

# 1 **Lake ecosystem tipping points and climate feedbacks**

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22 **Abstract**

23 Lakes and ponds experience anthropogenically-forced changes that may be non-linear and  
24 sometimes initiate ecosystem feedbacks leading to tipping points beyond which impacts become  
25 hard to reverse. In many cases climate change is a key driver, sometimes in concert with other  
26 stressors. Lakes are also important players in the global climate by ventilating a large share of  
27 terrestrial carbon back to the atmosphere as greenhouse gases, and will likely provide substantial  
28 feedbacks to climate change. In this paper we address various major changes in lake ecosystems,  
29 and discuss if tipping points can be identified, predicted, or prevented, as well as the drivers and  
30 feedbacks associated to climate change. We focus on potential large-scale effects with regional  
31 or widespread impacts, such as eutrophication-driven anoxia and internal phosphorus-loading,  
32 increased loading of organic matter from terrestrial to lake ecosystems (lake “browning”), lake  
33 formation or disappearance in response to cryosphere shifts or changes in precipitation to  
34 evaporation ratios, switching from nitrogen to phosphorus limitation, salinization, and the spread  
35 of invasive species where threshold-type shifts occur. We identify systems and drivers that could  
36 lead to self-sustaining feedbacks, abrupt changes and some degree of resilience, as opposed to  
37 binary states not subject to self-propelling changes or resilience. Changes driven by warming,  
38 browning, and eutrophication can cause increased lake stratification, heterotrophy (browning),  
39 and phytoplankton or macrophyte mass (eutrophication), which separately or collectively drive  
40 benthic oxygen depletion, internal phosphorus-loading and in turn increase greenhouse gas  
41 (GHG) emissions. Several of these processes can feature potential tipping point-thresholds,  
42 which further warming will likely make easier to surpass. We argue that the full importance of  
43 the vulnerability of lakes to climate and other anthropogenic impacts, as well as their feedback to  
44 climate is not yet fully acknowledged, so there is a need both for science and communication in  
45 this regard.

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## 47           1. Introduction

48 In natural sciences, the hysteretic behaviour of lakes (Scheffer et al. 2007) has informed the  
49 concept of tipping points at the ecosystem level, following the development of the alternative  
50 stable states theory in shallow lakes (Scheffer et al. 1993). Given the global vulnerability of  
51 freshwaters and the pervasive nature of major pressures acting upon them (e.g. nutrient pollution,  
52 over-extraction, and climate change), tipping points in these systems could have significant  
53 societal impacts, including on human and environmental health, clean water and food  
54 production, and climate regulation. The capacity to detect discontinuous ecosystem responses to  
55 pressure changes in natural systems has been challenged (e.g. Hillebrand et al. 2020; Davidson et  
56 al. 2023). Nevertheless, there are several studies that have reported the occurrence of tipping  
57 points even if they are difficult to detect (Lade et al, 2021; Seekell et al. 2022), such as shifts  
58 from one alternative state to another in small shallow lakes, the most common lake type globally  
59 (Messenger et al., 2016).

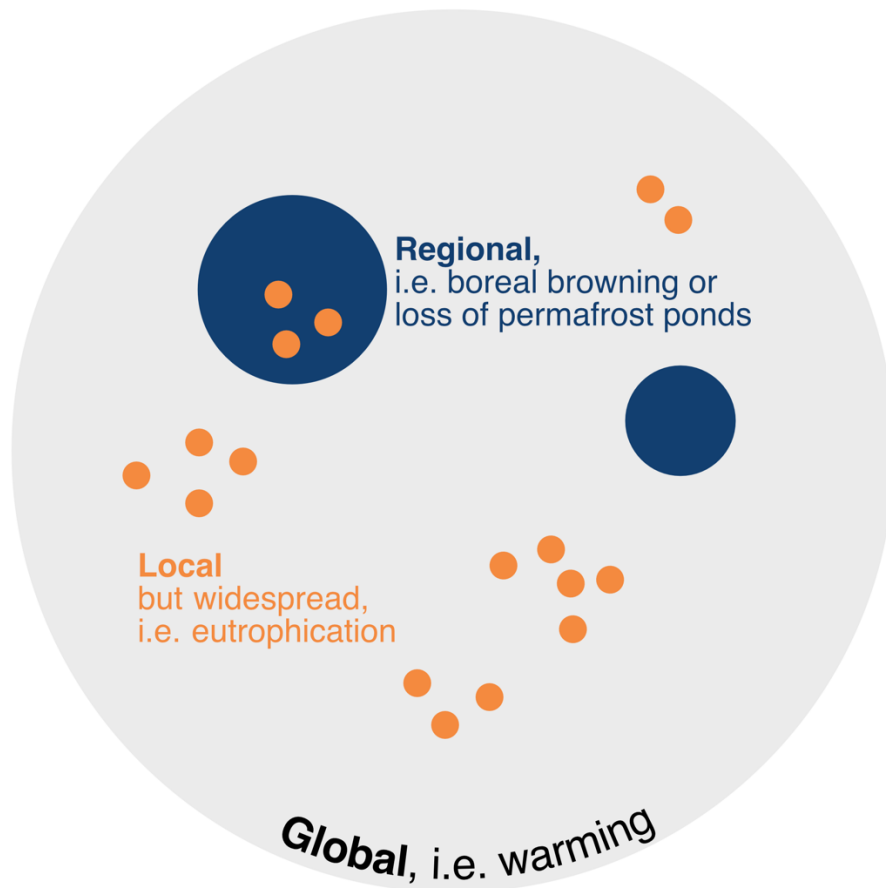
60           Some types of changes can be classified as *binary*, i.e. either-or situations at the system  
61 level. Increased temperature and/or reduced precipitation may induce negative water balance and  
62 shrinking of water volumes to the level where lakes or ponds simply disappear. Many lakes  
63 worldwide are facing reduced water volumes, but perhaps most striking is the widespread loss of  
64 high-latitude waterbodies, from Arctic or sub-Arctic ponds to wetlands or bogs. Such phenomena  
65 may qualify as one type of tipping point, but are not self-propelled by internal feedbacks *per se*,  
66 but rather by higher evaporation to precipitation ratios (Smol and Douglas 2007) or permafrost  
67 thaw (Smith et al. 2005; Webb et al. 2022; Smol 2023). While most tundra-ponds and other  
68 small waterbodies hardly qualify as *lakes* (Richardson et al. 2022), we mostly use the word lake  
69 through the text for simplicity, yet it will be evident from the context of wording where we  
70 specifically refer to ponds.

71           The question of what constitutes a “sudden” system shift, alternative stable states and  
72 hysteresis depends too on what is considered a relevant time span; days, years, decades or  
73 centuries. Also, systems may have alternative states that are not necessarily fixed over long time-  
74 spans, hence the phrase “stable” should be used with caution, so to the term ‘hysteresis’ infers  
75 little on the strength of the regulating processes. Uncertainty also remains on the geographical  
76 extent of tipping points in lakes and the wider relevance for the Earth’s climate system. Single  
77 lakes or local areas may experience non-linear or abrupt changes caused by local drivers, but we

78 here focus on potential tipping points of global or regional relevance, and with relevance to the  
79 climate system.

80 Empirical analyses, process modelling and experimental studies are advanced for shallow  
81 lakes, providing a good understanding of lake ecosystem behaviour around tipping points. There  
82 are related concepts in the literature (regime shifts, catastrophic shifts, forward switches, etc.),  
83 and there clearly many aspects of abrupt changes in nature and society that could be labelled a  
84 tipping point (Carrier-Belleau et al. 2022), but here we adopt the definition of a tipping point  
85 occurring when self-sustaining change in a system is triggered beyond a forcing threshold,  
86 typically starting with positive feedback loops, then entering a runaway phase before finally the  
87 tipping-point brings the system into a different alternative state (Nes et al. 2016; Lenton et al.  
88 2023). For example, the well documented increase of phosphorus (P) loading across European  
89 lakes in the last century (e.g. from agricultural and waste water pollution) has uncovered critical  
90 loading thresholds beyond which lakes can shift rapidly from a clear water, submerged  
91 macrophyte rich state to a turbid, phytoplankton dominated state (Scheffer et al., 2001; Jeppesen  
92 et al., 2005; Tátrai et al. 2008), and vice versa, when nutrient loading decreases. One of the  
93 theoretical implications is that to induce a switch back to the initial state the nutrient loading  
94 should be reduced to a lower threshold before the shift might be possible (hysteresis). Adding to  
95 such well-described and mechanistically well understood changes, there is a range of phenomena  
96 that may be perceived as tipping points. Hence, to provide structure to this complexity a range of  
97 tipping point candidates should be scrutinized against a common assessment approach. To  
98 qualify as tipping points, phenomena should not just be isolated phenomena in single lakes, but  
99 be more general and hold for specific (and widespread) types of lakes or waterbodies. Such  
100 phenomena may thus in the future occur across geographically distinct lake populations  
101 experiencing similar environmental change. In this way, the potential for identifying regional or  
102 global scale changes can be framed (Fig. 1).

