

Review: Climate-induced hydrological changes and the ecology of tropical freshwater biota

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Abstract. Sunardi, Wiegleb G. 2016. *Climate-induced hydrological changes and the ecology of tropical freshwater biota. Biodiversitas 17: 322-331.* Climate change is believed to pose adverse effects to biodiversity of aquatic systems, with no exception of those in tropical areas. However, next few decades species extinction is suggested as dark future as we lack researches uncovering how climate change threatens the aquatic biota. Unluckily, the tropical freshwater systems are expected to suffer more severe impacts of climate change, from heavy floods or extended drought than do the boreal areas. A comprehensive understanding of biota' performance in face of climatic pressures, will guide the further necessary researches. This paper presents a review on the available researches addressing ecological effects of the most influential climatic parameter in tropic area, the hydrological regime, on freshwater biota. The research reveal that the extreme water fluctuations induced by climate change have negatively affected the performance of freshwater biota. In the next few decades, climate change seems to remain as one of the main threats for freshwater ecosystems, and is responsible for the lost of its biodiversity.

Keywords: Biota performance, climate change, drought, flood, tropical freshwater

INTRODUCTION

Global change has been shown and predicted to have major effects on biodiversity at global, regional, and local scales, although global change constitutes a number of different forms of anthropogenic impacts (Sala et al. 2000), including land use alterations, nitrogen deposition, and invasions of exotic species. In other words, changes in climate and climate variability would, somehow, significantly affect natural ecosystems, and may pose additional threats to the already-stressed ecosystems. Furthermore, the effect of future climate change on biodiversity has been predicted to be unprecedented as well, with 15-37% of terrestrial species possibly becoming extinct due to climate change alone in the next 50 years (Thomas et al. 2004), and a similarly dark future has been suggested for freshwater species in the next few decades (Xenopoulos et al. 2005).

Freshwater ecosystems are vulnerable to global change. Important global climate variables that are expected to change in the next decades with respect to freshwater habitat are air temperature and precipitation (Mitchell et al. 1990). Changes in these variables will affect water temperature, water quantity and water quality variables of freshwater environments which are the three primary linkages between climate and freshwater organisms (Regier and Meisner 1990).

Climate change pushes species out of their ecological synchrony and environmental landscape. This influences

not only species distributions or community structure, but also the services they provide to ecosystems. Understanding how species' performances change along with the environmental gradients is worthwhile. This is particularly important in aquatic systems, where shifts in habitat quality associated with environmental perturbations threaten the integrity of aquatic biota (Strayer et al. 2004).

The magnitude of impacts and responses of aquatic ecosystems, however, differ between boreal and tropical area with regard to the global change. This is due to climatic variation between the two areas. In the tropics, the annual variation in air temperature is smaller, but there is a large and predictable annual variation in precipitation (Lowe-McConnell 1987). The seasonal precipitation cycle produces wide ranges in river flow rates and water levels, which directly alters the amount of freshwater habitat available for biota and indirectly alters many critical characteristics of that habitat (eg, O₂ levels, turbidity, food availability, etc.).

The increase in global temperature, nevertheless, will lead to a more vigorous hydrological cycle, with changes in precipitation and evapotranspiration rates regionally variable. Warming accelerates land-surface drying as heat goes into evaporation of moisture, and this increases the potential incidence and severity of droughts, which has been observed in many places worldwide (Dai et al. 2004). In tropical systems, evaporation and evapotranspiration often already exceed precipitation in the dry season (Irion and Junk 1997). In weather systems, convergence of

increased water vapor leads to more intense precipitation and the risk of heavy rain and snow events, but may also lead to reductions in duration and/or frequency of rain events, given that total amounts do not change much (see Trenberth 2005). In such case, the tropical areas are expected to suffer more severe impacts of climate change, from heavy floods or extended drought than do the boreal areas.

Despite the increase of research on the topic, we lack a comprehensive understanding of the consequences of extreme precipitation fluctuation (as distinct from seasonal climate) on the ecology of tropical freshwater biota. The aim of the present article is to complement the existing information by reviewing current knowledge of climate change effects, paying particular attention on hydrological regime, on species' performances in tropical freshwater.

