Lake ecosystem tipping points and climate feedbacks

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Abstract

Lakes experience anthropogenically-forced changes that may initiate ecosystem feedbacks, in some cases reaching tipping points beyond which impacts become hard to reverse. Lakes are also important players in the global climate by ventilating a large share of terrestrial carbon back to the atmosphere as greenhouse gases, and will likely provide substantial feedbacks to climate change. In this paper we address various major changes in lake ecosystems, and discuss if tipping points can be identified, predicted, or prevented in them, along with their associated feedbacks to climate change. Potential tipping dynamics assessed include eutrophication-driven anoxia and internal phosphorus-loading, increased loading of organic matter from terrestrial to lake ecosystems (lake “browning”), lake formation or disappearance in response to cryosphere shifts, switching from nitrogen to phosphorus limitation, salinization, and the spread of invasive species. We also address other types of abrupt, or threshold-type shifts in lakes and ponds, and conclude on which tipping points are locally or regionally relevant. We identify a key set of co-drivers that could lead to self-sustaining feedbacks, with warming, browning, and eutrophication leading to increased lake stratification, heterotrophy, and algal mass, which separately or collectively drive benthic oxygen depletion and in turn increased greenhouse gas emissions (helping to drive further warming and organic matter loading) and internal phosphorus-loading (driving further eutrophication). Several of these processes can feature tipping points, which further warming will likely make easier to reach. We argue that the full importance of the vulnerability of lakes to climate and other anthropogenic impacts, as well as their feedback to climate is not yet fully acknowledged, so there is a need both for science and communication in this regard.
Introduction

In natural sciences, the hysteretic behaviour of lakes (Scheffer et al. 2007) has informed the concept of tipping points at the ecosystem level, following the development of the alternative stable states theory in shallow lakes (Scheffer et al. 1993). Given the global vulnerability of freshwaters and the pervasive nature of major pressures acting upon them (e.g. nutrient pollution, over-extraction, and climate change), tipping points in these systems could have significant societal impacts, including on human and environmental health, food production, and climate regulation. The capacity to detect discontinuous ecosystem responses to pressure changes in natural systems has been challenged (e.g. Hillebrand et al. 2020; Davidson et al. 2023).

Nevertheless, there are several studies that have reported the occurrence of tipping points even if they are difficult to detect (Lade et al, 2021), such as shifts from one alternative state to another in small shallow lakes, the most populous lake type globally (Messager et al., 2016).

Widespread loss of water-bodies, from Arctic or sub-arctic ponds to wetlands or bogs might qualify as one type of tipping point, but are not self-propelled by internal feedbacks themselves rather than by permafrost thaw (Smol and Douglas 2007). The question of “sudden” system shifts, alternative stable states and hysteresis depends too on what is considered a relevant time span; days, years, decades or centuries. Also, systems may have alternative states that are not necessarily fixed over long time-spans, hence the phrase “stable” should be used with caution, just like there are strong and weak hysteresis. Uncertainty also remains on the geographical extent of tipping points in lakes and the wider relevance for the Earth’s climate system – we here focus on potential tipping points of global or regional relevance, and with relevance to global change.

Empirical analyses, process modelling and experimental studies are advanced for shallow lakes, providing a good understanding of lake ecosystem behaviour around tipping points. There are related concepts in the literature (regime shifts, catastrophic shifts, forward switches, etc.), but here we adopt the definition of a tipping point occurring when self-sustaining change in a system is triggered beyond a forcing threshold, typically starting with positive feedback loops, then entering a runaway phase before finally the tipping-point brings the system into a different alternative state (Nes et al. 2016). For example, the well documented increase of phosphorus (P) loading across European lakes in the last century (e.g. from agricultural and waste water pollution) has uncovered critical loading thresholds beyond which lakes can shift rapidly from a
clear water, submerged macrophyte rich state to a turbid, phytoplankton dominated state (Scheffer et al., 2001; Jeppesen et al., 2005; Tárai et al. 2008), and vice versa, when nutrient loading decreases. One of the theoretical implications is that to induce a switch back to the initial state the nutrient loading should be reduced to a lower threshold before the shift might be possible (hysteresis). Adding to such well-described and mechanistically well understood changes, there is a wide range of local or single lake shifts that may be categorized as tipping points. The question remains as to whether tipping points are merely isolated phenomena in single lakes, or specific types of lakes, or whether they are, or may be in the future, manifest

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

across geographically distinct populations of lakes experiencing similar environmental change, with the potential for regional or global extent (Fig. 1).

