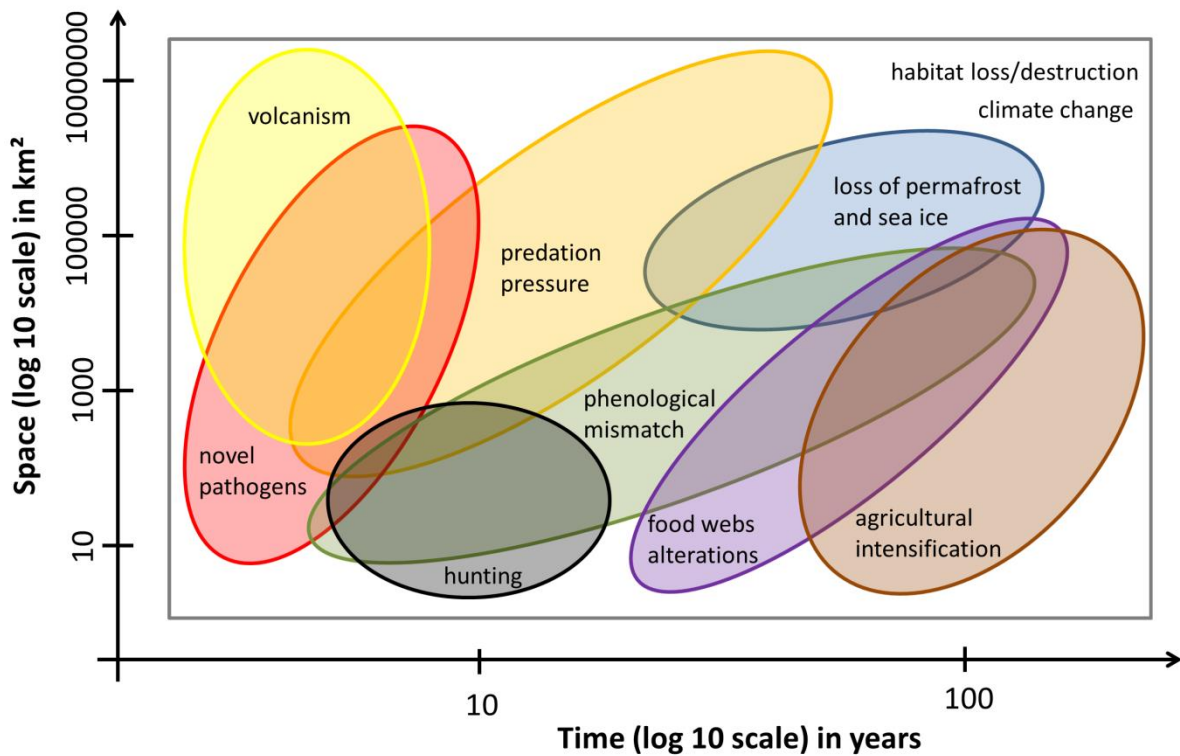
520

**(C) SPATIAL PATTERNS**



521

**(D) DISTURBANCE REGIMES OF STRESSORS**



522

523

**Figure 3. Temporal and Spatial Dynamics of Changes in Profitability of Migration for**

524

**the Arctic and North Temperate Long-distance Migrants.** Visualisations are following the

525 theory of pulse dynamics and disturbance in ecology [18]. **Temporal dynamics of migration**  
526 **profitability for acute stressors (A) and chronic stressors (B).** Various potential  
527 trajectories of stressors (e.g. changing amount or availability of food, pathogens prevalence or  
528 predation pressure) are projected.