KEY CLIMATE-RELATED PARAMETERS

Aquatic ecosystems are vulnerable to changes in quantity and quality of their water supply, and it is expected that climate change will have a pronounced effect on global freshwater through elevated temperature and alterations in hydrological regimes with great global variability. As a matter of fact, aquatic organisms have to deal with a wide variety of environmental factors simultaneously; however, temperature, water quantity and water quality are regarded as the most fundamental climate-related factors (Figure 1).

In tropical regions, fluctuations in rainfall often represent the strongest seasonal variation, and change the environment to an extent comparable to temperature in temperate areas (Jacobsen and Encalada 1998). Variation in rainfall that affects stream discharge is among the most important sources of natural disturbances (Taylor et al. 1996). Flow regimes range from spates or peak flows during the rainy season through to zero flow in the dry season. The shape and size of river channels, the distribution of riffle and pool habitats, and the stability of the substrate are all largely determined by the interaction between the flow regime and local geology and landform (Newbury and Gaboury 1993). In turn, this complex interaction between flows and physical habitat becomes the major determinant of the distribution, abundance, and diversity of stream and river organisms (Nilsson and Svedmark 2002).

It is reported that effects of climatic variability on hydrology can be particularly devastating, causing changes in water chemistry, stream size, water temperature, streambed structure, streambed substrate and stream flow (see Medeiros and Maltchik 2001; Starks et al. 2014). Hence, such environmental variation can dramatically alter the living conditions and aquatic habitats within the water, affecting much of the aquatic fauna inhabiting streams (Moyle and Vondracek 1985; Taylor and Warren 2001). Biota, in particular those living in vegetated riparian areas, respond differentially to water-level dynamics, either directly or indirectly. Direct effects on the biological communities include physical disturbance by wave activity.

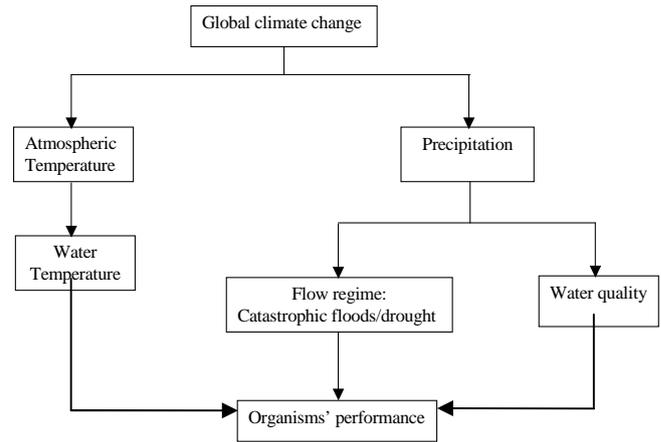


Figure 1. Key climate-related parameters affecting global freshwater organisms

Indirect effects include the reworking of substrates (which can enhance or restrict colonization by vegetation, and which in turn depends on silt accumulation to establish roots), and alteration of habitats suitable for aquatic flora and fauna. Any significant changes in the water level of a lake will affect not only the physical processes, but also the biological productivity. Therefore, water level fluctuation might have an overriding effect on Lake Ecosystem health and integrity (Furey et al. 2006; Leira and Cantonati 2008).

In addition, ambient temperature is undoubtedly among the most important for ectotherm ontogeny because of its pervasive effects on biological rate processes, altering nearly all physiological functions (Deutsch et al. 2008; Kingsolver 2009). It has been known for more than a century that biochemical reaction rates, metabolic rates, and nearly all other rates of biological activity increase exponentially with temperature. In tropical area, however, seasonal temperature variability is small (Meisner and Shuter 1992), and it seems does not pose profound effects on organism' performance as does in boreal areas. Therefore, this paper will not deal with the effects of inclined temperature on freshwater organisms.

IMPACTS OF FLOOD ON BIOTA

Water flow in aquatic ecosystem, particularly lotic, is often subject to a high short-term variability. This short-term variability may be caused by high precipitation events in the catchment. Hydroperiod and flood frequency (Medley and Havel 2007), as well as high water flow (Godlewska et al. 2003) have been recognized as significant factors structuring communities.

