1	Improving predictions of climate change-land use change interactions
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18 Abstract

19 Climate change and land use change often interact, altering biodiversity in unexpected 20 ways. Research into climate change-land use change interactions has so far focused on 21 quantifying biodiversity outcomes, rather than identifying the underlying ecological 22 mechanisms, making it difficult to predict interactions and design appropriate conservation 23 responses. We propose a risk-based framework to further our understanding of climate 24 change-land use change interactions. By identifying the factors driving the exposure and vulnerability of biodiversity to land use change, and then examining how these factors are 25 26 altered by climate change (or vice versa), this framework will allow the effects of different 27 interaction mechanisms to be compared across geographic and ecological contexts, 28 supporting efforts to reduce biodiversity loss from interacting stressors.

29

30 Predicting biodiversity change when stressors interact

31 Climate change and land use change are two major drivers of biodiversity change [1, 2, 3]. 32 Predicting the effects of climate change and land use change on biodiversity is necessary to inform effective conservation strategies and ultimately safeguard biodiversity and the 33 34 benefits that humans derive from it [4]. The impacts of both drivers on species and ecosystems have been extensively studied in the past, mostly separately from each other 35 [5], and are relatively well understood. However, there is a rapidly growing body of evidence 36 showing that climate change and land use change do not always affect biodiversity 37 38 independently from each other, meaning that climate change alters the impact of land use 39 change on biodiversity, and vice versa [6]. It is these combined effects, or so-called climate change–land use change (CC–LUC) interactions, that are comparatively less well understood. 40

Most research into CC–LUC interactions has focused on identifying situations in which the combined impact of climate change and land use change could have dramatic negative effects on species or ecosystems [7]. For instance, land use change often reduces habitat availability and landscape connectivity, thereby reducing carrying capacity and dispersal between neighbouring populations, and increasing their sensitivity to extreme events. Specifically, populations fragmented or isolated by land use change are at a higher risk of decline and extinction as extreme climatic events become more frequent due to climate change (Figure 1, [8]). However, since climate change does not always exacerbate the
effects of land use change on biodiversity and vice versa [9, 10], it is equally important to
predict neutral or positive, as well as negative, outcomes to help improve targeting of
management and policy interventions.

52 Climate change and land use change, and their interactions, operate at different scales, posing challenges to effective conservation planning, resourcing, and management. At the 53 regional to global level, accounting for CC-LUC interactions could change conservation 54 55 prioritisation hierarchies of ecosystems and species (e.g., [11]), highlighting the need to identify species and ecosystems at the highest risk of adverse outcomes from CC-LUC 56 interactions. At the site level, CC-LUC interactions could affect which biodiversity 57 management options have the greatest effectiveness [12]. Understanding the potential 58 impacts of CC-LUC interactions on biodiversity will therefore provide critical information to 59 guide effective conservation interventions and to mitigate against the impacts of 60 anthropogenic global change at local and regional scales [13]. 61

62 Despite substantial progress in our understanding of interactions between climate change and global change stressors [14], including decades of research into CC–LUC interactions, 63 we currently have little ability to predict when and where these interactions are going to 64 happen, and how they are likely to affect biodiversity [7]. Predicting CC-LUC interactions is 65 challenging because climate change, land use change and biodiversity are all 66 multidimensional concepts [15, 16, 17], resulting in a high number of possible interactions. 67 68 For instance, climate change can entail changes in average temperature, shifts in season, or 69 a change in the frequency of extreme events, which may interact with a multitude of land use change effects, ranging in intensity from land conversion such as deforestation to more 70 71 subtle changes in land management (e.g. altering fertiliser regimes). As a result, predicting the presence, type and magnitude of CC–LUC interactions by looking at each potential driver 72 73 combination in turn is unlikely to provide comprehensive insights into the effects of multiple drivers and their interactions. Additionally, CC–LUC interactions are likely to be shaped by 74 interspecific interactions and trophic cascades [18, 19]. This is further complicated by the 75 76 fact that biodiversity responses at different organisational scales (e.g. individual behaviour,

population size, species composition) can play out over different timescales, and that CC–
LUC interactions can change over time [20].

To address the challenges in predicting and managing CC–LUC interactions, we 1)
summarise recent research into CC–LUC interactions, 2) demonstrate the need to expand
this research, which is currently focused on quantifying biodiversity outcomes, by focusing
on the mechanisms underpinning these interactions, and finally 3) propose a risk-based
framework as a way to efficiently identify key mechanisms governing the outcome of CC–
LUC interactions in different ecological contexts.

85

86 Climate change–land use change interactions: current state of play

87 What we know so far

To identify the main gaps in our understanding of CC-LUC interactions, we collated a 88 89 representative sample of peer-reviewed studies (including empirical studies, meta-analyses and reviews) that explicitly discuss or quantify an interaction between climate change and 90 91 land use change in the context of their effects on terrestrial and freshwater biodiversity (see 92 Annex 1 in the Supplementary Material for methodology). We excluded the marine realm since land use change does not directly affect large parts of the oceans. We did not consider 93 studies which only show that climate change alters the rate of land use change (or vice 94 95 versa; [6]). Although such studies identify situations in which biodiversity is affected by 96 combined climate and land use change (and that there is thus a chance for CC-LUC 97 interactions to occur), they do not directly consider how the impacts of climate change on 98 biodiversity are altered by land use change (and vice versa).