529 **(C) Spatial dynamics of acute and chronic stressors. (D) Disturbance regimes of selected**  
530 **acute and chronic stressors in space and time.** Note that the discussed migratory benefits  
531 (food supply, pathogens prevalence and predation pressure) can be disrupted at any spatio-  
532 temporal scale. Examples from Table 1 are visualised as pictograms, where more negative  
533 changes underline more severe population dynamics consequences (A–C). But note that some  
534 of the case studies occupy wider space in reality. Similarly, selected stressors presented at  
535 spatio-temporal scale may not be limited to the mapped regions only (D). See also Online  
536 Supplemental Information Figures S2–S3.

537 **Table 1.** Case and comparative studies demonstrating recent disruptions of three historical  
538 advantages in Northward long-distance migration for terrestrial animals towards the North  
539 temperate and Arctic regions<sup>a</sup>.

Study, location and taxa	Description of current problems	Refs.
	<b>Food supply and availability</b>	
1) Alaskan and Canadian Arctic, six shorebird species (comparative study)	Despite high variability, generally prevailing mismatch between arthropod prey abundance and timing of breeding in shorebirds was connected to the snowmelt time, mismatches were more profound in Eastern locations, associated with steeper population declines of shorebird species there	[24], Figure 2
2) Scotland, United Kingdom Arctic skua <i>Stercorarius parasiticus</i> (case study)	Breeding population size of Arctic skuas declined by 81% between 1992–2015 alongside sharp declines in populations of their prey species black-legged kittiwake <i>Rissa tridactyla</i> , common guillemot <i>Uria aalge</i> , Atlantic puffin <i>Fratercula arctica</i> , Arctic tern <i>Sterna paradisaea</i> , linked to human and climate change impacts on food webs	[28]
3) The Netherlands, 10 migrating passerines (case study)	Mismatches for insectivorous passerine species and their prey was detected, with negative consequence for populations of migrating forest birds	[76]
4) Taimyr, Arctic Russia, red knot <i>Calidris canutus</i> (case study)	Reduced body size as a result of potential malnutrition during early life (mismatch with arthropod prey) was found with a negative consequence for survival at winter grounds in Mauritania, Africa	[77]
5) Kolguev Island, Kolokolkova Bay, Arctic Russia barnacle goose <i>Branta leucopsis</i> (case study)	The barnacle goose can skip stopover sites to advance its arrival to warming Arctic breeding grounds, but needs to refuel before egg-laying, resulting in a phenological mismatch between plants and offspring hatching late, reducing gosling survival	[27]
6) Barrow, Alaska, six shorebird species (case study)	Variable phenological mismatch was found but generally not sufficient food supply for families of three shorebird species	[23]
7) Svalbard, Norway snow bunting <i>Plectrophenax nivalis</i> (case study)	Changes in ambient temperature and precipitation on breeding grounds influence breeding productivity, suggesting decline in mean nestling body mass from 1998 to 2012	[78]
8) Canada caribou <i>Rangifer tarandus</i> (several case studies)	Phenological mismatches between plants and caribou at their summer grounds were suggested and discussed in several populations. From long-term perspective, caribou could benefit from increasing productivity in the Arctic, but altered plant community composition could be dominated by potentially less nutritious species	[17]
9) North America common green darner <i>Anax junius</i> (case study)	Dragonflies' migration is triggered by temperature and warming climate is expected to induce earlier spring flights, trigger later autumn flights and potentially shorten migratory distances and change wintering grounds and prey supplies	[79]
	<b>Pathogens and parasites</b>	
10) Eurasian Arctic, gulls, terns, auks, shorebirds and ducks (several case studies)	Spreading of helminth parasites and their increased impact on Arctic birds was described with examples of new host species colonisation, where parasites can reach maturity, although new hosts are phylogenetically unrelated to the main host	[33], Figure 2
11) Victoria Island, Canada, caribou <i>Rangifer tarandus</i> (case study)	Two species of protostrongylid nematodes have emerged for the first time in caribou, milder climates have facilitated spread of both parasites	[75], Figure 2