103



104

105 Fig. 1. Impacts at levels that may qualify for tipping points at relevant spatial scales. Regional or  
 106 biome-wise effects could be loss of ponds and lakes due to permafrost thaw and/or increased  
 107 loadings of DOM in the boreal biome or salinization. Also local, but widespread changes such as  
 108 anthropogenic eutrophication of lakes in populated or intense agricultural areas would have  
 109 large-scale impacts. Lakes worldwide shows a warming trend, hence a global impact.

110

111

112 It is well established that lakes are sensitive to the effects of climate change, including  
 113 warming and changes in precipitation and storminess (e.g., Adrian et al., 2009; Meerhoff et al.,  
 114 2022). Emerging evidence suggests that lakes and ponds may also play an important role in  
 115 climate regulation, through both the emission of greenhouse gases (i.e. CO<sub>2</sub> and CH<sub>4</sub>,  
 116 predominantly CH<sub>4</sub>, Downing et al., 2021) and carbon burial (Anderson et al., 2020). Lakes and  
 117 rivers are impacted by climate change and other anthropogenic pressures globally, but they also  
 118 provide strong feedbacks to the global climate systems and carbon (C) cycle (Cole et al. 2007;  
 119 Tranvik et al. 2009), despite comprising a small part of global water extent. While global

120 estimates of net greenhouse gas (GHGs) emissions from lakes remain poorly constrained, there is  
121 general consensus that a significant fraction of terrestrially fixed C is degassed to the atmosphere  
122 via surface waters. Cole et al. (2007) conservatively estimated that inland waters annually  
123 receive some 1.9 Pg C y<sup>-1</sup> from the terrestrial landscape, of which at least 0.8 Pg C y<sup>-1</sup> is  
124 returned to the atmosphere through water to atmosphere GHG exchange. Later estimates revised  
125 this global GHG exchange term, to include evasion rates, at 2.1 Pg C yr<sup>-1</sup>, from lakes, rivers and  
126 reservoirs (Raymond et al. 2013). Notably, boreal lakes are important conduits of CO<sub>2</sub> release to  
127 the atmosphere, estimated to be equivalent to the annual CO<sub>2</sub> release from forest fires, globally  
128 (Hastie et al. 2017). Under a high CO<sub>2</sub>-emission scenario and as a result of increased terrestrial  
129 NPP, CO<sub>2</sub> emissions from boreal lakes are projected to increase by 107%, showing the coupling  
130 between the terrestrial and aquatic C cycle (Hastie et al. 2017).

131 This significant role of surface waters for GHG-emissions is also highly relevant, but  
132 poorly constrained, in both national and global C-budgets (Lindroth and Tranvik 2021). The  
133 relationship between inputs of organic C and nutrients is a key determinant of the balance  
134 between heterotrophic and autotrophic processes, determining the biodiversity, community  
135 composition and food web structure, as well as the productivity-to-respiration (P:R) ratio. And  
136 so, it is relevant to consider the extent to which potential tipping points may drive, or be driven  
137 by, climate change, leading to higher level feedbacks to the Earth's climate system.

138 Here, we discuss candidate tipping points in freshwaters reported in the literature (based  
139 on literature searches including the term 'tipping point' and either 'lake' or 'pond') as well as  
140 experience and insights of the author team. The discussion on each is constrained to waterbody  
141 categories with the potential for global or at least regional or biome-scale relevance. In this  
142 context we also constrain the discussion to potential tipping points that are more generic, at least  
143 carrying regional or biome-wide impact, and that could have feedbacks to the climate, while not  
144 necessarily being driven or triggered by climate change *per se*.

145 We identify 6 candidate categories for tipping points at a relevant scale in this context  
146 (regional to global impact), and for each of the categories we discuss whether observed changes  
147 can be categorised as tipping points according to the definition above. We also address climatic  
148 and other drivers and consequences, including potential feedbacks to the climate system, and  
149 wider societal implications, with emphasis on the most relevant and influential categories.

150 **2. Candidates and categories of lake tipping points**

151 In principle, any abrupt or sudden stress imposed on a waterbody could result in specific  
 152 impacts, i.e. toxic waste or toxic treatments (e.g. rotenone to kill off undesired species; runoff of  
 153 herbicides inadvertently killing aquatic plants), hydrological alterations by impoundment or  
 154 canals, and stocking or immigration of new (often exotic) species. In some cases, when stressors  
 155 are removed the system will return abruptly to its original state. To qualify as a tipping point,  
 156 here, we consider that system response should be self-sustaining and involve positive feedbacks,  
 157 in line with the criteria set out in Nes et al. (2016) and Lenton et al. (2023). To be relevant in a  
 158 wider context, the tipping point should be more generic to certain types of impact, certain types  
 159 of waterbodies, and potentially also have feedbacks to the climate in terms of GHG-emissions.  
 160 We have identified 6 stressors that may trigger a freshwater ecosystem to cross a tipping point  
 161 (Table 1) and scrutinise them in turn below.

162

163 Table 1. Candidate events from the literature with potential to occur at local to regional scales, their association with  
 164 climate change, and whether tipping points and hystereses have been associated with them. Brackets indicate higher  
 165 uncertainty.  
 166

Type of event	Local, common	Regional	Climate driver	Climate feedback	Tipping point	Hysteresis
Eutrophication driven water anoxia and internal P-loading	x		x	x	x	x
Increased loadings of DOM		x	x	x	(x)	(x)
Disappearance/ appearance of waterbodies		x	x	x	x	(x)
Switch between N and P limitation		x	x	(x)		
Salinization		x	x	x		(x)
Spread of invasive species	x	(x)	(x)			(x)

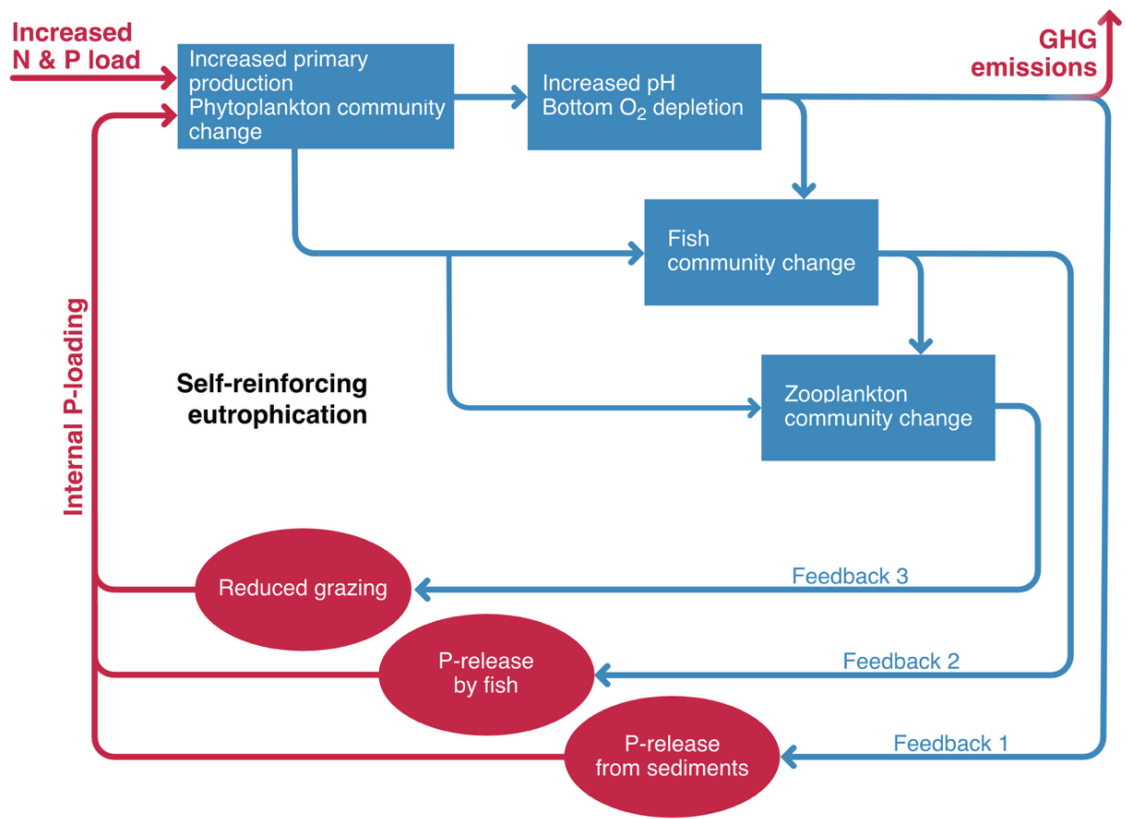
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169 **2.1. Eutrophication, anoxia and internal P-loading**