We considered 69 studies focusing on the combined effects of climate change and land use
change on biodiversity (see Annex 1 in the Supplementary Material for a complete list).
These studies addressed numerous features of biodiversity, including the distribution of
individual species (e.g., [21, 22], species abundance (e.g., [23]), response to disturbance
dynamics (e.g., [24]), species diversity (e.g., [25, 26]), or ecosystem composition and
processes (e.g., [27, 28]). Across these studies, we found two predominant empirical

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approaches to investigating CC–LUC interactions. First, some analyses compared biodiversity 105 106 outcomes between scenarios of no climate and land use change, either climate or land use 107 change, and combined climate and land use change (e.g., [29, 30, 31]). Second, other 108 analyses tested a dose-response relationship between climate, land use, an interaction 109 term, and biodiversity variables using a statistical model (e.g. [32, 33, 34]). Only 8 of the 110 empirical studies directly investigated interaction mechanisms (Table 1, [35, 36, 9, 37, 38, 39, 40, 41]). Interestingly, however, every review retrieved by our literature search (n = 11) 111 explicitly discussed mechanisms through which climate change could alter the impact of 112 113 land use change on biodiversity (and vice versa, Table 1).

114 Studies of CC–LUC interactions are drawn from different research fields with an emphasis on either climate, land use, or biodiversity science, and thus would benefit from a shared, 115 unifying framework to interpret and extract general patterns from the results. Previous 116 117 attempts to provide such a framework - based on studies of interactions between different stressors (including, but not limited to, climate change and land use change) - have focused 118 119 on classifying interactions based on how realised outcomes differ from expected outcomes, 120 i.e. those occurring in the absence of an interaction [42, 43, 44]. These classifications tend to distinguish between (a) independent effects (cases where climate change does not change 121 the effect of land use change on biodiversity, or vice versa), (b) antagonistic effects (cases 122 123 where climate change reduces the strength of the effect that land use change has on biodiversity, or vice versa), and (c) synergistic effects (cases in which climate change 124 125 increases the strength of the effect of land use change on biodiversity). Sometimes, a socalled dominance effect is included whereby climate change reduces the effect of land use 126 127 change to zero, or vice versa (e.g. [45, 46]), although dominance effects are more commonly 128 framed as an alternative null model describing an independent effect [47, 48].

129 Issues with the current approach

The current approach to researching CC–LUC interactions makes it difficult to synthesise insights from empirical studies that can predict the prevalence and effect of CC–LUC interactions. One reason for this is that there is no standard approach to formally define interaction types: what may be termed e.g. "synergy" by one study may not be considered an interaction at all, or an antagonistic interaction, by another [7]. To overcome this

challenge, however, it is not enough to develop a consensus on how interactions are 135 classified based on the difference between expected and observed outcomes. What 136 outcomes are "expected" always depends on the chosen null model, i.e. the expected 137 138 biodiversity outcome if no CC-LUC interaction occurs. This means that the choice of null model affects whether an interaction is classified as independent, antagonistic or 139 synergistic. Often, however, null models are not explicitly chosen but imposed by the choice 140 141 of statistical methods. As a result, there are now efforts to standardise null model choice in stressor interaction research to account for known differences in the mechanisms driving 142 143 the effects of single stressors on biodiversity [48] and thus to enable direct comparison of 144 results and the synthesis of insights across studies.

However, standardising the way we measure and classify outcomes of CC-LUC interactions 145 is by itself insufficient for the development of predictive power. For this, we need an 146 147 improvement in our understanding of the mechanisms underlying CC–LUC interactions. Since climate change, land use change and biodiversity each have multiple dimensions, 148 149 interactions that are classified as synergistic (or antagonistic, or independent, respectively) 150 are likely to include cases from many different geographic and ecological contexts, which may not be directly comparable. For instance, change in species richness, abundance or 151 152 interactions due to habitat loss may depend on climate change, but how it depends on 153 climate change varies between biomes and taxonomic groups [49]. The type, strength and direction of CC-LUC interactions is therefore shaped by a range of different biological or 154 ecological processes (Figure 1, Table 1) – put differently, the "surprising" outcomes that 155 characterise CC–LUC interactions likely result from different mechanisms, depending on 156 157 geographic and ecological context.

158

159 Using risk-based frameworks to predict interactions

The mechanistic pathways by which climate change and land use change interact are best identified using a framework based on risk, as this can improve our ability to predict the outcomes of CC–LUC interactions on biodiversity. Risk is the likelihood of an adverse outcome resulting from an external hazard, and can be conceptualised as a function of the

exposure to this hazard, as well as the intrinsic vulnerability of any particular entity to it [50, 164 51], where vulnerability is determined by sensitivity and adaptive capacity [52]. In a 165 biodiversity context, species, communities or ecosystems with high exposure and high 166 167 vulnerability are at a higher risk of an adverse outcome than other species, communities or 168 ecosystems (e.g. [53], Figure 2). Overall risk can be estimated by (i) identifying indicators for 169 each risk component (exposure, sensitivity, adaptive capacity [54]), so that each indicator 170 represents a process that affects the risk of an adverse outcome, then (ii) deriving an overall risk estimate, typically by combining scores from different risk components either 171 172 qualitatively [55] or quantitatively [56].