12) Central Kazakhstan saiga antelope <i>Saiga tatarica</i> (case study)	More than 200,000 saiga antelopes died in May 2015 from hemorrhagic septicemia caused by the bacterium <i>Pasteurella multocida</i> type B, following unusually high temperatures and humidity. The mass mortality event was spread across numerous calving grounds, reducing the regional population size of saigas by 85%	[35,36]
13) North America, monarch butterfly <i>Danaus plexippus</i> (several case studies)	More northerly hatched butterflies are recently more negatively affected by the parasite protozoan <i>Ophryocystis elektroscirrha</i> and fewer of them reach wintering sites in Mexico. Moreover, recently observed climate and human-induced shift of migratory to sedentary behaviour in several populations will likely lead to greater infection prevalence and can contribute to the species observed declines	[80]
14) Canadian Arctic, common eider <i>Somateria mollissima</i> (case study)	Recent outbreak of avian cholera caused by the bacterium <i>Pasteurella multocida</i> was recorded, with mortality rates of birds ranged from 1% to 43% of the local breeding populations	[34]
15) Alaska, bird populations at three locations (case study)	Avian malaria was detected in migratory as well as resident species of birds, for the first time documented avian <i>Plasmodium</i> transmission in the North American Arctic	[37]
16) Wisconsin, USA Eastern red bat <i>Lasiurus borealis</i> (case study)	Migratory bats were found with fungus <i>Pseudogymnoascus destructans</i> during June–September, illustrating the potential of detrimental White-nose syndrome to be transferred and dispersed among bats also at Northern breeding grounds during summer months	[81]
17) Europe painted lady butterfly <i>Vanessa cardui</i> (case study)	Painted lady butterflies are known for seasonal migrations from North Africa and South Europe to temperate and Arctic Europe to avoid high levels of parasitism from numerous Hymenoptera and Diptera parasitoids, however with rising ambient temperatures, parasitoids-free refuges might shrink	[82]
18) Kazakhstan saiga antelope <i>Saiga tatarica</i> (case study)	Saigas are being infected with gastrointestinal nematodes <i>Marshallagia marshalli</i> during their seasonal migration by grazing on pastures used by domesticated sheep	[83]
	<b>Predation</b>	
19) Global, 111 shorebird species (comparative study)	Significant increases of nest predation was found in the North temperate and Arctic regions during last 70 years, rapid especially in the Arctic and after year 2000	[45], Figure 2
20) Western Europe, five shorebird species (comparative study)	Significant increases of nest predation was detected during four decades until 2006, accompanied with decline in chick survival over the same period	[47]
21) Northern Sweden, pied flycatcher <i>Ficedula hypoleuca</i> (case study)	Increased nest predation was found over long term study following higher densities of mustelid predators: in 1965–1986 just 6% of the clutches on average were predated, whilst 26% in the period 1991–2017	[84]
22) Arctic, ducks, geese, gulls and auks (several case studies)	Polar bear <i>Ursus maritimus</i> , which is now with a disappearance of sea ice more often trapped on the land, has increased predation pressure on breeding colonies of Arctic birds	[52,53]
23) Svalbard, brant geese <i>Branta bernicla</i> (case study)	Recorded significant decrease of nests and young numbers on islands was associated with higher predator impact from polar bears and expanding great skuas <i>Stercorarius skua</i>	[85]
24) Svalbard, common eider <i>Somateria mollissima</i> (case study)	Recently observed high egg losses were associated with increasing predator pressure and declining eider populations. But historic predation rates were also high	[86]
25) Canadian Arctic, shorebirds (case studies)	Increased nest predation was indirectly caused by overabundant geese changing vegetation structure and nest detectability for predators	[65,87]

540 <sup>a</sup>These examples are illustrative and not exhaustive. Note also that the highlighted interactions  
541 may not be disruptive in all contexts. The Arctic and North temperate regions consist of  
542 various environmental mosaics that are highly dynamic in time or space, and local situations  
543 at particular location might counter the global trend. Patterns of high variability are obvious  
544 from detailed comparative studies on phenological mismatch [24] or nest predation [45].