170 Eutrophication is one of the most pervasive stressors on fresh water and coastal systems.  
 171 Although it may naturally occur due to inputs from the watershed or from biota translocating  
 172 nutrients from connecting ecosystems, eutrophication is a largely human-induced phenomenon.  
 173 The main causes of cultural eutrophication have varied across time and regions. However, it is  
 174 widely accepted that the main current cause of eutrophication is the change in land use in

175 watersheds, and, particularly agricultural activities driving diffuse nutrient pollution (as well as  
 176 other agrochemicals) (Moss 2008; Schulte-Uebbing et al., 2022). Agriculture, with myriad  
 177 impacts on fresh waters that go well beyond nutrient pollution (Moss, 2008), has been identified  
 178 as a major driver of ecosystem shifts and tipping points (Gordon et al., 2008).



179  
 180  
 181 Fig. 2. Feedback loop diagram for eutrophication, demonstrating key feedbacks that can amplify  
 182 P-loading, and beyond a tipping point cause self-sustaining change. Eutrophication and internal  
 183 mobilization of P cause high algal biomass, decrease benthic oxygen or anoxia, and consequently  
 184 also increased greenhouse gas emissions. Blue denotes primary responses, red the secondary  
 185 feed-back responses as well as the key driver and consequence (nutrient loading and GHG-  
 186 release).

187  
 188 The mobilisation of P from lake bed sediments, a process known as internal loading  
 189 (Sondergaard et al., 2001), plays a key role in hysteresis in lakes following the reduction of P  
 190 loading from the catchment (Boström et al. 1982; Jeppesen et al. 1991; Spears & Steinman  
 191 2020). In this context, hysteresis can be strengthened by eutrophication-driven biological  
 192 changes in fish composition and size structure that have cascading effects on zooplankton and



193 phytoplankton as well as strong impacts on fish-mediated nutrient cycling (Brabrand et al. 1990).  
194 This in turn will maintain a system with deep water anoxia and high nutrient load, supporting the  
195 release of GHGs (Fig. 2).

196

### 197 *Feedbacks and Tipping points*

198 The phenomenon of eutrophication is local, but widespread, and likely to worsen in its  
199 manifestations as a result of climate change (Moss et al., 2011; Meerhoff et al., 2022). In  
200 particular, the process of internal loading may be enhanced by lake warming (Jeppesen et al.,  
201 2009) due to an increased metabolism of bacteria and an acceleration of biochemical reactions.  
202 Warming also increases stratification and the duration and strength of stratification, also  
203 promoting anoxia (Maberly et al. 2020; Woolway et al. 2020; 2022). As a case example, this  
204 phenomenon is well documented by the recent study of the Danish, shallow and highly eutrophic  
205 lake Ormstrup (Davidson et al. 2024). Increases in precipitation, and high intensity rainfall  
206 events, are also expected to significantly increase runoff of P from agricultural catchments to  
207 surface freshwaters (Ockenden et al., 2017), further promoting eutrophication and its  
208 manifestations.

209 The different states of shallow lakes can feedback differently on climate by either  
210 reducing or increasing GHG emissions (Hilt et al. 2017). Clear and turbid lakes differ in their  
211 CO<sub>2</sub> emissions due to the magnitude of CO<sub>2</sub> uptake by primary producer photosynthesis. Efflux  
212 of CO<sub>2</sub> appears to decrease when submerged macrophytes establish after the reduction of  
213 nutrient loading (Jeppesen et al., 2016). Submerged-macrophyte dominated shallow lakes tend to  
214 emit lower CH<sub>4</sub> by ebullition and diffusion than phytoplankton dominated, turbid lakes (Colina  
215 et al., 2022; Davidson et al. 2018). The turbid state in particular feeds back on climate since  
216 warming and eutrophication-induced water anoxia could offset increased CO<sub>2</sub>-fixation by  
217 blooms or by macrophytes as lower oxygen levels stimulate methane (CH<sub>4</sub>) emission, with CH<sub>4</sub>  
218 emissions from eutrophic systems expected to increase with 6-20% with each degree of warming  
219 (Aben et al. 2017).

220 The eutrophication and warming-associated shift from submerged macrophyte dominance  
221 to phytoplankton or floating plant dominance may also strongly increase greenhouse gas  
222 emissions, particularly CH<sub>4</sub> (Aben et al. 2022). Cyanobacterial blooms, a typical manifestation of  
223 eutrophication and high internal P-loading, can both promote CO<sub>2</sub> sequestration and produce

224 CH<sub>4</sub>. CH<sub>4</sub> can be produced even under oxic conditions as a by-product of photosynthesis (Bižić  
225 et al., 2020). Blooms often create anoxic layers in surface sediments or through the water column  
226 after their collapse, favouring the production and transport of CH<sub>4</sub> via methanogenesis (Li et al.,  
227 2021; Yan et al., 2017). Cyanobacterial blooms are thus considered a key mechanism by which  
228 eutrophication has a positive feedback on climate change (Bižić 2021; Yan et al., 2017).  
229 Although increased inputs on N from atmospheric deposition or catchment runoff are the main  
230 causes of elevated N<sub>2</sub>O release from lakes (Yang et al. 2015), warming also impacts aquatic N<sub>2</sub>O  
231 emissions. N<sub>2</sub>O emissions are estimated to increase by 8 – 14% for each degree of warming  
232 (Velthuis and Veraart 2022), highlighting another strong climate feedback.

233 Despite the fact that nutrient loading is still the major driver of eutrophication (including  
234 algal blooms; e.g. Bonilla et al., 2023), climate change is also expected to promote  
235 eutrophication (Moss et al., 2011; Meerhoff et al., 2022). Indeed, interaction between  
236 temperature and trophy has been observed to produce synergistic emission responses in  
237 experimental lakes (Davidson et al., 2018) and warming alters resident microbial communities to  
238 favour methanogenesis over methanotrophy (Zhu et al., 2020). It is thus likely that warming  
239 decreases the nutrient thresholds for a tipping point leading to a shift to an alternative state in  
240 shallow lakes and ponds.

241 The predominantly amplifying influence of climate change on eutrophication-driven  
242 tipping points in lakes provides a mechanism for coherent threshold exceedance beyond the local  
243 scale, with more widespread eutrophication-induced tipping points expected with further  
244 warming. However, despite the dearth of studies that generate bi-directional carbon flux data to  
245 assess the balance between emission and burial in lakes, it remains unknown whether the climate  
246 feedback can be buffered by projected eutrophication-driven increases in lake carbon burial  
247 (Anderson et al. 2020). Moreover, robust projections are lacking for climate impacts on  
248 eutrophication, with no emergent regional to global warming threshold identifiable beyond  
249 which a nonlinear increase in these localised tipping points occur (Grasset et al. 2020). In  
250 general, tipping points becomes harder to predict in a warmer climate (Kosten et al. 2009).