173 Risk-based frameworks have previously been used to identify the risk of single stressors such as climate change on species [57, 53], and have been adapted to include observed 174 175 outcomes of interactions between two stressors (e.g. [58]). Building on this work, we 176 propose a novel application of risk frameworks that identifies the mechanisms driving such interactions, and incorporates them into the assessment. Specifically, candidate interaction 177 178 mechanisms are systematically identified (and then tested) by asking how climate change 179 could alter the exposure and vulnerability of a species, community or ecosystem to land use change, i.e. how climate change can affect the components that determine risk of an 180 adverse outcome in response to land use change (and vice versa, Figure 2). 181

To illustrate this, consider risks from CC-LUC interactions to populations of a large predator, 182 such as African wild dogs (Lycaon pictus). This species declines in anthropogenically 183 184 modified landscapes due to reduced prey populations. Climate change (specifically 185 increased temperatures) is predicted to increase sensitivity of wild dogs to land use intensification by restricting the number of hours they can hunt [59]. Such time restrictions 186 around hunting compound the risk from reduced prey availability, and thus increase the 187 overall risk posed by land use change to this population. CC-LUC interactions could also 188 189 affect adaptive capacity. For instance, a species' ability to adapt to climate change by shifting its range can be impeded by habitat fragmentation, increasing the overall risk 190 191 posed by climate change (Figure 2). These mechanisms, which relate to changes in intrinsic vulnerability (sensitivity and adaptive capacity), correspond to "modification effects" [6], i.e. 192 true CC–LUC interactions (Table 1). 193

The risk framework approach we propose can also explicitly account for the direct effects of climate change and land use change on each other via effects on exposure, which need to be considered to estimate the overall impact on biodiversity. For instance, if the exposure of an ecosystem to climate change is determined by the magnitude of rainfall change, then land use change in the form of large-scale deforestation, which affects regional rainfall patterns, could increase the exposure of this particular ecosystem to climate change. Such interaction mechanisms correspond to Didham et al.'s [6] "chain effects".

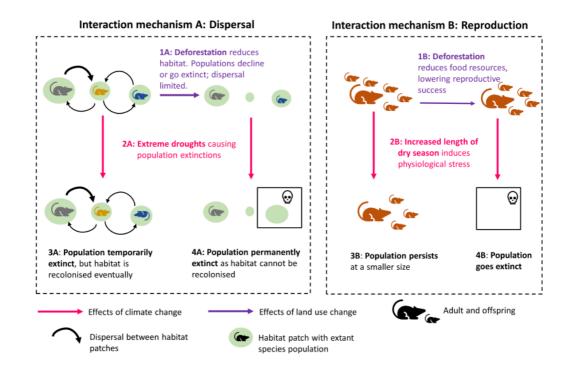
201 To account for CC–LUC interaction mechanisms within this framework, it is necessary to 202 identify risk components (exposure, sensitivity and adaptive capacity) with regard to both 203 climate change and land use change, as well as suitable risk indicators to estimate each component. Potential indicators may be drawn from existing frameworks and databases 204 that identify and quantify threats to biodiversity, such as the IUCN Red List of Species or 205 206 Ecosystems [60] and existing climate change or land use change risk assessments (e.g. [61, 57, 62]). Once risk components are known, candidate interaction mechanisms can be 207 208 identified based on known sets of possible interaction mechanisms (Table 1) as well as local 209 and expert knowledge. Which of these interaction mechanisms affect biodiversity in a given context can then be tested empirically. Interaction mechanisms that are shown to have 210 important effects on overall risk levels to biodiversity can subsequently be integrated into 211 212 risk assessments, either by modifying risk scores, or by including interaction mechanisms in quantitative risk models. 213

214 An important aspect of our risk framework is that it can be applied to any dimension of 215 biodiversity, such as genetic diversity or community composition. Indeed, the process explicitly considers all ways by which climate change may impact biodiversity's response to 216 land use change, as well as the ways by which land use change may impact biodiversity's 217 218 response to climate change, to ensure that the largest range of potential CC–LUC interaction 219 mechanisms are identified (see Table 1). The scope and flexibility of our framework can thus 220 be harnessed to provide conservation decision makers with context-specific information about all interaction mechanisms posing risks to all aspects of biodiversity at any given scale 221 222 or context.

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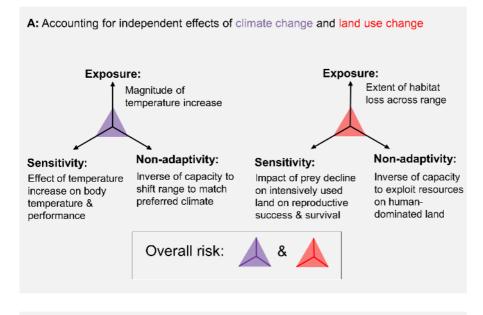
224 Concluding Remarks

- 225 Interactions between climate change and land use change can significantly shape
- biodiversity. So far, predicting their occurrence and impact has been hampered by a focus
- 227 on classifying the outcomes of interactions, rather than understanding the mechanisms by
- 228 which they operate. To advance our understanding of CC–LUC interactions, and to improve
- 229 our ability to mitigate their potentially negative impacts on biodiversity across different
- 230 geographic and taxonomic contexts, we recommend that future research focuses on
- investigating how the exposure and sensitivity of biodiversity to land use change, as well as
- its capacity to adapt to such change, is altered by climate change, and vice versa (see
- 233 Outstanding Questions). A key step towards this goal will involve interdisciplinary
- 234 cooperation e.g. among ecologists, physiologists, agronomists, and climate scientists as
- 235 insights from a range of fields are required to advance our understanding of how CC–LUC
- 236 interactions affect biodiversity, and to develop more effective risk assessment procedures
- to support environmental management worldwide.