545 **Glossary**

546 **Acute stressors (pulses):** abrupt changes in ecological parameters, e.g. food/prey abundance  
547 or pathogens pressure, following (mis)match in the given year or disease outbreak. This  
548 stressing event is changing the environment temporally, returning (pulsing) back to original  
549 state.

550 **Breeding grounds:** specific locations within the species breeding distribution range used for  
551 reproduction. For long-distance migrants, breeding grounds are often separated from non-  
552 breeding areas which include migration routes with stopover sites, staging sites and wintering  
553 grounds.

554 **Chronic stressors (presses):** gradual and directional changes in ecological parameters,  
555 persisting stress impacting ecosystems at a longer temporal scale, not returning to the original  
556 state, such as increased predation pressure over the years, loss of alternative prey and food  
557 web alterations or loss of permafrost/sea ice.

558 **Ecological traps and degraded environments:** ecological traps emerge when organisms  
559 make settlement decisions in a given location based on cues that were correlated formerly  
560 with habitat quality in a situation when better habitat alternatives are available nearby. The  
561 use of unreliable cues can lead to reduced reproductive output. In contrast to ecological traps,  
562 if there are no suitable alternatives in the surrounding area, then the entire landscape  
563 represents a degraded environment, with negative consequences for the population dynamics  
564 of species settling in the area.

565 **Long-distance migration:** migration when animals of the given species migrate regularly  
566 over 1,000 km between breeding and wintering grounds.

567 **Migration:** seasonal movements of individual animals or whole populations between  
568 breeding and wintering grounds.

569 **Migratory routes:** geographic routes along which animals migrate; for birds they are usually  
570 referred to as flyways.

571 **Stopover and staging sites:** important locations along migratory routes used by migrating  
572 animals for resting and energy refuelling, migrating animals can be found in high  
573 concentrations at those places.

574 **Trophic/phenological mismatch:** different rates of change of the seasonal timing of key  
575 phases in life cycles of interacting species, resulting in trophic asynchrony where the peak  
576 requirements of a predator species are offset from peaks in the abundance and availability of  
577 the prey.

578 **Supplemental Information**

579 **Animal Migration to Northern Latitudes: Environmental Changes and Increasing**  
580 **Threats**

581 Vojtěch Kubelka<sup>1-4\*</sup>, Brett K. Sandercock<sup>5</sup>, Tamás Székely<sup>2,3</sup> and Robert P. Freckleton<sup>1\*</sup>

582 <sup>1</sup>Department of Animal and Plant Sciences, University of Sheffield, Alfred Denny Building,  
583 Western Bank, Sheffield S10 2TN, UK