It is well established that lakes are sensitive to the effects of climate change, including warming and changes in precipitation and storminess (e.g., Adrian et al., 2009; Meerhoff et al., 2022). Emerging evidence suggests that lakes and ponds may also play an important role in climate regulation, through both the emission of greenhouse gases (i.e. predominantly CH$_4$, Downing et al., 2021) and carbon burial (Anderson et al., 2020). Lakes and rivers are impacted by climate change and other anthropogenic pressures globally, but they also provide strong feedbacks to the global climate systems and carbon (C) cycle, (Cole et al. 2007; Tranvik et al. 2009), despite comprising a small part of global water extent While global estimates of net greenhouse gas (GHGs) emissions from lakes remain poorly constrained, there is general consensus that a significant fraction of terrestrially fixed C is degassed to the atmosphere via surface waters. Cole et al. (2007) conservatively estimated that inland waters annually receive some 1.9 Pg C yr$^{-1}$ from the terrestrial landscape, of which at least 0.8 Pg C yr$^{-1}$ is returned to the atmosphere through water to atmosphere GHG exchange. Later estimates revised this global GHG exchange term, to include evasion rates, at 2.1 Pg C yr$^{-1}$, from lakes, rivers and reservoirs (Raymond et al. 2013). Notably, boreal lakes are important conduits of CO$_2$ release to the atmosphere, estimated to be equivalent to the annual CO$_2$ release from forest fires, globally (Hastie et al. 2017). Under a high CO$_2$-emission scenario and as a result of increased terrestrial NPP, CO$_2$ emissions from boreal lakes are projected to increase by 107%, showing the coupling between the terrestrial and aquatic C cycle (Hastie et al. 2017).

This significant role of surface waters for GHG-emissions is also highly relevant, but poorly constrained both in national and global C-budgets (Lindroth and Tranvik 2021). The balance between inputs of organic C and nutrients is a key determinant of the balance between heterotrophic and autotrophic processes, and thus not only determine the biodiversity, community composition and food web structure, but also the productivity-to-respiration (P:R) ratio. And so, it is relevant to consider the extent to which potential tipping points may drive, or be driven by, climate change, leading to higher level feedbacks to the Earth’s climate system.

Here, we discuss tipping points in freshwaters reported in the literature, focusing on lakes and ponds, with the potential for global or at least regional or biome-scale relevance. In this context we will constrain the discussion to potential tipping points that are more generic, at least
with some regional or biome-wise impact, and that could have feedbacks to the climate, while not necessarily being driven or triggered by climate change per se. We identify 6 candidate categories for tipping points at a relevant scale in this context (regional to global impact), and for each of the categories we discuss whether observed changes can be categorised as tipping points according to the definition above. We also address climatic and other drivers and consequences, including potential feedbacks to the climate system, and wider societal implications, with emphasis on the most relevant and influential categories.

**Candidates and categories of lake tipping points**

In principle many abrupt or sudden changes imposed on a waterbody could result in specific impacts, i.e. toxic waste or toxic treatments (e.g. rotenone to kill off undesired species; runoff of herbicides inadvertently killing aquatic plants), hydrological alterations by impoundment or canals, and stocking of new (often exotic) species. To qualify as tipping point, there should be self-sustaining dynamics and positive feedbacks involved, and to be relevant in a wider context, the tipping point should be more generic to certain types of impact, certain types of waterbodies, and potentially also have feedbacks to the climate in terms of GHG-emissions. We have identified 6 stressors that may trigger a freshwater ecosystem to cross a tipping point (Table 1), and scrutinise them one by one.

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

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

Eutrophication is one of if not the most pervasive impacts on fresh waters and coastal systems. Although it may naturally occur due to inputs from the watershed or from biota translocating nutrients from connecting ecosystems, eutrophication is a largely human-induced phenomenon. The main causes of cultural eutrophication have varied across time and regions. However, it is widely accepted that the main current cause of eutrophication is the change in land use in the watersheds and particularly agricultural activities acting as diffuse sources of nutrients (among other agrochemicals) (Moss 2008; Schulte-Uebbing et al., 2022). Agriculture, with myriad impacts on fresh waters that go well beyond nutrient pollution (Moss, 2008), has been pointed as a major driver of ecosystem shifts and tipping points (Gordon et al., 2008).

Fig. 2. Feedback loop diagram for eutrophication, demonstrating key feedbacks that can amplify P-loading (and beyond a tipping point drive self-sustaining change) and drive increased greenhouse gas emissions.
The mobilisation of P from sediments, a process known as internal loading (Sondergaard et al., 2001), plays a key role in hysteresis in lakes recovering from cultural eutrophication (Boström et al. 1982; Jeppesen et al. 1991; Spears & Steinman 2020). Eutrophication-driven changes in biota, such as changes in fish composition and size structure with cascading effects on zooplankton and phytoplankton as well as strong impacts if fish mediated nutrient cycling (Brabrand et al. 1990), also strengthen hysteresis and maintain a system with deepwater anoxia and high nutrient load, supporting the release of GHGs (Fig. 2).