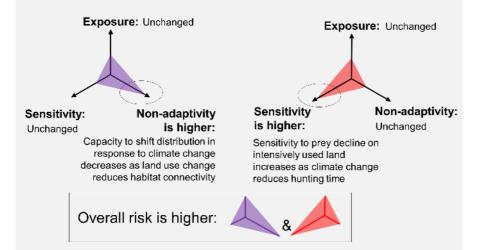
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Figure 1: Multiple mechanisms drive climate change-land use change interactions. In this 239 example, a combination of climate change and land use change drives population extinction 240 in both scenarios, but interaction mechanisms differ. In scenario A (left panel), deforestation 241 reduces habitat availability (green patches), reducing the size of three hypothetical 242 populations and, in some cases, leading to their extinction. Dispersal between these 243 244 populations is also reduced (1A). Climate change may also drive population declines, for 245 example by increasing the frequency of extreme droughts (2A). In absence of deforestation, these declines may be reversed by dispersal and recolonisation (inverted rodent icon, 3A). 246 However, in conjunction with deforestation, recolonisation of habitat patches is impossible, 247 leading to local extinction of some populations (skull icon; 4A). In scenario B (right panel), 248 habitat clearance reduces availability of food resources, leading to lower reproductive 249 success (fewer offspring), and therefore population decline, in a hypothetical population 250 (1B). Climate change may also reduce reproductive rates in this population, for instance by 251 252 increasing aridity, inducing physiological stress (2B). Habitat clearance again mediates the effect of climate change on the population: in its absence the population declines in size, 253 but persists (3B), whereas climate change in conjunction with habitat clearance leads to 254 population collapse and local extinction (4B). 255



Risk to a hypothetical population of African wild dogs (Lycaon pictus)

B: Accounting for climate change and land use change interactions



256

257 Figure 2: Using the concept of risk to conceptualise interactions between biodiversity

258 stressors. These diagrams illustrate the approach as applied to a hypothetical African wild

- 259 dog (Lycaon pictus) population. The risk of biodiversity change in response to a single
- 260 stressor is determined by different risk components; the overall risk increases as each
- 261 component increases. These components are *exposure* to a hazard (the rate or magnitude
- of the stressor that biodiversity experiences), and vulnerability of biodiversity to this hazard,
- which is determined by *sensitivity* (the magnitude of the biodiversity response to a unit of a
- given stressor), and *adaptive capacity* (the capacity of biodiversity to undergo changes in

265 response to a hazard that allow it to persist). Following [63], we use non-adaptability (NA), i.e. the inverse of adaptive capacity, to visualise this relationship, so that increases along this 266 267 axis represent increases in overall risk. Each risk component represents an environmental, 268 biological or ecological process that shapes biodiversity. If different stressors do not interact, the risk from a given stressor is independent from the presence of another (A). 269 Stressor interactions can be conceptualised as mechanisms by which a second stressor 270 alters processes that affect each risk component (B). In this example, land use change 271 decreases the African wild dogs' ability to adapt to climate change by limiting range shifts, 272 and climate change increases their sensitivity to land use change by limiting the time 273 274 available to hunt prey, which are already depleted owing to land use change. The 275 interaction of these effects increases overall risk from global change.

Table 1: Overview of known or hypothesised climate change–land use change interaction mechanisms. Examples are given of mechanisms

by which climate change can alter the sensitivity of biodiversity to land use change, or its capacity to adapt to land use change (and vice versa).

Asterisk (*) indicates references not captured by the systematic literature search.

Interaction mechanism	Description	References
Microclimate refugia	Land use change alters the structure of the vegetation canopy and the litter layer,	[8] [33] [36] [37] [64]
	as well as drainage patterns, and thus can create microclimates that either	
	accentuate or reduce sensitivity to climate change.	
Disturbance responses	Climate change reduces the resistance and/or resilience of ecosystems to	[8] [65] [66]*
	disturbance caused by land use change (e.g. by delaying recovery from habitat	
	disturbance), and vice versa, thereby increasing risk.	
Range shifts	Land use change can hinder adaptive range shifts, including access to climate	[8] [9] [41] [67] [68]
	refugia, reducing the habitat available to a species affected by climate	[69] [70] [71]
	change. Conversely, climate change can prevent the expansion of species into	
	habitat that land use change has made suitable (e.g. due to forest clearance or	
	abandonment of cultivation).	
Natural selection	Land use change can reduce local effective population size or gene flow, potentially	[72]* [73]*
	reducing or counteracting selection for genotypes that increase fitness under	
	climate change, and thus reducing adaptive capacity. Conversely, climate change	

	can lead to genetic homogenisation of populations, potentially reducing their	
	capacity to adapt to new ecological conditions caused by land use change.	
Genetic constraints	Co-adaptation to climate change and land use change could be difficult because of	[8] [74]*
	antagonistic pleiotropy (i.e. the same genes confer high fitness under climate	
	change but low fitness under land use change, or vice versa), or epistasis (i.e.	
	genetic interdependence) of traits conferring high fitness in the presence of one	
	driver but low fitness in the presence of another. This mechanism reduces the	
	capacity of a population to adapt to either stressor in the presence of the other.	
Metapopulation dynamics	Land use change can lower the size of habitat patches and increase the effective	[4] [8] [67] [68] [69]
	distance between them. Thus, species populations may decline or disappear within	[75] [76] [77] [78]
	patches, and incur reduced connectivity or genetic transfer between patches (e.g.	
	by constraints on dispersal of individuals or propagules), increasing the sensitivity of	
	metapopulations to climate change.	
Community filtering	Species can be co-tolerant or co-sensitive to climate change and land use change,	[38] [39] [79] [80]
	suggesting that the sensitivity of communities to subsequent climate change	
	depends on whether they have already been "filtered" by land use change, and vice	
	versa.	
Portfolio effect	Land use change can increase sensitivity of species communities to climate change	[40]
	by decreasing species richness and functional diversity. This is because such	