251

## 252 **2.2. Increased loadings of DOM in the boreal biome**

253 Over thousands to millions of years, the feedback between terrestrial vegetation and aquatic  
254 productivity has been essential for the evolution of the atmosphere and the global climate

255 (Beerling 2007). Vegetation serves not only as a major C pool and eventually a source of total  
256 organic carbon (TOC) in boreal areas, but it also promotes root exudates of CO<sub>2</sub> and organic C.  
257 This enhances weathering rates thereby increasing the flux of nutrients (P, N, Si, Fe, Ca and  
258 carbonate (CO<sub>3</sub>)) (Humborg et al. 2004; Hessen et al. 2009) to surface waters. The availability of  
259 nutrients subsequently enhances aquatic productivity, and thereby C-sequestration. In addition,  
260 the carbonate species are important for buffering capacity towards acidification in fresh water  
261 and marine systems. On different timescales there is thus a range of feedback mechanisms  
262 between terrestrial and aquatic ecosystems that demands a better understanding. Tracking past  
263 history (Holocene) tree-line, forest cover and lake sediments, revealed a strong and consistent  
264 link between climate, forest cover and lake TOC (Rühland et al. 2003; Rosén 2005). Thus, at  
265 least on the centennial scale, there is a strong temporal TOC-link between terrestrial and aquatic  
266 systems. Allochthonous C derived either directly as leachate from litterfall and roots or indirectly  
267 via partial decomposition of organic matter in the soils, constitutes the (by far) dominant pool of  
268 dissolved organic matter (DOM) in boreal freshwaters, hence forest cover and fraction of bogs  
269 and wetland areas in the catchment are key determinants for the concentration and color of this  
270 terrestrially derived chromophoric DOM (Dillon and Molot 1997; Kortelainen et al. 2006;  
271 Larsen et al. 2011a).

272         Since terrestrially derived C is a main determinant of freshwater C, any changes in  
273 terrestrial primary production and export of organic C will invariably also increase aquatic  
274 outputs of CO<sub>2</sub>. Increased terrestrial productivity has been linked to “CO<sub>2</sub>-fertilization” (Huang  
275 et al. 2007) yet these CO<sub>2</sub> effects will be constrained by N-availability. Elevated N-deposition  
276 due to human emissions has driven an increase in the forest C sink in tandem with the CO<sub>2</sub>-  
277 fertilization effect, while at the same time also increased the deficiency of P (and other key  
278 elements allocated to tree biomass) (De Vries et. al. 2006).

279         Increased export of terrestrially derived DOM to lakes and rivers in boreal regions  
280 (“browning”) is a widespread phenomenon partly linked to reduced acidification, but also driven  
281 by land-use changes (notably afforestation) and climate change (CO<sub>2</sub>-fertilization of forests,  
282 warming and hydrology) (de Wit et al., 2016; Creed et al. 2018; Monteith et al., 2023). An  
283 empirically based space-for-time model of changes in the Normalised Difference Vegetation  
284 Index (NDVI) under a 2° C climate scenario predicts continued browning of boreal lakes (Larsen  
285 et al. 2011b). Forest dynamics are slow, however, hence space-for-time scenarios projecting

286 increased flux of TOC from catchments owing to increased forest cover could require centuries  
287 to play out. Thus, catchment properties governing *production* of TOC such as forest size and  
288 fraction of bog and wetland areas could very well be temporally decoupled from the export,  
289 especially considering the large stock of organic matter typically present in boreal catchments.

290 Time series analysis (30 years) of data from 70 Norwegian catchments and lakes  
291 provided evidence for a tight temporal coupling between the decadal increase in land “greening”  
292 (with NDVI as a proxy) and lake browning (with TOC as a proxy) (Finstad et al. 2016), and the  
293 browning on northern lakes can, to a large extent, be attributed to recent afforestation (Kritzberg  
294 2017; Skerlep et al. 2020). The prominent “greening” by increased vegetation cover trend in  
295 many boreal and alpine regions (Guay et al. 2014) and increase in forest volume (cf. Opdahl et  
296 al. 2023) will thus have bearings on lakes and rivers in these regions. There are a number of  
297 confounding explanatory drivers for this greening: warming, elevated CO<sub>2</sub>, accumulated nitrogen  
298 deposition and changes in grazing activities as well as forestry practices are all implicated. An  
299 extended growing season has also been recorded (Barichivich et al. 2013), and elevated levels of  
300 CO<sub>2</sub> per se may contribute to this (Piao et al. 2006). Collectively, these changes in environmental  
301 drivers and pressures yield an increase in terrestrial net primary production (NPP), notably at  
302 high latitudes (Forkel et al. 2016). Since a significant fraction of the terrestrial NPP will be  
303 exported to surface waters as DOM, it means that terrestrial greening could lead to freshwater  
304 browning.

305 The role of forest cover is further accentuated by a need for a carbon-negative future (i.e.  
306 net drawdown of CO<sub>2</sub> from the atmosphere) where widespread afforestation is the only currently  
307 feasible means of reducing atmospheric concentrations of CO<sub>2</sub> beyond the continued action of  
308 natural carbon sinks (MacDougall et al., 2020). However, such afforestation also comes with  
309 climate costs, both in terms of decreased albedo (Betts and Ball 1997; Bathiany et al. 2010;  
310 Lawrence et al. 2022) and as argued above, the potential for increased production and degassing  
311 of GHGs from surface waters. Enhanced primary production in forested catchments stimulated  
312 by reactive N deposition has, by increasing the pool of C available for fluvial export, been linked  
313 to increased C burial in northern lakes over the past two centuries (Heathcote et al. 2015). Again,  
314 this highlights the need for improved understanding of the balance between C emissions and  
315 burial in lakes in response to browning (Williamson et al., 2015) and other identified stressors in  
316 order to better constrain climate feedbacks. Browning will also promote darkening of coastal

317 waters with as yet unknown climate feedbacks (Opdal et al. 2023). The question that remains is  
318 whether these terrestrial and aquatic responses are directly coupled in time, or if there is a  
319 delayed aquatic response in the order of decades or even millennia. Another question is how the  
320 CO<sub>2</sub> in itself could boost these processes, and how this skewed C-supply to autotrophs could  
321 affect land-aquatic interactions?

322         Wide-scale shifts in boreal lakes caused by increased loadings of DOM can promote a  
323 prolonged and more intensified stratification period (implications summarized above, described  
324 for DOM by Spears et al., 2017), amplified by warming. Increased terrestrial DOM loadings  
325 intensify net heterotrophy in the systems (i.e. through increased light attenuation and increased  
326 access to organic C for heterotrophic bacteria) (Hessen et al. 1990; Karlsson et al. 2007; Thrane  
327 et al 2014; Horppila et al. 2023). A well explored case study as an example of these impacts,  
328 which is also linked directly to the tipping point concept, is the Swedish, boreal brownwater lake  
329 Härsvatten where a long-term study clearly links loadings of DOC to anoxia (Spears et al. 2017).  
330 While at present the thresholds around these effects have not been well constrained, the impacts  
331 may be significant at the global scale for GHG emissions (Tranvik et al. 2009) and regionally for  
332 coastal primary producers (Opdal et al. 2019). Given the strong empirical links between drivers  
333 and consequences, it means that impacts and feedbacks can be predicted qualitatively, while not  
334 yet quantitatively.

335         The temporal aspect also deserves further attention. If the main source of browning is  
336 afforestation, responses will proceed slowly compared with cases where reduced acid deposition  
337 is the main driver, yet both drivers operate on decadal timescales. In the latter case, the browning  
338 could represent a *re-browning* (Meyer-Jacob et al. 2020).