Feedbacks and Tipping points

The phenomenon of eutrophication is local, but widespread, and likely to worsen in its manifestations as a result of climate change (Moss et al., 2011; Meerhoff et al., 2022). In particular, the process of internal loading may be enhanced by lake warming (Jeppesen et al., 2009) due to an increased metabolism of bacteria and speed of biochemical reactions. Warming also increases stratification and thermal stability promoting anoxia (Maberly et al. 2020; Woolway et al. 2020). Increases in precipitation, and high intensity rainfall events, are also expected to significantly increase runoff of P from agricultural catchments to surface freshwaters (Ockenden et al., 2017), further promoting eutrophication and its manifestations.

The different states of shallow lakes can feedback differently on climate by either reducing or increasing GHG emissions (Hilt et al. 2017). Clear and turbid lakes differ in their CO₂ emissions due to the magnitude of CO₂ uptake by primary producer photosynthesis. Efflux of CO₂ appears to decrease when submerged macrophytes establish after the reduction of nutrient loading (Jeppesen et al., 2016). Submerged-macrophyte dominated shallow lakes tend to emit lower CH₄ by ebullition and diffusion than phytoplankton turbid lakes (Colina et al., 2022; Davidson et al. 2018). The turbid state in particular feeds back on climate since warming and eutrophication induced water anoxia could offset increased CO₂-fixation by blooms or by macrophytes as lower oxygen levels stimulate methane (CH₄) emission, with CH₄ emissions from eutrophic systems expected to increase with 6-20% with each degree of warming (Aben et al. 2017).

In addition, the eutrophication and warming-associated shift from submerged macrophyte dominance to phytoplankton or floating plant dominance may also strongly increase greenhouse gas emissions, particularly CH₄ (Aben et al. 2022). Cyanobacterial blooms, a typical...
manifestation of eutrophication and high internal P-loading, can both promote CO$_2$ sequestration and produce CH$_4$. CH$_4$ can be produced even under oxic conditions as a by-product of photosynthesis (Bižić et al., 2020). Blooms often create anoxic layers at the surface of aquatic systems or along the water column after their collapse of blooms, favouring the release of CH$_4$ via methanogenesis (Li et al., 2021; Yan et al., 2017). Cyanobacterial blooms are thus considered a key mechanism by which eutrophication has a positive feedback on climate change (Bižić 2021; Yan et al., 2017). Although increased inputs on N from atmospheric deposition or catchment runoff are main causes of elevated N$_2$O release from lakes (Yang et al. 2015), warming also impacts aquatic N$_2$O emissions. N$_2$O emissions are estimated to increase with 8 – 14% for each degree of warming (Velthuis and Veraart 2022), highlighting another strong ecosystem-climate feedback.

Despite the fact that nutrient loading is still the major driver of eutrophication manifestations such as blooms (e.g. Bonilla et al., 2023), climate change is expected to promote eutrophication (Moss et al., 2011; Meerhoff et al., 2022). Indeed, interaction between temperature and trophy has been observed to produce synergistic emission responses in experimental lakes (Davidson et al., 2018) and warming alters resident microbial communities to favour methanogenesis over methanotrophy (Zhu et al., 2020). It is thus likely that warming decreases the nutrient thresholds for a tipping point leading to a shift to an alternative state in shallow lakes and ponds. The predominantly amplifying influence of climate change on eutrophication-driven tipping points in lakes provides a mechanism for coherent tipping beyond the local scale, with more widespread eutrophication-induced tipping points expected with further warming. However, whether any of eutrophication’s climate feedback effect can be buffered by projected eutrophication-driven increases in lake carbon burial (Anderson et al. 2020) remains uncertain, and there is a dearth of studies that generate bi-directional carbon flux data to assess the balance between emission and burial in lakes. Moreover, robust projections are lacking for climate impacts on eutrophication, with no emergent regional to global warming threshold identifiable beyond which a nonlinear increase in these localised tipping points occur (Grasset et al. 2020), which is amplified by the fact that tipping points become harder to predict in a warmer climate (Kosten et al. 2009).
2. Increased loadings of DOM in the boreal biome