declines decrease the so-called portfolio effect whereby apparent high redundancy	
provides greater insurance or resilience in the face of climate change.	
Antagonistic species (e.g. predator, pathogen, dominant competitor) can benefit	[32, 35, 81] [82]*
from changes to habitat associated with land use change, increasing sensitivity to	
climate change for associated species (e.g. prey, host, subordinate	
competitor). Similarly, the risk of disease can be elevated by climate change	
(especially warming temperatures), reducing the resilience of populations to land	
use change.	
Climate change can disrupt mutualistic interactions by driving asymmetric range	[83]* [84]*
shifts or asynchronous phenology, for example between plants and their	
pollinators, thereby reducing population size and theoretically increasing sensitivity	
to land use change. Similarly, land use change can theoretically fragment	
populations of co-dependent mutualists and increase their sensitivity to	
phenological mismatches or other effects of climate change.	
A species community can adapt to climate change by shifting community trait	[85]*
distributions to match the new climatic conditions. Land use change could decrease	
the capacity of communities to adapt by limiting the arrival of new species whose	
traits match the new climatic conditions.	
	 provides greater insurance or resilience in the face of climate change. Antagonistic species (e.g. predator, pathogen, dominant competitor) can benefit from changes to habitat associated with land use change, increasing sensitivity to climate change for associated species (e.g. prey, host, subordinate competitor). Similarly, the risk of disease can be elevated by climate change (especially warming temperatures), reducing the resilience of populations to land use change. Climate change can disrupt mutualistic interactions by driving asymmetric range shifts or asynchronous phenology, for example between plants and their pollinators, thereby reducing population size and theoretically increasing sensitivity to land use change. Similarly, land use change can theoretically fragment populations of co-dependent mutualists and increase their sensitivity to phenological mismatches or other effects of climate change. A species community can adapt to climate change by shifting community trait distributions to match the new climatic conditions. Land use change could decrease the capacity of communities to adapt by limiting the arrival of new species whose

280 281	References
282 283 284	 IPCC (2014) Climate Change 2014: Synthesis Report. Contribution of Working Groups I, II and III to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change, IPCC.
284 285 286 287	 Newbold, T. <i>et al.</i> (2016) Has land use pushed terrestrial biodiversity beyond the planetary boundary? A global assessment. <i>Science</i> 353, 288-291.
288 289 290	3. Visconti, P. <i>et al.</i> (2016) Projecting global biodiversity indicators under future development scenarios. <i>Conserv. Lett.</i> 9, 5-13.
291 292 293	 Driscoll, D.A. <i>et al.</i> (2012) Priorities in policy and management when existing biodiversity stressors interact with climate-change. <i>Clim. Change</i> 111, 533-557.
294 295 296	5. Mazor, T. <i>et al.</i> (2018) Global mismatch of policy and research on drivers of biodiversity loss. <i>Nat. Ecol. Evol.</i> 2, 1071.
297 298 299	6. Didham, R. <i>et al.</i> (2007) Interactive effects of habitat modification and species invasion on native species decline. <i>Trends Ecol. Evol.</i> 22, 489-496.
300 301 302	 Côté, I.M. <i>et al.</i> (2016) Interactions among ecosystem stressors and their importance in conservation. <i>Proc. R. Soc. B</i> 283, 20152592.
303 304 305 306	8. Oliver, T.H. and Morecroft, M.D. (2014) Interactions between climate change and land use change on biodiversity: attribution problems, risks, and opportunities. <i>Wiley Interdiscip. Rev. Clim. Change</i> 5, 317-335.
307 308 309 310	9. Keshtkar, H. and Voigt, W. (2016) Potential impacts of climate and landscape fragmentation changes on plant distributions: Coupling multi-temporal satellite imagery with GIS-based cellular automata model. <i>Ecol. Inform.</i> 32, 145-155.
311 312 313 314	 Hamilton, C. M. <i>et al.</i> (2018) Slow and steady wins the race? Future climate and land use change leaves the imperiled Blanding's turtle (Emydoidea blandingii) behind. <i>Biol. Conserv.</i> 222, 75-85.
315 316 317 318	 Brown, C. J. <i>et al.</i> (2014) Interactions between global and local stressors of ecosystems determine management effectiveness in cumulative impact mapping. <i>Divers. Distrib.</i> 20, 538-546.
319 320 321	 Brown, C. J. <i>et al.</i> (2013) Managing for interactions between local and global stressors of ecosystems. <i>PloS one</i> 8, e65765.
322 323 324	 Staudt, A. <i>et al.</i> (2013) The added complications of climate change: understanding and managing biodiversity and ecosystems. <i>Front. Ecol. Environ.</i> 11, 494-501.
325 326 327 328	 Hooper, M. <i>et al.</i> (2013) Interactions between chemical and climate stressors: A role for mechanistic toxicology in assessing climate change risks. <i>Environment. Toxicol. Chem.</i> 32, 32- 48.