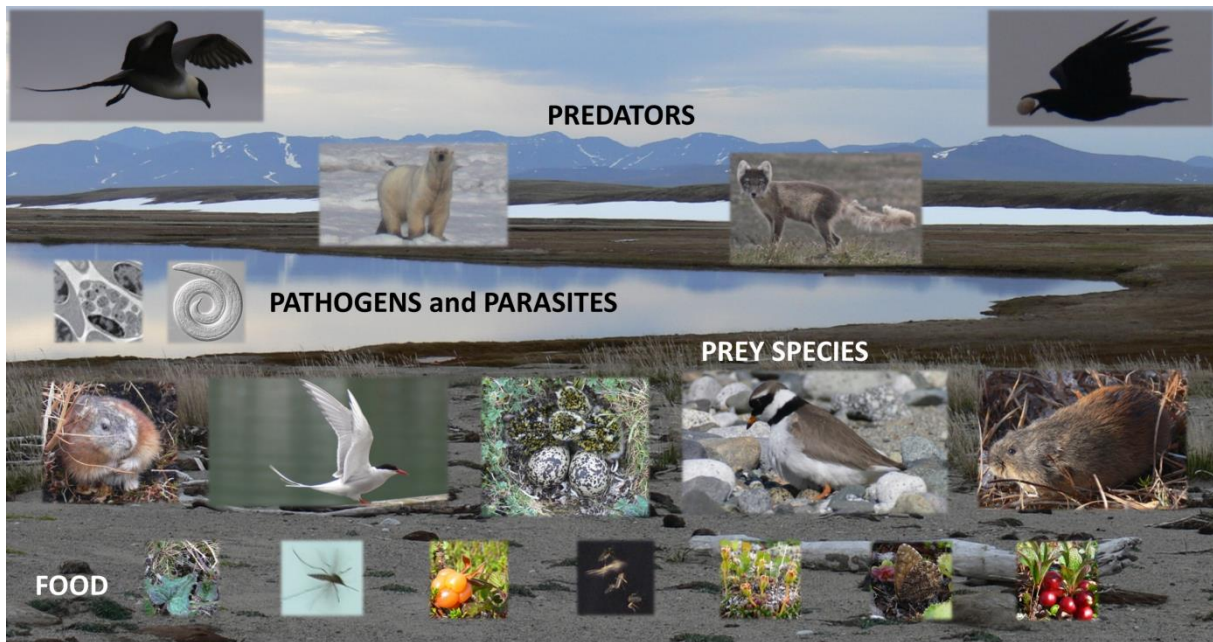
584 <sup>2</sup>Milner Centre for Evolution, University of Bath, Claverton Down, Bath BA2 7AY, UK.

585 <sup>3</sup>Department of Evolutionary Zoology and Human Biology, Faculty of Science, University of  
586 Debrecen, Egyetem tér 1, Debrecen, Hungary

587 <sup>4</sup>Department of Biodiversity Research, Global Change Research Institute, Czech Academy of  
588 Sciences, Bělidla 986/4a, Brno, 603 00, Czech Republic

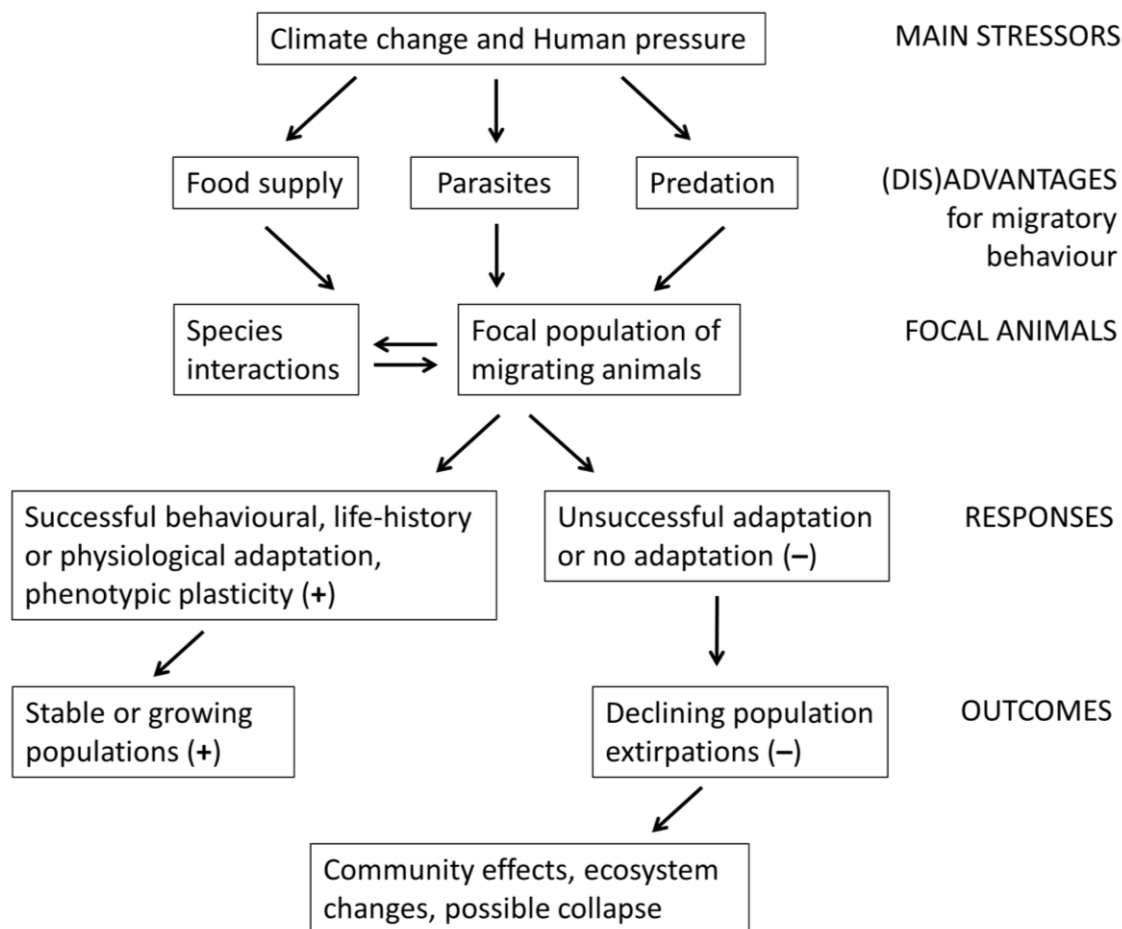
589 <sup>5</sup>Department of Terrestrial Ecology, Norwegian Institute for Nature Research, Høgskoleringen  
590 9, Trondheim, 7485, Norway