Over thousands to millions of years, the mutual feedback between terrestrial vegetation and aquatic productivity has been essential for the evolution of the atmosphere and the global climate (Beerling 2007). Vegetation serves not only as a major C pool and eventually a source of total organic carbon (TOC) in boreal areas, but it also promotes root exudates of CO$_2$ and organic C. This enhances weathering rates thereby increasing the flux of nutrients (P, N, Si, Fe, Ca and carbonate (CO$_3$)) (Humborg et al. 2004; Hessen et al. 2009) to surface waters. The availability of nutrients subsequently enhances aquatic productivity, and thereby C-sequestration. In addition, the carbonate species are important for buffering capacity towards acidification in freshwater and marine systems. On different timescales there is thus a range of feedback mechanisms between terrestrial and aquatic ecosystems that demands a better understanding. Tracking past history (Holocene) tree-line, forest cover and lake sediments, revealed a strong and consistent link between climate, forest cover and lake TOC (Rosén 2005). Thus, at least on the centennial scale, there is a strong temporal TOC-link between terrestrial and aquatic systems. Allochthonous C derived either directly as leachate from litterfall and roots or indirectly via partial decomposition of organic matter in the soils, constitutes the by far dominant pool of dissolved organic matter (DOM) in boreal freshwaters. Forest cover and fraction of bogs and wetland areas in the catchment are key determinants for the concentration and color of this terrestrially derived DOM (Dillon and Molot 1997; Kortelainen et al. 2006; Larsen et al. 2011a), of which TOC is the main constituent.

Since terrestrially derived C is a main determinant of freshwater C, any changes in terrestrial primary production and export of organic C will invariably also increase aquatic outputs of CO$_2$. Increased terrestrial productivity has been linked to a “CO$_2$-fertilization” (Huang et al. 2007) yet these CO$_2$ effects will be constrained by N-availability. Elevated N-deposition due to human emissions has driven a ~12% increase in the forest C sink in tandem with the CO$_2$-fertilization effect, while at the same time also increased the deficiency of phosphorus (and other key elements allocated to tree biomass).

Increased export of terrestrially derived dissolved organic matter (DOM) to lakes and rivers in boreal regions (“browning”) is a widespread phenomenon partly linked to reduced acidification, but also driven by land-use changes (notably afforestation) and climate change (CO$_2$-fertilization of forests, warming and hydrology) (de Wit et al., 2016; Creed et al. 2018;
Monteith et al., 2023). An empirically based space-for-time model of changes in NDVI under a 2°C climate scenario predicts a continued profound browning of boreal lakes (Larsen et al. 2011b). Forest dynamics are slow, however, hence space-for-time scenarios projecting increased flux of TOC from catchments owing to increased forest cover could require centuries to play out. Thus, catchment properties governing production of TOC such as forest size and fraction of bog and wetland areas could very well be temporally decoupled from the export, especially considering the large stock of organic matter typically present in boreal catchments.

Time series analysis (30 years) of data from 70 Norwegian catchments and lakes provided however evidence also for a tight temporal coupling between the decadal increase in land “greening” (with NDVI as a proxy) and lake browning (with TOC as a proxy) (Finstad et al. 2016), and the browning on northern lakes can to a large extent be attributed a recent afforestation (Kritzberg 2017; Skerlep et al. 2020.). The prominent “greening” by increased vegetation cover trend in many boreal and alpine regions (Guay et al. 2014) and increase in forest volume (cf. Opdahl et al. 2023) will thus have bearings on lakes and rivers in these regions. There are a number of confounding explanatory drivers for this greening: warming, elevated CO₂, accumulated nitrogen deposition and changes in grazing activities as well as forestry practices. An extended growing season has also been recorded (Barichivich et al. 2013), and elevated levels of CO₂ per se may contribute to this (Piao et al. 2006). In sum, these changes in the environmental drivers and pressures yield an increase in terrestrial net primary production, notably at high latitudes (Forkel et al. 2016). Since a significant fraction of the terrestrial NPP will be exported to surface waters as DOM, it means that terrestrial greening could lead to freshwater browning.

The role of forest cover is further accentuated by a need for a carbon-negative future (i.e. net drawdown of CO₂ from the atmosphere) where widespread afforestation is the only currently feasible means of reducing atmospheric concentrations of CO₂ beyond the continued action of natural carbon sinks (MacDougall et al., 2020). However, such afforestation also comes with climate costs, both in terms of decreased albedo (Betts and Ball 1997; Bathiany et al. 2010; Lawrence et al. 2022) and as argued above, the potential for increased production and degassing of GHGs from surface waters. Enhanced primary production in forested catchments stimulated by reactive nitrogen deposition has, by increasing the pool of C available for fluvial export, been linked to increased carbon burial in northern lakes over the past two centuries (Heathcote et al.
Again, this highlights the need for improved understanding of the balance between carbon emissions and burial in lakes in response to browning (Williamson et al., 2015) and other identified stressors in order to better constrain climate feedbacks. Browning will also promote darkening of coastal waters with as yet unknown climate feedbacks (Opdal et al. 2023). The question that remains unsettled is whether these terrestrial and aquatic responses are directly coupled in time, or if there is a delayed aquatic response in the order of decades or even millennia. Another issue is how the CO$_2$ in itself could boost these processes, and how this skewed C-supply to autotrophs could affect land-aquatic interactions.