329 1 330 331	L5.	Noss, R. (1990) Indicators for monitoring biodiversity: a hierarchical approach. <i>Conserv. Biol.</i> 4, 355-364.
332 1 333	L6.	Kuemmerle, T. <i>et al.</i> (2013) Challenges and opportunities in mapping land use intensity globally. <i>Curr. Opin. Env. Sust.</i> 5, 484-493.
336	L7.	Garcia, R. <i>et al.</i> (2014) Multiple dimensions of climate change and their implications for biodiversity. <i>Science</i> 344, 1247579.
339	L8.	Valiente-Banuet, A. and Verdú, M. (2013) Human impacts on multiple ecological networks act synergistically to drive ecosystem collapse. <i>Front. Ecol. Environ.</i> 11, 408-413.
342	L9.	Bregman, T. <i>et al.</i> (2015) Species interactions regulate the collapse of biodiversity and ecosystem function in tropical forest fragments. <i>Ecology</i> 96, 2692-2704.
345	20.	Smith, M. D. <i>et al.</i> (2009) A framework for assessing ecosystem dynamics in response to chronic resource alterations induced by global change. <i>Ecology</i> 90, 2898-2907.
348	21.	Reino, L. <i>et al.</i> (2018) Modelling landscape constraints on farmland bird species range shifts under climate change. <i>Sci. Total Environ.</i> 625, 1596-1605.
351	22.	Trisurat, Y. <i>et al.</i> (2015) Assessing potential effects of land use and climate change on mammal distributions in northern Thailand. <i>Wildlife Res.</i> 41, 522-536.
354	23.	Lemoine, N. <i>et al.</i> (2007) Effects of climate and land-use change on species abundance in a central European bird community. <i>Conserv. Biol.</i> 21, 495-503.
357	24.	Bremer, L. <i>et al.</i> (2018) Bringing multiple values to the table: assessing future land-use and climate change in North Kona, Hawai'i. <i>Ecol. Soc.</i> 23, 33.
360	25.	Hoiss, B. et al. (2013) Combined effects of climate and management on plant diversity and pollination type in alpine grasslands. <i>Divers. Distrib.</i> 19, 386-395.
363	26.	Heubes, J. <i>et al.</i> (2013) The projected impact of climate and land use change on plant diversity: An example from West Africa. <i>J. Arid Environ.</i> 96, 48-54.
366	27.	Zhou, G. <i>et al.</i> (2002) Responses of grassland ecosystems to precipitation and land use along the Northeast China Transect. <i>J. Veg. Sci.</i> 13, 361-368.
367 368 2 369 370	28.	Aleman, J. <i>et al.</i> (2017) Tree cover in Central Africa: determinants and sensitivity under contrasted scenarios of global change. <i>Sci. Rep.</i> 7, 41393.
371 2 372 373	29.	Mantyka-Pringle, C S. <i>et al.</i> (2014) Understanding and predicting the combined effects of climate change and land-use change on freshwater macroinvertebrates and fish. <i>J. Appl. Ecol.</i> 51, 572-581.
374 375 3 376 377 378	30.	Kuemmerlen, M. <i>et al.</i> (2015) An attack on two fronts: predicting how changes in land use and climate affect the distribution of stream macroinvertebrates. <i>Freshw. Biol.</i> 60, 1443-1458.

380 381	31.	Lang, Y. <i>et al.</i> (2017) Responses of the water-yield ecosystem service to climate and land use change in Sancha River Basin, China. <i>Phys. Chem. Earth</i> 101, 102-111.
	22	Very studies and (2017) interesting effects of lead was and alignets an underst being
382	32.	Young, H. et al. (2017) Interacting effects of land use and climate on rodent-borne
383		pathogens in central Kenya. Philos. T. R. Soc. B. 372, 20160116.
384		
385	33.	Conenna, I. et al. (2017) Interactive effects of climate and forest canopy cover on Goshawk
386		productivity. <i>J. Ornithol. 158,</i> 799-809.
387		
388	34.	Benot, M. et al. (2014) Stronger short-term effects of mowing than extreme summer
389		weather on a subalpine grassland. <i>Ecosystems</i> 17, 458-472.
390		
391	35.	Jaramillo, J. et al. (2013) Climate change or urbanization? Impacts on a traditional coffee
392		production system in East Africa over the last 80 years. <i>PloS one</i> 8, e51815.
393		
394	36.	Senior, R. A. et al. (2018) Tropical forests are thermally buffered despite intensive selective
395		logging. <i>Glob. Change Biol. 24,</i> 1267-1278.
396		
397	27	Carón, M. et al. (2018) Regeneration responses to climate and land-use change of four
398	57.	subtropical tree species of the southern Central Andes. <i>Forest Ecol. Manag.</i> 417, 110-121.
		subtropical tree species of the southern central Andes. <i>Porest Ecol. Multug.</i> 417, 110-121.
399	20	Frickloff L O at $r/(2015)$ Thermolyniaka gradiets to be presented to be hitst conversion in
400	38.	Frishkoff, L. O. <i>et al.</i> (2015) Thermal niche predicts tolerance to habitat conversion in
401		tropical amphibians and reptiles. Glob. Change Biol. 21, 3901-3916.
402		
403	39.	Frishkoff, L. O. <i>et al.</i> (2016) Climate change and habitat conversion favour the same
404		species. <i>Ecol. Lett.</i> 19, 1081-1090.
405		
406	40.	Klaus, V. et al. (2016) Plant diversity moderates drought stress in grasslands: Implications
407		from a large real-world study on 13C natural abundances. Sci. Total Environ. 566, 215-222.
408		
100	11	
409	41.	Gimona, A. et al. (2012) Woodland networks in a changing climate: threats from land use
410	41.	Gimona, A. <i>et al.</i> (2012) Woodland networks in a changing climate: threats from land use change. <i>Biol. Conserv. 149,</i> 93-102.
410 411		change. Biol. Conserv. 149, 93-102.
410		
410 411		change. Biol. Conserv. 149, 93-102.
410 411 412		change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol.</i>
410 411 412 413	42.	change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol.</i>
410 411 412 413 414	42.	change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11,</i> 1278-1286.
410 411 412 413 414 415	42.	 change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11,</i> 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in
410 411 412 413 414 415 416	42. 43.	 change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11,</i> 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in
410 411 412 413 414 415 416 417	42. 43.	 change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11,</i> 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple
410 411 412 413 414 415 416 417 418	42. 43.	 change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11,</i> 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315.
410 411 412 413 414 415 416 417 418 419	42. 43. 44.	 change. <i>Biol. Conserv. 149</i>, 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11</i>, 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple stressors. <i>Ecol. Evol.</i> 5, 1538-1547.
410 411 412 413 414 415 416 417 418 419 420 421	42. 43. 44.	 change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11,</i> 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple
410 411 412 413 414 415 416 417 418 419 420 421 422	42. 43. 44.	 change. <i>Biol. Conserv. 149,</i> 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11,</i> 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple stressors. <i>Ecol. Evol.</i> 5, 1538-1547. Sala, O. <i>et al.</i> (2000) Global biodiversity scenarios for the year 2100. <i>Science</i> 287, 1770-
410 411 412 413 414 415 416 417 418 419 420 421 422 423	42. 43. 44. 45.	 change. <i>Biol. Conserv. 149</i>, 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11</i>, 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple stressors. <i>Ecol. Evol.</i> 5, 1538-1547. Sala, O. <i>et al.</i> (2000) Global biodiversity scenarios for the year 2100. <i>Science</i> 287, 1770-1774.
410 411 412 413 414 415 416 417 418 419 420 421 422 423 424	42. 43. 44. 45.	 change. <i>Biol. Conserv. 149</i>, 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11</i>, 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple stressors. <i>Ecol. Evol.</i> 5, 1538-1547. Sala, O. <i>et al.</i> (2000) Global biodiversity scenarios for the year 2100. <i>Science</i> 287, 1770-1774. Halpern, B. S. <i>et al.</i> (2008) Managing for cumulative impacts in ecosystem-based
410 411 412 413 414 415 416 417 418 419 420 421 422 423 424 425	42. 43. 44. 45.	 change. <i>Biol. Conserv. 149</i>, 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11</i>, 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple stressors. <i>Ecol. Evol.</i> 5, 1538-1547. Sala, O. <i>et al.</i> (2000) Global biodiversity scenarios for the year 2100. <i>Science</i> 287, 1770-1774.
410 411 412 413 414 415 416 417 418 419 420 421 422 423 424 425 426	42.43.44.45.46.	 change. <i>Biol. Conserv. 149</i>, 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11</i>, 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple stressors. <i>Ecol. Evol.</i> 5, 1538-1547. Sala, O. <i>et al.</i> (2000) Global biodiversity scenarios for the year 2100. <i>Science</i> 287, 1770-1774. Halpern, B. S. <i>et al.</i> (2008) Managing for cumulative impacts in ecosystem-based management through ocean zoning. <i>Ocean Coast. Manag.</i> 51, 203-211.
410 411 412 413 414 415 416 417 418 419 420 421 422 423 424 425	42.43.44.45.46.	 change. <i>Biol. Conserv. 149</i>, 93-102. Darling, E. S. and Côté, I. M. (2008) Quantifying the evidence for ecological synergies. <i>Ecol. Lett. 11</i>, 1278-1286. Crain, C M. <i>et al.</i> (2008) Interactive and cumulative effects of multiple human stressors in marine systems. <i>Ecol. Lett.</i> 11, 1304-1315. Piggott, J. J. <i>et al.</i> (2015) Reconceptualizing synergism and antagonism among multiple stressors. <i>Ecol. Evol.</i> 5, 1538-1547. Sala, O. <i>et al.</i> (2000) Global biodiversity scenarios for the year 2100. <i>Science</i> 287, 1770-1774. Halpern, B. S. <i>et al.</i> (2008) Managing for cumulative impacts in ecosystem-based