591 \*Corresponding authors: Kubelka, V. (kubelkav@gmail.com), Freckleton, R. P.  
592 (r.freckleton@sheffield.ac.uk)



593

594 **Figure S1. Key Interactions in the Arctic Ecosystem with Respect to Potential Benefits of**  
 595 **Migratory Behaviour.** Example organisms are depicted (all photos by Vojtěch Kubelka  
 596 except pathogens and parasites). **Predators.** Migratory or sedentary species of predators, from  
 597 left to right: long-tailed skua *Stercorarius longicaudus*, polar bear *Ursus maritimus*, Arctic  
 598 fox *Vulpex lagopus* and common raven *Corvus corax*. **Pathogens and Parasites,** interacting  
 599 with Arctic predators and prey species. *Plasmodium circumflexum* causing avian malaria on  
 600 the left and unidentified species of Protostrongylidae, Nematoda on the right (photos modified  
 601 from Salakij *et al.* (2012) [S1] and Kutz *et al.* (2007) [S2] respectively). **Prey species.** Arctic  
 602 rodents as the main prey species and migratory birds where nests and chicks represent suitable  
 603 alternative prey for Arctic predators. From left to right: Northern collared lemming  
 604 *Dicrostonyx groenlandicus*, Arctic tern *Sterna paradisaea*, nest with young of an American  
 605 golden plover *Pluvialis dominica*, common ringed plover *Charadrius hiaticula*, and a North  
 606 American brown lemming *Lemmus trimucronatus*. **Food.** Species of seasonally abundant  
 607 insects and plants representing food sources for prey species and their offspring. Figure S1 is  
 608 related to Figure 1.