Wide-scale shifts in boreal lakes caused by increased loadings of DOM can promote a prolonged and more intensified stratification period (implications summarized above, described for DOM by Spears et al., 2017), amplified by warming. Increased terrestrial DOM loadings intensify net heterotrophy in the systems (i.e. through increased light attenuation and increased access to organic C for heterotrophic bacteria) (Hessen et al. 1990; Karlsson et al. 2007; Thrane et al. 2014; Horppila et al. 2023). While at present the thresholds around these effects have not been well constrained, the impacts may be significant at the global scale for GHG emissions (Tranvik et al. 2009) and regionally for coastal NPP (Opdal et al. 2019). Given the strong empirical links between drivers and consequences, it means that impacts and feedback can be predicted qualitatively, while not yet quantitatively.

Feedbacks and tipping points

The links and feedbacks between climate to land to lakes and back to climate in terms of increased GHG-emissions is conceptually well understood, and also the main drivers for the specific GHGs (CO$_2$, CH$_4$ and N$_2$O) in boreal areas is understood (Yang et al. 2015; Wik et al. 2016; Valiente 2022). However, the question as of whether these feedbacks can result in tipping points by becoming self-sustaining beyond a threshold is not yet settled. Most boreal lakes are net heterotrophic and thus conduits of CO$_2$, often also CH$_4$, due to high concentrations of DOM and common deep-water of sediment anoxia. A shift from net autotrophy to net heterotrophy would classify as a binary shift, yet with a strong, positive climate feedback. If it eventually leads to oxygen depletion and cascading feedbacks then it would qualify as a tipping point, yet with a time delay between the two events, and where the latter is the critical tipping event. There is also a commonly reported unimodal response in lakes to increased loadings of DOM, typically
around 5 mg DOC l\(^{-1}\) (Karlsson et al. 2007; Thrane et al. 2014), where increases in DOM below the threshold may promote NPP and thus CO\(_2\) drawdown due to N and P associated with DOM, while reduced NPP and increased degassing of CO\(_2\) (and CH\(_4\)) will take place above. We thus propose two types of large-scale potential tipping points, one related to anoxia, the other to DOM-concentrations, yet both are related to increasing load of terrestrially derived DOM across the boreal region.

3. Disappearance/appearance of waterbodies

A global reduction in lake water storage (Yao et al., 2023), and the climate-driven creation or disappearance of water bodies is a crucial issue. Loss of water-bodies due to overuse, warming or draught pose a major threat to vulnerable, freshwater resources, also by deteriorating water quality or salinization (cf. below). The most dramatic warming has already taken place in the high Arctic with temperature increases up to 3 °C over the past few decades (Wang et al. 2022), and onset of permafrost thaw (Langer et al. 2016). Both current and future permafrost thaw and glacier melting can both create new waterbodies and drain old, providing a strong link to the fate of the cryosphere (Smith et al. 2005; Olefeldt et al. 2021). Such small, but numerous waterbodies residing on permafrost over large geographical scales in Eurasia and North-America are currently among the most vulnerable water-bodies globally (Smol and Douglas 2007; Heino et al. 2020). They host species-poor but specific communities of invertebrates (Rautio et al. 2011; Walseng et al. 2021) of vital importance for birdlife and other biota. Warming may also affect these waterbodies indirectly via glacier melt, increased inputs of organic C, fertilisation by increasing populations of geese (caused by climate change), and consequently changes in microbial communities and increased GHG emissions (Eiler et al. 2023). Thus, by their share number these systems may also serve as increasingly important conduits of greenhouse gases and historical soil carbon stocks to the atmosphere (Laurion et al. 2010; Negandhi et al. 2016), and play an important role in mediating nutrient delivery to the polar oceans (Emmerton et al., 2008), potentially affecting global NPP (Terhaar et al., 2021). While the main problem is loss of water bodies resting on (thawing) permafrost (Smol and Douglas 2007) there are also cases where collapsing palsas and thermokarst areas create new waterbodies, and these waterbodies may themselves represent a positive feedback by accelerating the thaw (Langer et al. 2016; Turetsky et al., 2020).
Some essential feedbacks to climate change are involved in the change of Arctic waterbodies; e.g. reduced ice and snow cover in the Arctic will promote further permafrost thaw. More organic carbon entering water bodies from their terrestrial surroundings, combined with warming and eventually bird induced eutrophication will promote GHG emissions. It is important to make clear that some of the impacts are contrasting, i.e. the loss of waterbodies may at first increase GHG emissions (Keller et al. 2020; Paranaiba et al. 2021) but will eventually reduce GHG emissions. Permafrost thaw and drainage of water-logged areas will increase CO₂-emissions, but could reduce CH₄-emissions. Sudden release of methane-hydrates upon permafrost thaw is a possibility, yet hard to predict and quantify, and not specifically linked to aquatic habitats.