339

#### 340 *Feedbacks and tipping points*

341 The links and feedbacks between climate to land to lakes and back to climate in terms of  
342 increased GHG-emissions is conceptually well understood, and also the main drivers for the  
343 specific GHGs (CO<sub>2</sub>, CH<sub>4</sub> and N<sub>2</sub>O) in boreal areas are understood (Yang et al. 2015; Wik et al.  
344 2016; Valiente 2022). However, the question of whether these feedbacks can result in tipping  
345 points by becoming self-sustaining beyond a threshold is not yet settled. Most boreal lakes are  
346 net heterotrophic and thus conduits of CO<sub>2</sub>, often also CH<sub>4</sub>, due to high concentrations of DOM  
347 and common deep-water anoxia. A shift from net autotrophy to net heterotrophy would classify

348 as a binary shift, yet with a strong, positive climate feedback. If it eventually leads to oxygen  
349 depletion and cascading feedbacks then it would qualify as a tipping point. However, there  
350 would be a time delay between the two events, with the latter being the critical tipping event.  
351 There is also a commonly reported unimodal response in lakes to increased loadings of DOM,  
352 typically around 5 mg DOC l<sup>-1</sup> (Karlsson et al. 2007; Thrane et al 2014), where increases in  
353 DOM below the threshold may promote NPP and thus CO<sub>2</sub> drawdown due to N and P associated  
354 with DOM, while reduced NPP and increased degassing of CO<sub>2</sub> (and CH<sub>4</sub>) will take place above.  
355 We thus propose two types of large-scale potential tipping points, one related to anoxia, the other  
356 to DOM-concentrations, yet both are related to increasing load of terrestrially derived DOM  
357 across the boreal region.

358

### 359 **2.3. Disappearance/appearance of waterbodies**

360 A global reduction in lake water storage (Yao et al., 2023), and the climate-driven creation or  
361 disappearance of water bodies is a crucial issue. Loss of water-bodies due to overuse, warming  
362 or drought pose a major threat to vulnerable, freshwater resources, also by deteriorating water  
363 quality or salinization (cf. below). The most dramatic warming has already taken place in the  
364 high Arctic with temperature increases up to 3 °C over the past few decades (Wang et al. 2022),  
365 further promoting the onset of permafrost thaw (Langer et al. 2016). Current and future  
366 permafrost thaw and glacier melting can both create new waterbodies and drain old, providing a  
367 strong link to the fate of the cryosphere (Smith et al. 2005; Olefeldt et al. 2021). Such small, but  
368 numerous waterbodies residing on permafrost over large geographical scales in Eurasia and  
369 North-America are currently among the most vulnerable water-bodies globally (Smol and  
370 Douglas 2007; Heino et al. 2020). They host species-poor but specific communities of  
371 invertebrates (Rautio et al. 2011; Walseng et al. 2021) of vital importance for birdlife and other  
372 biota. Warming may also affect these waterbodies indirectly via glacier melt, increased inputs of  
373 organic C, fertilisation by increasing populations of geese (caused by climate change), and  
374 consequently changes in microbial communities and increased GHG emissions (Eiler et al.  
375 2023). Thus, by their sheer number these systems may also serve as increasingly important  
376 conduits of GHGs and historical soil carbon stocks to the atmosphere (Laurion et al. 2010;  
377 Negandhi et al. 2016) and play an important role in mediating nutrient delivery to the polar  
378 oceans (Emmertson et al., 2008), potentially affecting global NPP (Terhaar et al., 2021). While

379 the main problem is loss of water bodies affected by warming-induced increased  
380 evapotranspiration rates (Smol and Douglas 2007) and permafrost thaw (Smith et al. 2005), there  
381 are also cases where collapsing palsas and thermokarst areas create new waterbodies, and these  
382 waterbodies may themselves represent a positive feedback by accelerating the thaw (Langer et al.  
383 2016; Turetsky et al., 2020).

384 Since most of these potentially lost waterbodies are small and nameless ponds, it is hard  
385 to point to specific cases, but the works cited above provide a number of telling examples. While  
386 the focus in this context is negative water balance or loss of high-latitude waterbodies, this is  
387 actually a widespread problem causing shrinking of many lakes. In Arctic areas, responses to  
388 warming may differ substantially between perennial lakes and ephemeral wetlands, related to  
389 ambient temperature and permafrost depth (Vulis et al. 2021). Although loss or gain of water  
390 bodies does not classify as a tipping event in the very strict sense, i.e. there is not obvious strong,  
391 self-reinforcing factors involved, it still is a climate driven change with potentially large,  
392 widespread and irreversible consequences, and with repercussions on climate in terms of changes  
393 GHG-emissions from vast areas.

394

#### 395 *Feedbacks and tipping points*

396 Some essential feedbacks to climate change are involved in the change of Arctic waterbodies;  
397 e.g. reduced ice and snow cover in the Arctic will promote further permafrost thaw. Certain  
398 Arctic areas have experienced a major increase in breeding birds, notably gees, that promote  
399 increased loadings of organic C and nutrients (Hessen et al. 2017). More organic carbon entering  
400 water bodies from their terrestrial surroundings, combined with warming and eventually bird  
401 induced eutrophication promotes GHG emissions (Wei et al. 2023). It is important to make clear  
402 that some of the impacts are contrasting, i.e. the loss of waterbodies may at first increase GHG  
403 emissions (Keller et al. 2020; Paranaiba et al. 2021) but will eventually reduce GHG emissions.  
404 Permafrost thaw and drainage of water-logged areas will increase CO<sub>2</sub>-emissions but could  
405 reduce CH<sub>4</sub>-emissions. Sudden release of methane-hydrates upon permafrost thaw is a  
406 possibility, yet hard to predict and quantify, and not specifically linked to aquatic habitats.

407 Few changes are as irreversible as complete habitat loss, and the climate-driven loss of  
408 numerous water-bodies residing on permafrost over large geographical scales in Eurasia and  
409 North America (due to permafrost thaw) with climate feedbacks in terms of changed GHG-

410 emissions is possible. In fact, as argued by Smol and Douglas (2007); “The final ecological  
411 threshold for these aquatic ecosystems has now been crossed: complete desiccation”. If strictly  
412 adhering to the tipping point criteria as an event occurring when self-sustaining change in a  
413 system is triggered beyond a forcing threshold, typically starting with positive feedback loops  
414 and a runaway phase before finally the tipping-point brings the system into a different alternative  
415 state, loss of waterbodies is not strictly a tipping point, but a binary shift. Abrupt permafrost  
416 thaw, which can drive abrupt self-sustained formation or draining of thermokarst lakes, is  
417 categorised as a “regional impact” climate tipping element by Armstrong McKay et al. (2022).  
418 We extend this categorisation to include the lakes associated with these abrupt thaw processes,  
419 seeing them as a coupled permafrost-lake systems with tipping dynamics involving both  
420 components (Turetsky et al., 2020). Despite the scale considered here, the extent of open water  
421 globally is relatively easy to quantify using remote sensing, and it is possible to make predictions  
422 based on time-series and empirical relationships between temperature increase, permafrost thaw  
423 and loss of water-bodies. Quantifying potential climate feedbacks related to processing of  
424 organic C to CO<sub>2</sub> and CH<sub>4</sub> should be possible to predict within orders of magnitude, with initial  
425 analysis suggesting abrupt thaw involving thermokarst lake formation and draining could double  
426 the warming impact of gradual permafrost thaw (Turetsky et al., 2020).

427

#### 428 **2.4. Switch from N to P-limitation**

429 Imbalance in biogeochemical cycles has become a major concern both on the local and global  
430 scale. Anthropogenic emissions of CO<sub>2</sub> now appear as the major environmental challenge for  
431 ecosystems and human well-being in the foreseeable future. In relative terms, however, the  
432 anthropogenic effects on the global N-cycle are even more pronounced. Transformation of  
433 atmospheric N<sub>2</sub> to more reactive reduced or oxidized forms of inorganic N by the fertilizer  
434 industry and combustion processes has dramatically changed. Recent analyses of the global N-  
435 cycle (Bodirsky et al. 2014; Zhang et al. 2020) suggest that various human activities currently  
436 convert similar N<sub>2</sub> to total natural ecosystem fixation, and that both the use of N and P are far  
437 beyond “safe boundaries” (Rockström et al. 2023).