610

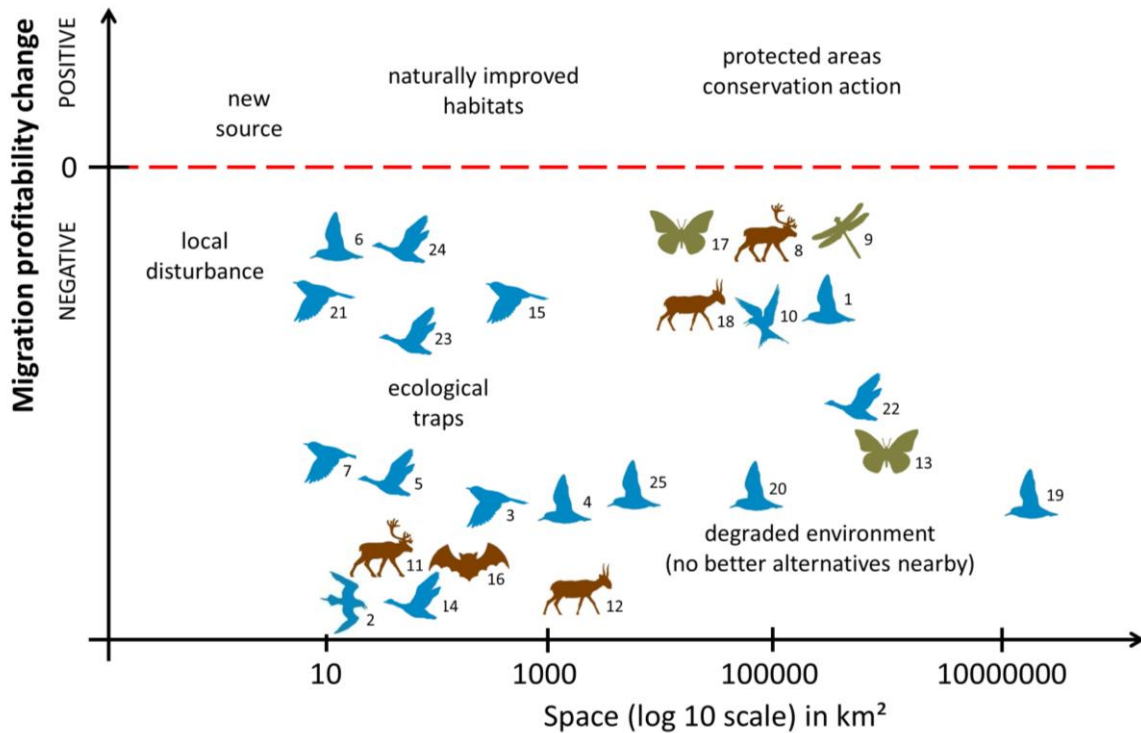
611 **Figure S2. Implications of a Changing Environment at Arctic and North Temperate**612 **Breeding Grounds for Migratory Animals.** A simplified chain of changes without possible

613 feedback loops for organismal responses to acute and chronic stressors: altered food supply,

614 parasitism and predation patterns. (Dis)advantages for migratory behaviour have three

615 different levels: 1) No changes or improvement; 2) Altered patterns but still advantageous

616 from migratory perspective – usually *sources* (+); 3) Ecological traps or degraded617 environments – usually *sinks* (-). Figure S2 is related to Figure 3.



618

619 **Figure S3. Spatial Dynamics of Profitability in Migration with Disturbances or Habitat**

620 **Improvements at Various Spatial Scales.** Acute and chronic stressors can precipitate in

621 negative impacts, ranging from local disturbances and regional ecological traps to large scale

622 degraded environments. Examples from Table 1 are visualised as pictograms but some of

623 them occupy wider space in reality. Figure S3 is related to Figure 3.

624 **Supplemental References**

625 S1 Salakij, J. *et al.* (2012) *Plasmodium circumflexum* in a Shikra (*Accipiter badius*):

626 Phylogeny and ultra-structure of the haematozoa. *Jpn. J. Vet. Res.* 60, 105–109

627 S2 Kutz, S.J. *et al.* (2007) Serendipitous discovery of a novel protostrongylid (Nematoda:

628 *Metastrongyloidea*) in caribou, muskoxen, and moose from high latitudes of North

629 America based on DNA sequence comparisons. *Can. J. Zool.* 85, 1143–1156