Few changes are as irreversible as complete habitat loss, and the climate-driven loss of numerous water-bodies residing on permafrost over large geographical scales in Eurasia and North-America (due to permafrost thaw) with climate feedbacks in terms of changed GHG-emissions is possible. In fact, as argued by Smol and Douglas (2007); “The final ecological threshold for these aquatic ecosystems has now been crossed: complete desiccation”. If strictly adhering to the tipping point criteria as an event occurring when self-sustaining change in a system is triggered beyond a forcing threshold, typically starting with positive feedback loops and a runaway phase before finally the tipping-point brings the system into a different alternative state, loss of waterbodies is not strictly a tipping point, but a binary shift. Abrupt permafrost thaw, which can drive abrupt self-sustained formation or draining of thermokarst lakes, is categorised as a “regional impact” climate tipping element by Armstrong McKay et al. (2022).

We extend this categorisation to include the lakes associated with these abrupt thaw processes, seeing them as a coupled permafrost-lake systems with tipping dynamics involving both components (Turetsky et al., 2020). Despite the scale considered here, the extent of open water globally is relatively easy to quantify using remote sensing, and it is possible to make predictions based on time-series and empirical relationships between temperature increase, permafrost thaw and loss of water-bodies. Quantifying potential climate feedbacks related to processing of organic C to CO₂ and CH₄ should be possible to predict within orders of magnitude, with initial analysis suggesting abrupt thaw involving thermokarst lake formation and draining could double the warming impact of gradual permafrost thaw (Turetsky et al., 2020).
4. Switch from N to P-limitation

Imbalance in biogeochemical cycles has become a major concern both on the local and global scale. Anthropogenic emissions of CO₂ now appear as the major environmental challenge for ecosystems and human well-being in the foreseeable future. In relative terms, however, the anthropogenic effects on the global N-cycle are even more pronounced. Transformation of atmospheric N₂ to more reactive reduced or oxidized forms of inorganic N by fertilizer industry and combustion processes has dramatically changed, and recent analyses of the global N-cycle (Bodirsky et al. 2014; Zhang et al. 2020) suggest that various human activities currently convert similar N₂ to total natural ecosystem fixation, and that both the use of N and P are far beyond “safe boundaries” (Rockström et al. 2023).

Increased N-deposition may affect surface waters in fundamentally different ways. It will increase the emissions of N₂O (Yang et al. 2015), and increased deposition of inorganic N promotes soil and water acidification through increased NO₃ in surface waters (Stoddard 1994). It will however also affect elemental ratios in lakes and rivers (Hessen et al. 2009). The relative proportions of these elements will determine the nature of elemental limitation for both autotrophs and a range of heterotrophs, and could thus profoundly affect community composition and ecosystem processes. One effect of such skewed inputs of N over P would be an intensified P-limitation in surface waters or even large-scale shifts from N to P-limitation (Elser et al. 2009). Conversely, increased N-loss by denitrification, eventually associated with increased internal P-loading may shift systems from P to N-limitation (Weyhenmeyer et al. 2007). Societal implications include an increased prevalence of toxin producing cyanobacteria, purported to be promoted in extent by warming (Paerl et al., 2008) and favouring non-N-fixing toxin producing species where reduced-N concentrations are high relative to oxidized-N (Hoffman et al., 2022).

Additionally, a threshold on toxic effects on sensitive freshwater species has been proposed (i.e. 2 mg L⁻¹; Camargo et al., 2006; Moss et al., 2013), above which a marked decline in biodiversity is expected.

Feedbacks and tipping points:

Changes in N- versus P-limitation of NPP are associated with changes in community structure, both for the phytoplankton and macrophyte communities. While the shift from one limiting
nutrient to another representing no doubt represent a binary shift and abrupt transition, it is not
driven by self-propelling events or positive or negative feedbacks, since a shift from N to P-
limitation typically is caused by N-deposition or agricultural use of fertilizers. While increased
N-loading per se could promote climate feedbacks in terms of N₂O, the switch from N to P-
limitation or vice versa is neither driven by climate or have strong feedbacks on climate. There is
also no inherent hysteresis, and when drivers change the system may immediately return to the
other limiting nutrient. For these reasons we do not classify this category as a tipping point
according to the definition above.