430 431 432	48.	Schäfer, R. B. and Piggott, J. J. (2018) Advancing understanding and prediction in multiple stressor research through a mechanistic basis for null models. <i>Glob. Change Biol.</i> 24, 1817-1826.
433 434 435 436	49.	Mantyka-Pringle, C. S. <i>et al.</i> (2012) Interactions between climate and habitat loss effects on biodiversity: a systematic review and meta-analysis. <i>Glob. Change Biol.</i> 18, 1239-1252.
437 438 439	50.	Turner, B. <i>et al.</i> (2003) A framework for vulnerability analysis in sustainability science. <i>Proc. Natl. Acad. Sci.</i> 100, 8074-8079.
440 441 442	51.	Polsky, C. <i>et al.</i> (2007) Building comparable global change vulnerability assessments: the vulnerability scoping diagram. <i>Glob. Environ. Change</i> 17, 472-485.
443 444 445 446	52.	IPCC (2014) Climate Change 2014: Impacts, Adaptation, and Vulnerability. Part A: Global and Sectoral Aspects. Contribution of Working Group II to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change. IPCC.
447 448 449	53.	Young, W. B. & Foden, B.E (2016) IUCN SSC guidelines for assessing species' vulnerability to climate change., IUCN, Ed., Cambridge, England and Gland, Switzerland.
450 451 452	54.	Schröter, D. <i>et al.</i> (2005) Assessing vulnerabilities to the effects of global change: an eight step approach. <i>Mitig. Adapt. Strateg. Glob. Change</i> 10, 573-595.
453 454 455	55.	Bland, L.M. <i>et al.</i> (2017) Guidelines for the application of IUCN Red List of Ecosystems Categories and Criteria, version 1.1., IUCN, Gland, Switzerland.
456 457 458	56.	Žurovec, O., Čadro, S., and Sitaula, B. K. (2017). Quantitative assessment of vulnerability to climate change in rural municipalities of Bosnia and Herzegovina. <i>Sustainability 9</i> , 1208.
459 460 461	57.	Ameca y Juárez, E. I. <i>et al.</i> (2013) Assessing exposure to extreme climatic events for terrestrial mammals. <i>Conserv. Lett.</i> 6, 145-153.
462 463 464	58.	Mantyka-Pringle, C. S. et al. (2015) Climate change modifies risk of global biodiversity loss due to land-cover change. <i>Biol. Cons.</i> 187, 103-111.
465 466 467	59.	Woodroffe, R., Groom R. and J. W. McNutt, JW. (2017) Hot dogs: High ambient temperatures impact reproductive success in a tropical carnivore. <i>J Animal Ecol</i> 86, 1329-1338.
468 469 470	60.	IUCN (2019) The IUCN Red List of Threatened Species. Version 2019-3. [Online]. Available: <u>http://www.iucnredlist.org</u> . [Accessed 19 2 2020].
471 472 473	61.	Pereira, H. M. <i>et al.</i> (2004) A framework for assessing the relative vulnerability of species to land-use change. <i>Ecol. Appl.</i> 14, 730-742.
474 475 476	62.	Lee, C. <i>et al.</i> (2018) A new framework to assess relative ecosystem vulnerability to climate change. <i>Conserv. Lett.</i> 11, e12372.