438 Increased N-deposition may affect surface waters in fundamentally different ways. It will  
439 increase the emissions of N<sub>2</sub>O (Yang et al. 2015), and increased deposition of inorganic N  
440 promotes soil and water acidification through increased NO<sub>3</sub> in surface waters (Stoddard 1994).



441 It will, however, also affect elemental ratios in lakes and rivers (Hessen et al. 2009). The relative  
442 proportions of these elements will determine the nature of elemental limitation for both  
443 autotrophs and a range of heterotrophs, and could thus profoundly affect community composition  
444 and ecosystem processes. One effect of such skewed inputs of N over P would be an intensified  
445 P-limitation in surface waters or even large-scale shifts from N to P-limitation (Elser et al. 2009).  
446 Rather than pinpointing specific lakes as examples, contrasting areas with high vs low N-  
447 deposition Elser et al (2009) provide good regional examples, where Colorado offers examples  
448 of regions with either high or low N-deposition, while southern regions of Norway and Sweden  
449 experience up to ten-fold elevated levels of N-deposition compared with central regions.

450         Conversely, increased N-loss by denitrification, eventually associated with increased  
451 internal P-loading may shift systems from P to N-limitation (Weyhenmeyer et al. 2007). Societal  
452 implications include an increased prevalence of toxin producing cyanobacteria, purported to be  
453 promoted in extent by warming (Paerl et al., 2008) and favouring non-N-fixing toxin producing  
454 species where reduced-N concentrations are high relative to oxidized-N (Hoffman et al., 2022).  
455 Additionally, a threshold on toxic effects on sensitive freshwater species has been proposed (i.e.  
456 2 mg L<sup>-1</sup>; Camargo et al., 2006; Moss et al., 2013), above which a marked decline in biodiversity  
457 is expected.

458

459 *Feedbacks and tipping points:*

460 Changes in N- versus P-limitation of NPP are associated with changes in community structure,  
461 both for the phytoplankton and macrophyte communities. While the shift from one limiting  
462 nutrient to another represents a binary shift and abrupt transition, it is not driven by self-  
463 propelling events or positive or negative feedbacks, since a shift from N- to P-limitation typically  
464 is caused by N-deposition or agricultural use of fertilizers. While increased N-loading per se  
465 could promote climate feedbacks in terms of N<sub>2</sub>O, the switch from N to P-limitation or vice  
466 versa is neither driven by climate nor does it exhibit strong feedbacks on climate. There is also  
467 no inherent hysteresis, and when drivers change the system may immediately return to the other  
468 limiting nutrient. For these reasons we do not classify this category as a tipping point according  
469 to the definition above.

470

471       **2.5. Salinization**

472 Salinization is a prevalent threat to freshwater rivers, lakes and wetlands world-wide, particularly  
473 in arid and semi-arid regions and coastal areas. It is caused by a range of anthropogenic actions  
474 including water extraction, pollution and climate change (Herbert et al. 2015). The causes of  
475 salinization have historically been classified as being primary or secondary. Primary salinization  
476 refers to natural causes including wet and dry deposition of marine salts, weathering of rocks and  
477 surface or groundwater flows transporting salts from geological salt deposits. Secondary  
478 salinization refers to salinization caused by human activities such as irrigation with water rich in  
479 salts, rising of brackish and saline groundwater due to increased ground water extraction and  
480 increased seawater intrusion as a result of sea level rise. The distinction between natural and  
481 anthropogenic causes underlying salinization is becoming less clear cut due to climate change as  
482 anthropogenically caused changes in temperature, precipitation patterns and wind will affect the  
483 primary salinization processes (Oppenheimer et al. 2019). Salinization has severe consequences  
484 for aquatic communities (Jeppesen et al. 2015; Short et al. 2016; Cunillera-Montcusí et al. 2022).  
485 Salinization has a strong ecological impact often associated with osmotic stress and changes in  
486 biogeochemical cycles which often entails an increase in concentration of toxic sulfides (Herbert  
487 et al. 2015). In addition, studies focussing on the application of road salts indicate that  
488 salinization may disrupt lake water mixing and release of metals (Szkłarek et al. 2022 and  
489 references therein). Negative effects of increased salinity have been described for trophic levels  
490 ranging from microorganisms to fish and birds (reviewed by Cunillera-Montcusí et al. 2022). In  
491 addition, salinization also has a high societal impact particularly related to domestic and  
492 agriculture water supply in arid and semi-arid regions (Williams et al. 1999).

493       A strong example on climate driven salinisation and its impact on biota is the long-term  
494 study (1938 – 2004) in two Canadian lakes (Sereda et al. 2011). Concomitant with periodic  
495 declines in precipitation, lake elevation declined and salinity increased in Jackfish and Murray  
496 lakes from 1938 to 2004. The increase in salinity caused an estimated 30% loss in diversity of  
497 macrobenthos. If salinity exceed thresholds where key species or functional groups are wiped  
498 out, it will no doubt represent an abrupt ecosystem transition, yet still not a tipping point in the  
499 sense that it is self-propelled.

500

501 *Feedbacks and tipping points*

502 Regime shift from clear to turbid may occur at 6-8 per mil salinity in systems with intermediate  
503 to high nutrient loadings and have been associated with a change in zooplankton community  
504 composition from cladocerans to more salinity tolerant cyclopoid copepods (Jeppesen et al  
505 2007). Salinity induced regime shifts may also lead to dominance by microbial mats at the  
506 expense of submerged macrophytes (Davis et al. 2003, Sim et al. 2006). While there are species-  
507 specific tolerance thresholds to salinity, and these effects are expected to interact with other  
508 stressors - including eutrophication (Jeppesen et al. 2007, Kaijser et al. 2019), color and turbidity  
509 (Davis et al. 2003) - the process is not driven by feedbacks of increased salinization, but external  
510 factors like warming, water (over)use and road salting. Hysteresis after refreshing of salinized  
511 systems has been little studied but is likely strongly biogeochemical in nature as evidenced by  
512 previously brackish waters that have been flushed with freshwater for over 90 years and still  
513 contain high levels of chloride, sodium and sulfate (Van Dijk et al. 2019).

514 A weakened top-down control by zooplankton on phytoplankton occurring at moderate  
515 high salinities would be an indirect consequence of salinization, leading to a worsening of  
516 eutrophication symptoms (Gutierrez et al. 2018) and thus promoting indirect climate effects.  
517 Salinization however tends to decrease CH<sub>4</sub> emissions (Herbert et al. 2015, Chamberlain et al.  
518 2020, Gremmen et al. 2022). The decrease in CH<sub>4</sub> emission can be either caused by a decrease in  
519 CH<sub>4</sub> production - e.g. because methanogens are outcompeted by sulfate reducers or are  
520 negatively impacted by sulfide toxicity - or because an increase in methane oxidation (reviewed  
521 by Herbert et al. 2015). The salinity induced decrease in aquatic CH<sub>4</sub> emissions may imply a  
522 negative feedback with climate change, but only when this is not off-set by a decrease in carbon  
523 burial. Insight in this balance is currently limited (Chamberlain et al. 2020), and while no doubt  
524 salinization is widespread on regional scales and may reach threshold values for species and  
525 processes, we do not categorize it as a tipping point under the cited criteria.

526

## 527 2.6. Spread of invasive species

528 Freshwaters are especially vulnerable to species loss and population declines as well as species  
529 invasions due to their constrained spatial extent. Substantial ecosystem changes by reinforcing  
530 interactions between invasive species and alternative states (i.e. macrophyte *versus*  
531 phytoplankton dominance, as described above) may occur (Reynolds and Aldridge 2021). The  
532 spread of several invasive species can change community composition and ecological functions

533 in dramatic ways, and can be regarded as sudden transitions with major site-specific or regional  
534 impacts. Moreover, species invasions can be facilitated by climate change (Rahel and Olden,  
535 2008), and notably flooding and other hydrological events can facilitate species invasion with  
536 potentially far-reaching ecological consequences (Anufriieva and Shadrin 2018).