5. Salinization

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

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

Salinization tends to decrease CH$_4$ emissions (Herbert et al. 2015, Chamberlain et al. 2020, Gremmen et al. 2022). The decrease in CH$_4$ emission can be either caused by a decrease in CH$_4$ production - e.g. because methanogens are outcompeted by sulfate reducers or are negatively impacted by sulfide toxicity - or because an increase in methane oxidation (reviewed by Herbert et al. 2015). The salinity induced decrease in aquatic CH$_4$ emissions may imply a negative feedback with climate change, but only when this is not off-set by a decrease in carbon burial. Insight in this balance is currently limited (Chamberlain et al. 2020), and while no doubt salinization are widespread on regional scales and may reach threshold values for species and processes, we do not categorize it is a tipping point under the cited criteria.

6. Spread of invasive species

Freshwaters are especially vulnerable to species loss and population declines as well as species invasions due to their constrained spatial extent. Substantial ecosystem changes by reinforcing interactions between invasive species and alternative states (i.e. macrophyte versus phytoplankton dominance, as described above) may occur (Reynolds and Aldridge 2021). The spread of several invasive species can in dramatic ways change community composition and ecological functions, and per se be regarded as sudden transition with major site-specific or regional impacts. Moreover, species invasions can very well be facilitated by climate change (Rahel and Olden, 2008). While species invasions for good reasons are of major ecological and
societal concern, and can induce ecological tipping points in certain lakes, they are generally not self-propelling involving internal feedbacks. No doubt it may be appropriate to say that invaded system as subject of hysteresis, since also local extinction of species is far from trivial.

Feedbacks and tipping points

We do here not pursue the discussion feedbacks and potential tipping points further for this candidate category since we have constrained our definition of tipping points to situations with internal feedback and regional occurrence. It is however likely that species invasions interact with other drivers lowering the potential thresholds (of nutrients, temperature, browning, etc.) for a shift to occur, and vice versa, by impacting on previously occurring stabilizing mechanisms (Willcock et al. 2023). This is an area that deserves further research.

Discussion

Freshwaters are one of the most vulnerable ecosystems and resources globally, and will increasingly be so with continued global warming. They also link catchment properties and terrestrial changes to marine systems, and notably lakes serve as good sentinels of global change (Adrian et al. 2009). Population declines and species loss of freshwater species are happening at an alarming pace, and is another reason why knowledge on the ecological status of lakes is important. Drinkable freshwater is a scarce resource qualitatively, but also quantitatively (Yao et al. 2023). Predicting (and preventing) sudden shifts in water quality and quantity is therefore a high priority also from an anthropocentric perspective, and insights into feedbacks, thresholds and tipping points are highly relevant to lakes. Lake are also major players in the global climate, and besides being highly vulnerable to climate change, they can provide strong feedback to the climate by ventilating a substantial share of terrestrially fixed C back to the atmosphere as CO₂ and CH₄ (Cole et al. 2007; Tranvik et al. 2009; Raymond et al. 2013). Lakes are also subject to changes, sometimes sudden, due to climate change and other natural or anthropogenic drivers. In fact, some of the first and most striking examples on tipping points and regime shift come from lake studies (Scheffer et al. 1993; Jeppesen et al. 1998).

We argue that there are two key drivers that may shift lakes towards major ecological changes, as well as increased climate feedback by GHG emissions, namely eutrophication and browning (increased loadings of terrestrially derived DOM). Both these drivers are promoted by
warming, which per se may be seen as a separate driver. Both processes are also characterised to some degree by self-sustaining feedback loops, feedback to climate in terms of GHG-emissions, and are also strongly integrated with land surface impacts in the catchment (Fig. 3). Warming, browning, and eutrophication lead to increases in stratification, heterotrophy, and algal mass, which collectively drive benthic oxygen depletion and in turn increased GHG emissions (helping to drive further warming and DOM loading from land) and internal P loading (driving further eutrophication) (Meerhoff et al. 2022). Several of these processes can feature tipping points (eutrophication and potentially DOM loading), which warming will likely make easier to reach. Few processes have been more thoroughly described in terms of drivers, impacts and remedies than freshwater eutrophication. The drivers are well known (nutrient loadings, basically from agricultural activities, but locally also sewage), despite long-term controversies regarding the relative importance of nitrogen or phosphorus in promoting eutrophication (e.g., Smith & Schindler 2009, Paerl et al. 2016). There are also long traditions for predictive hydraulic models that link the load of phosphorus to algal blooming and benthic O₂-depletion (e.g., Vollenweider type models, Imboden 1974). Moreover, given the scarcity, increasing demands and increasing prices of P worldwide, there are indeed strong arguments to close the loop for P and reduce excess P (Spears et al. 2022). Due to the strong impact of O₂-depletion on sediment release of P and thus internal fertilization (Soendergaard et al. 2002), that will play in concert with food-web driven feedbacks (cf. Fig. 3), tipping points in this context can be identified, while the climate component is difficult to separate.