477 478 479	63.	Almeca y Juárez, E.I. <i>et al.</i> (2012) Natural population die-offs: causes and consequences for terrestrial mammals. <i>Tr. Ecol. Evol.</i> 27, 272-277.
480 481 482	64.	Greenwood, O. <i>et al.</i> (2016) Using in situ management to conserve biodiversity under climate change. <i>J. Appl. Ecol.</i> 53, 885-894.
483 484 485	65.	Bebi, P. <i>et al.</i> (2009) Snow avalanche disturbances in forest ecosystems—State of research and implications for management. <i>Forest Ecol. Manag.</i> 257, 1883-1892.
486 487 488	66.	Seidl, R. and Rammer, W. (2017) Climate change amplifies the interactions between wind and bark beetle disturbances in forest landscapes. <i>Lands. Ecol.</i> 32, 1485-1498.
489 490 491 492	67.	Opdam, P. and Wascher, D. (2004) Climate change meets habitat fragmentation: linking landscape and biogeographical scale levels in research and conservation. <i>Biol. Conserv.</i> 117, 285-297.
493 494 495 496	68.	Pitelka L. F. <i>et al.</i> (1997) Plant migration and climate change: a more realistic portrait of plant migration is essential to predicting biological responses to global warming in a world drastically altered by human activity. <i>Am. Sci.</i> 85, 464-473.
497 498 499	69.	Ewers, R. M. and Didham, R. K. (2006) Confounding factors in the detection of species responses to habitat fragmentation. <i>Biol. Rev.</i> 81, 117-142.
500 501 502	70.	García-Valdés, R. <i>et al.</i> (2015) Evaluating the combined effects of climate and land-use change on tree species distributions. <i>J. Appl. Ecol.</i> 52, 902-912.
503 504 505	71.	Barros, C. <i>et al.</i> (2017) Extreme climate events counteract the effects of climate and land-use changes in Alpine tree lines. <i>J. Appl. Ecol.</i> 54, 39-50.
506 507 508	72.	Vellend, M. (2004) Parallel effects of land-use history on species diversity and genetic diversity of forest herbs. <i>Ecology</i> 85, 3043-3055.
509 510 511	73.	Des Roches, S. et al. (2020) Climate-driven habitat change causes evolution in Threespine Stickleback. <i>Glob. Change Biol.</i> 26, 597-606
512 513 514	74.	Reusch, T. B and Wood, T. E. (2007) Molecular ecology of global change. <i>Mol. Ecol.</i> 16, 3973-3992.
515 516 517	75.	Maron, M. <i>et al.</i> (2015) Climate-induced resource bottlenecks exacerbate species vulnerability: a review. <i>Divers. Distrib.</i> 21, 731-743.
518 519 520	76.	Morelli, T. L. <i>et al.</i> (2012) Anthropogenic refugia ameliorate the severe climate-related decline of a montane mammal along its trailing edge. <i>Proc. R. Soc. B. 279, 4279-4286</i> .
521 522	77.	De Chazal, J. and Rounsevell, M. D. (2009) Land-use and climate change within assessments of biodiversity change: a review. <i>Global Environ. Change</i> 19, 306-315.

523		
524 525 526	78.	Farber, D. A. (2015) Separated at Birth: Addressing the Twin Crises of Biodiversity and Climate Change. <i>Ecology LQ</i> 42, 841.
527 528 529	79.	Vinebrooke, R. D. <i>et al.</i> (2004) Impacts of multiple stressors on biodiversity and ecosystem functioning: The role of species co-tolerance. <i>Oikos</i> 104, 451-457.
530 531 532	80.	Katayama, N. <i>et al.</i> (2014) Landscape heterogeneity–biodiversity relationship: effect of range size. <i>PloS one</i> 9, e93359.
533 534 535	81.	Bennett, J. M. <i>et al.</i> (2015) Climate drying amplifies the effects of land-use change and interspecific interactions on birds. <i>Landsc. Ecol. 30, 2031-2043.</i>
536 537 538 539	82.	Benning, T. L. <i>et al.</i> (2002) Interactions of climate change with biological invasions and land use in the Hawaiian Islands: modeling the fate of endemic birds using a geographic information system. <i>Proc. Natl. Acad. Sci.</i> 99, 14246-14249.
540 541 542	83.	Willis, C. G. et al. (2008) Phylogenetic patterns of species loss in Thoreau's woods are driven by climate change. <i>Proc. Natl. Acad. Sci.</i> 105, 17029-17033.
543 544 545	84.	Gómez-Ruiz, E. P. and Lacher Jr, T. E. (2019) Climate change, range shifts, and the disruption of a pollinator-plant complex. <i>Sci. Rep.</i> 9, 1-10.
546 547	85.	Norberg, J. and Moor, H. (2019) Amplitude and timescale of metacommunity trait-lag response to climate change. <i>BioRxiv</i> , 850289.
5/18		