537         There are numerous examples of ecological consequences in lakes following species  
538 invasions, and the major impacts of invasions by zebra mussel as well as the predatory  
539 cladoceran *Bythotrephes* in the Great lakes, serve as striking examples of major impacts at the  
540 regional scale even in very large lakes (Ricciardi and MacIsaac 2000). While species invasions  
541 are of major ecological and societal concern, and can induce ecological tipping points in certain  
542 lakes, they are generally not self-perpetuating involving internal feedbacks. No doubt it may be  
543 appropriate to say that invaded system may cause irreversible changes or hysteresis in specific  
544 lakes or lakes within regions.

545

#### 546 *Feedbacks and tipping points*

547 Climate, both in the context of warming that open for latitudinal and altitudinal spread of species  
548 (Hessen et. al. 2006) and hydrological events that likewise may promote invasions (Anufriieva  
549 and Shadrin 2018) may pose drastic changes in community composition and ecosystem functions  
550 to an extent that qualify as abrupt shifts. Species invasions may also interact with other drivers  
551 lowering the potential thresholds (of nutrients, temperature, browning, etc.) for a shift to occur,  
552 and vice versa, by impacting on previously occurring stabilizing mechanisms (Willcock et al.  
553 2023). Likewise, species shifts may have repercussions on GHG-emissions. We do not pursue  
554 the discussion feedbacks and potential tipping points further for this candidate category,  
555 however, since we have constrained our definition of tipping points to situations with internal  
556 feedback and regional occurrence. Given the widespread anthropogenic changes promoting  
557 invasive species in aquatic communities worldwide, the often abrupt and unpredictable shifts that  
558 may follow from this deserves further attention.

### 559 **3. Discussion**

560 Freshwaters are one of the most vulnerable ecosystems and resources globally and will  
561 increasingly be so in warming world. They also link catchment properties and terrestrial changes  
562 to marine systems, and notably lakes serve as good sentinels of global change (Adrian et al.

563 2009). Population declines and species loss in freshwaters are happening at an alarming pace  
564 underpinning the urgency for evidence of ecological tipping points in response to environmental  
565 change. Drinkable freshwater is a scarce resource both in terms of water quality and availability  
566 (Yao et al. 2023). Predicting (and preventing) sudden shifts in water quality and quantity is  
567 therefore a high priority also from an anthropocentric perspective, and insights into feedbacks,  
568 thresholds and tipping points are highly relevant to lakes. Lakes are also major players in the  
569 global climate, and besides being highly vulnerable to climate change, they can provide strong  
570 feedback to the climate by ventilating a substantial share of terrestrially fixed C back to the  
571 atmosphere as CO<sub>2</sub> and CH<sub>4</sub> (Cole et al. 2007; Tranvik et al. 2009; Raymond et al. 2013). Lakes  
572 are also subject to changes, sometimes sudden, due to climate change and other natural or  
573 anthropogenic drivers. In fact, some of the first and most striking examples on tipping points and  
574 regime shift come from lake studies (Scheffer et al. 1993; Jeppesen et al. 1998).

575 We argue that there are two key drivers that may shift lakes towards major ecological  
576 changes, as well as increased climate feedback by GHG emissions, namely eutrophication and  
577 browning. Both these drivers are promoted by warming, which may be seen as a separate driver.  
578 Both processes are also characterised to some degree by self-sustaining feedback loops, feedback  
579 to climate in terms of GHG-emissions, and are also strongly integrated with land surface impacts  
580 in the catchment (Fig. 3). Warming, browning, and eutrophication lead to increases in  
581 stratification, heterotrophy, and phytoplankton or macrophyte mass, which collectively drive  
582 benthic oxygen depletion and in turn increased GHG emissions (helping to drive further  
583 warming and DOM loading from land) and internal P loading (driving further eutrophication)  
584 (Meerhoff et al. 2022). Several of these processes can feature tipping points (eutrophication and  
585 potentially DOM loading), which warming will likely make easier to reach.

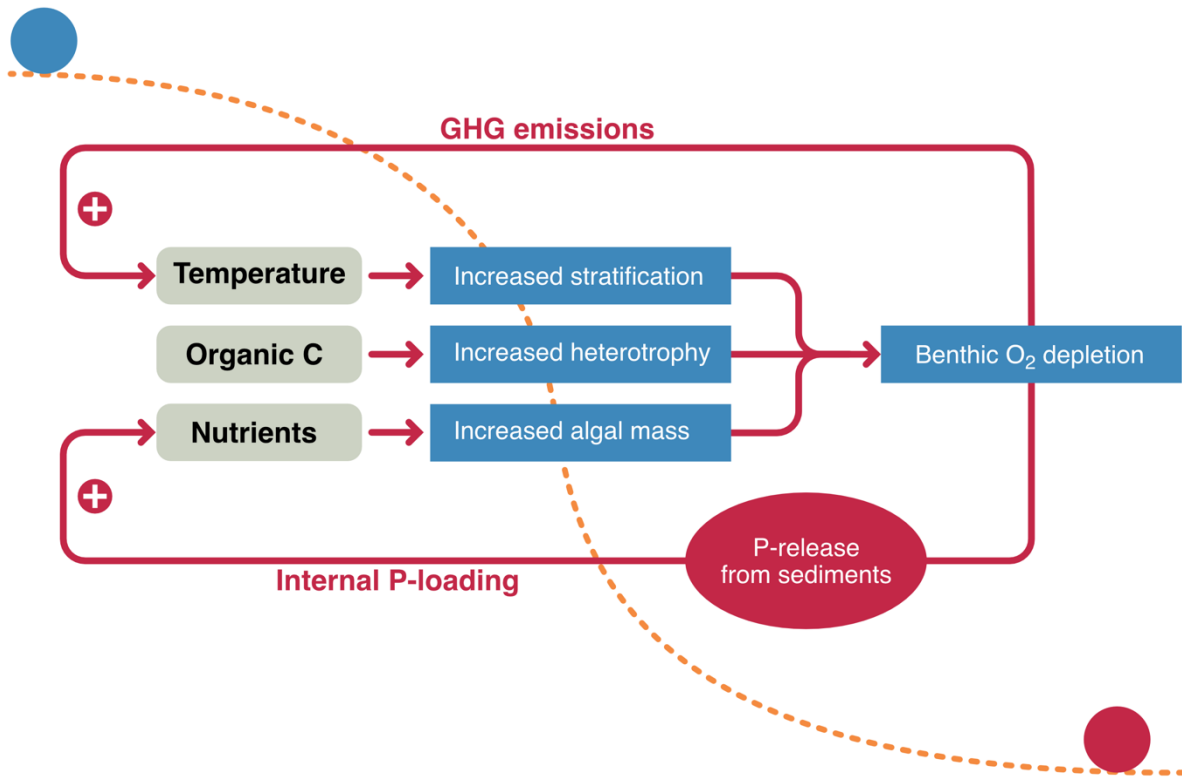
586 Few processes have been more thoroughly described in terms of drivers, impacts and  
587 remedies than freshwater eutrophication. The drivers are well known (nutrient loadings from  
588 agricultural activities and locally also from sewage discharges), despite long-term controversies  
589 regarding the relative importance of N or P in promoting eutrophication (e.g., Smith & Schindler  
590 2009, Paerl et al. 2016). There are also long traditions for predictive hydraulic models that link  
591 the load of P to algal blooms and benthic O<sub>2</sub>-depletion (e.g., Vollenweider type models, Imboden  
592 1974). Moreover, given the scarcity, increasing demands and increasing prices of P as a  
593 commodity worldwide, there are indeed strong arguments to close the loop for P and reduce

594 losses to the environment (Brownlie et al. 2022). Due to the strong impact of O<sub>2</sub>-depletion on  
595 sediment release of P and thus internal fertilization (Soendergaard et al. 2002), that will play in  
596 concert with food-web driven feedbacks (cf. Fig. 3), tipping points in this context can be  
597 identified, while the climate component is difficult to separate.