Browning shares many of these attributes in terms of increased net heterotrophy. Shift from net autotrophy with a net uptake of CO₂ to net heterotrophy with a net release of CO₂ (plus CH₄) also represents a binary situation, yet since most boreal lakes already are net heterotrophic owing to microbial conversion of organic C (Hessen et al. 1990; Cole et al. 1994; Larsen et al.
Fig. 3. The interactive role of eutrophication, DOM-export (browning) and warming on lakes. Separately or combined they promote benthic O₂-depletions which cause an internal feedback by P-loading from sediments and a climate feedback via release of greenhouse gases. The potential shift between states (blue to red circle) is indicated.

As a separate type of binary tipping point, widespread and with feedback of GHG-release, we propose the loss of water bodies, notably Arctic ponds. This is driven by permafrost...
thaw in the case of thermokarst-linked lake formation or disappearance (categorised as a regional tipping element in previous assessments (Armstrong McKay et al. 2022), but together the coupled permafrost-lake system can act as a localised tipping system with the lake providing key feedbacks to help drive self-sustaining thaw. This makes the tipping points easy to monitor (by remote sensing), and predictable in the sense that it will be closely linked to permafrost thaw. There are however feedbacks to the climate, with potentially high emissions during the drying process (Marcé et al. 2019; Turetsky et al. 2020) although the final disappearance of water bodies could in fact reduce GHG-emissions and thus serve as a negative feedback. A different situation would be the less widespread case of new waterbodies formed by collapsing palsas, in cases also retreating glaciers, but the combined net effect of permafrost thaw and increased release of CO$_2$ by oxidised organic C and the effect of disappearing waterbodies is not settled but should be a research question of high priority.

Gradients or tipping points – does it matter?

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

Shifts between ecological states do not necessarily involve alternative stable states with hysteresis. In fact, both the concepts of abruptness and irreversibility depends on time perspective. Over a lakes life-time perspective, shifts back and forth between states occurring over years or even decades are “sudden” in a relative sense. For example, Rühland et al. (2008) report apparent coherence in diatom community shifts post 1850 on hemispheric scales over 100 years or so. Similarly, a coherent, global increase in hypoxia in lakes have reported over a 100 years period (from about 1850) by Jenny et al. (2016). If the observational time step is increased to centuries, then it is likely that more large-scale examples will come through in paleo-studies. In fact, there are several examples on
coherence in lake responses to climate variability or climate change, some of which also can take place over short time spans (Stone et al. 2016; Isles et al. 2023). Finally, it is also worth pointing to the fact that multiple drivers may jointly drive lakes towards shifts or tipping points, as shown in Huang et al. (2022) and Willcock et al. (2023).

Taken together, there are at least two major reasons why an improved understanding of sudden changes in lake ecosystems are imperative; they are highly vulnerable to climate change and other anthropogenic stressors globally, and they serve as major feedbacks to the climate system by GHG emissions. Being well-mixed and semi-closed entities, still reflecting changes in catchment properties, they also serve as sentinels of global change (Adrian et al. 2019). For fresh waters in general, lakes are crucial in the hydrological cycling, and link the terrestrial and marine ecosystems. The major tipping point dynamics converge in oxygen depletion, primarily in deeper strata and the sediment surface, which promotes feedbacks and hysteresis in terms of internal P release as well as increased GHG-emissions.

High nutrient load, increased inputs of dissolved organic C and warming all drive oxygen depletion, and while many problems related to global warming boils down to the obvious recommendation of reduced use of fossil fuel and other GHG-emitting activities, reducing nutrient loading is comparatively simpler both for N and P, both elements that long time ago have crossed the “safe boundary” thresholds (Rockström et al. 2009; 2023). The incentives should be even larger for closing the P-loop, given the scarcity of this non-substitutable element and its role in eutrophication (Brownlie et al. 2022).

Regime shifts and tipping points are concepts closely linked to resilience (Andersen et al. 2008; Spears et al. 2017). Lakes represent excellent model case studies in this respect and have been used widely to demonstrate theories of ecological stability and resilience that are needed to underpin preventative management approaches and to guide science-based environmental policy.

The full importance of the vulnerability of lakes to climate and other anthropogenic impacts, as well as their feedback to climate is not yet fully acknowledged, so there is a need both for science and communication in this regard. However, we argue that the search for empirical evidence to underpin theory should not prevent societies and managers taking more action to protect fresh waters in the meantime.

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