598         Browning shares many of these attributes in terms of increased net heterotrophy. The  
599 shift from net autotrophy with a net uptake of CO<sub>2</sub> to net heterotrophy with a net release of CO<sub>2</sub>  
600 (plus CH<sub>4</sub>) also represents a binary situation. However, since most boreal lakes are already net  
601 heterotrophic owing to microbial conversion of organic C (Hessen et al. 1990; Cole et al. 1994;  
602 Larsen et al. 2011), most boreal lakes simply become more heterotrophic, hence there is no  
603 tipping point in this context. However, an increased degree of heterotrophy combined with  
604 increased thermal stability, will promote deep-water anoxia, thereby internal P-cycling and  
605 GHG-release. Since the key driver here is the external load of terrestrial DOM, the feedback  
606 component driving P-release is weaker than in the case of eutrophication. Nevertheless, an  
607 increased release of GHGs no doubt poses a feedback to the climate and hence the terrestrial  
608 systems that may promote further browning. These processes are amplified by climate change,  
609 and have global consequences in terms of GHG emissions. Given the high confidence in this  
610 case we recommend it as apriority for parametrization of models to underpin future predictions  
611 of the impacts of global tipping points in lakes on GHG emissions.

612

613



614  
 615 Fig. 3. The interactive role of eutrophication, DOM-export (browning) and warming on lakes.  
 616 Separately or combined they promote benthic O<sub>2</sub>-depletions which cause an internal feedback by  
 617 P-loading from sediments and a climate feedback via release of greenhouse gases. The potential  
 618 shift between states (blue to red circle) is indicated.  
 619

620  
 621 As a separate type of binary tipping point which is likely to be widespread and related to  
 622 GHG-release, we propose the loss of water bodies, notably Arctic ponds. This is driven by  
 623 permafrost thaw in the case of thermokarst-linked lake formation or disappearance (categorised  
 624 as a regional tipping element in previous assessments (Armstrong McKay et al. 2022). Together,  
 625 the coupled permafrost-lake system can act as a localised tipping system with the lake providing  
 626 key feedbacks to help drive self-sustaining thaw. This makes the tipping points easy to monitor  
 627 (by remote sensing), and predictable in the sense that it will be closely linked to permafrost thaw.  
 628 There are, however, feedbacks to the climate, with potentially high emissions during the drying  
 629 process (Marcé et al. 2019; Turetsky et al. 2020). A different situation would be the less  
 630 widespread case of new waterbodies formed by collapsing palsas, in cases of retreating glaciers.  
 631 Given the potential scale of occurrence, the net effect of permafrost thaw and increased release

632 of CO<sub>2</sub> balanced with the effect of disappearing waterbodies and potential changes in net GHG  
633 emissions requires further attention as a matter of high priority.

634

635 *Gradients or tipping points – does it matter?*

636 One could argue that what matters is whether a change or process is linear (and thus more  
637 predictable) or non-linear (and less predictable), and that the rest is semantics. This is truly not  
638 the case, since there are substantial differences in what is considered as a ‘tipping point’, not the  
639 least in terms of whether impacts are easily reversible or are effectively “locked in” (e.g.  
640 hysteresis). Still, from an ecosystem perspective, abrupt shifts, even if they do not qualify as  
641 tipping points, may have devastating effects that should urge us to invest more in preventing  
642 deterioration as we do not know where/if a sudden shift may occur. As argued by Moss et al.  
643 (2008): "the sort of precision demanded by legislators and lobbies will never be attainable and  
644 this has been a major weapon used to delay regulation of agricultural activities."

645         Shifts between ecological states do not necessarily involve alternative stable states with  
646 hysteresis. In fact, both the concepts of abruptness and irreversibility depends on one’s  
647 perspective of time. Over a lakes life-time, shifts back and forth between states occurring over  
648 years or even decades are “sudden” relative to the human lifespan. For example, Rühland et al.  
649 (2008) report apparent coherence in diatom community shifts post 1850 on hemispheric scales  
650 over 100 years or so. Similarly, a coherent, global increase in hypoxia in lakes has been reported  
651 over a 100 year period (from about 1850) by Jenny et al. (2016). Likewise regional patterns of  
652 species turnover ( $\beta$ -diversity) over 200 years demonstrated regional differences in species  
653 turnover, but also recent changes attributed to warming (Kahlert et al. 2020). As warming  
654 progresses, such studies forms good baselines for future changes. If the observational time step is  
655 increased to centuries, then it is likely that more large-scale examples will come through in  
656 paleo-studies. In fact, there are several examples on coherence in lake responses to climate  
657 variability or climate change, some of which take place over short time spans (Stone et al. 2016;  
658 Isles et al. 2023). Finally, multiple drivers may jointly push lakes towards shifts or tipping  
659 points, as shown in Huang et al. (2022) and Willcock et al. (2023).

660         Taken together, there are at least two major reasons why an improved understanding of  
661 sudden changes in lake ecosystems are imperative; they are highly vulnerable to climate change  
662 and other anthropogenic stressors globally, and they serve as major feedbacks to the climate



663 system through GHG emissions. Being well-mixed and semi-closed entities that reflect changes  
664 in catchment properties, they also serve as sentinels of global change (Adrian et al. 2019). For  
665 fresh waters in general, lakes are crucial in the hydrological cycling, and link the terrestrial and  
666 marine ecosystems. The major tipping point dynamics converge in oxygen depletion, primarily  
667 in deeper strata and the sediment surface, which promotes feedbacks and hysteresis in terms of  
668 internal P release as well as increased GHG-emissions. High nutrient load, increased inputs of  
669 dissolved organic C and warming all drive oxygen depletion, and while many problems related  
670 to global warming boil down to the obvious recommendation of reduced use of fossil fuel and  
671 other GHG-emitting activities, reducing nutrient use and losses to within the carrying capacity of  
672 the system (i.e. from ecosystem to global scales) is comparatively simpler both for N and P when  
673 compared to C (Rockström et al. 2009; 2023). The incentives should be even greater for closing  
674 the P-loop, given the potential for scarcity of this non-substitutable element and its role in lake  
675 eutrophication (Brownlie et al. 2022).

676         Regime shifts and tipping points are concepts closely linked to resilience (Andersen et al.  
677 2008; Spears et al. 2017). Lakes represent excellent model case studies in this respect and have  
678 been used widely to demonstrate theories of ecological stability and resilience that are needed to  
679 underpin preventative management approaches and to guide science-based environmental policy.  
680 The full importance of the vulnerability of lakes to climate and other anthropogenic impacts, as  
681 well as their feedback to climate is not yet fully acknowledged, so there is a need both for  
682 science and communication in this regard. However, we argue that the search for empirical  
683 evidence to underpin theory should not prevent societies and managers taking more action to  
684 protect fresh waters in the meantime.

685

#### 686         **4. Conclusions**

687 Anthropogenic forcing may induce non-linear, abrupt changes in freshwater ecosystems, and  
688 awareness of such potential threshold effects are important. Here we focus on lake and pond  
689 systems that may be subject to tipping points, where self-reinforcing feedback and some type of  
690 post-change hysteresis can be identified. To qualify in this context these changes must also be  
691 relevant or larger scales (i.e. regions or biomes), or a large number of systems. Two types of  
692 potential tipping points were identified based on these criteria; eutrophication and browning. The  
693 first is an example of a widespread phenomenon, the second occurring in lakes in the boreal

694 biome. In both cases, climate is involved as a driver, and the changes in terms of deep-water  
695 anoxia and internal P-cycling may boost the emission of GHG-gases from such systems. While  
696 not tipping points according to the criteria applied here, two types of binary shifts is also  
697 discussed; loss of water-bodies, notably in the Arctic areas, caused by permafrost thaw or  
698 negative water balance as well as shifts between N and P-limitation caused by N-deposition.  
699 Notably the first is driven by climate change and will also have repercussions on climate by  
700 changes in GHG-emissions. Finally, salinization and species invasions are also changes that may  
701 occur over large scales with potentially abrupt ecosystem changes, and where changes in  
702 temperature and precipitation as important drivers.

703

704 *Acknowledgements:* This work has benefitted from discussions withing the group behind the  
705 Tipping Point report presented at COP 28 meeting in Dubai, and the ms has benefitted  
706 substantially from inputs and suggestions by John Smol and an anonymous reviewer on the first  
707 draft.

708

709 *Author contribution:* DH conceived the idea and wrote the first draft, all authors have contributed  
710 to the writing and approved the final manuscript.

711 *Code and data availability:* Not relevant

712 *Competing interests:* None.

713 *Special issue statement:* Part of the ESD tipping points special issue

714

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