This suggests that hydrology has a significant effect on both species richness and community structure of planktonic communities in rivers, streams, floodplain ponds and lakes. Flood events often act as disturbances that interrupt the succession in planktonic community (Mulyaert et al. 2001); even a short flood event may have effects on the planktonic ecosystem that last for weeks.

Nevertheless, the research showed that the responses of the planktonic communities differ from site to site showing its dependence to other biotic and abiotic factors. Some groups of plankton may respond positively or negatively, while some others do not show clear response; e.g. Chlorophyll a concentration and abundance of bacteria, oligotrich ciliates and crustacean zooplankton did not respond clearly to the flood event (Mulyaert and Vyverman 2006).

It is evident that flood waters transport large amounts of suspended solids and nutrients into lakes. An increase in the concentration of suspended matter in a lake leads to greater light attenuation and thus to a decrease in primary production (Lloyd et al. 1987). High concentrations of abiotic turbidity can light-limit phytoplankton photosynthesis and therefore restrict biomass development (Holst and Dokulil 1987; Dokulil 1994). In most cases, an increase in nutrient input causes a consequent rapid increase in algal biomass. This is especially true of oligotrophic lakes (Thomas 1973). However, in several lakes, an impoverishment of the algal standing stock and a decrease in the phosphorus concentration are observed following events involving the discharge of suspended sediment into the lake, despite the fact that the turbid inflow transports a large load of particulate phosphorus into the lake (Sampl 1986). This is due to the phytoplankton is sedimented out after flood event, as the phytoplankters and suspended matter do coagulate efficiently (Elber and Schanz 1990), result in a decrease in primary production and phytoplankton biomass. Additionally, the negative responses can occur under condition of prolonged floods because the flood pulse can have dilution effects on nutrient, so that a significantly lower phytoplankton biomass is established (Keckeis et al. 2003; Mihaljevi et al. 2009).

Grobbelaar (1992) and Dokulil (1994) suggest that the ratio of mixing to euphotic depth is one of the most important factors affecting overall productivity mainly in turbid waters. In these water bodies the aphotic portion is large compared to the euphotic zone and determines the relative time spent in the dark by the algae. The ratio of euphotic to mixing depth is considered to be the most important factor affecting overall productivity, while nutrients are of secondary importance only. Nutrients influence productivity only when a more favorable underwater light regime prevails, such as prior to the flooding. Energy available for phytoplankton growth is strongly regulated by underwater light availability which depends on the critical mixing depth, fluctuating light intensities and algal circulation patterns.

Godlewska et al. (2003) reported that shifts in phytoplankton distribution (from hypolimnion into the whole water column) and species composition were observed. In one case, this is because of high water flow eliminated large species of cladocerans and copepods, and favored development of rotifers; while in another case, plankton animals may concentrate at different depths in the water column before and after the flood because they are transported to different locations by water currents. Dirnberger and Threlkeld (1986) found that during the

period of flooding most zooplankton populations declined and the distributions of remaining individuals deepened; greatest population losses occurred between 0 and 7m depths where most zooplankton had been concentrated prior to the flood. The changes in distribution may be results from search for the right compromise between two conflicting demands: to maximize feeding and to avoid predators (Gliwicz 1986).

On the other hand, floods can have positive impacts on planktonic communities. Dispersal among patches is important to the long-term viability of species in metapopulations, and flood connections can enhance the viability of certain species (Jenkins 1995). Immigrants with differential competitive ability can be introduced into communities which releases local communities from competitive exclusion, shifts local dynamics, and enhance long-term persistence (Leibold et al. 2004). Flooding can introduce new species (Havel et al. 2000), but intense flooding can wash out entire populations (Baranyi et al. 2002). Turbulence, which increases in high flood, can also have direct effects on reduced grazing rate (Miquelis et al. 1998) and food selectivity of zooplankton (Vanderploeg 1994).

In river, increase in discharge leads to increased drifts. Downstream invertebrate drift is a normal feature in lotic systems and facilitates the recolonization of denuded areas of a stream (Brittain and Eikeland 1988). However, disturbances notably flooding have an important role in regulating the distribution, abundance and coexistence of macroinvertebrate (Resh et al. 1988). Significant reductions in macroinvertebrate density have been recorded after scouring floods (Robinson et al. 2004). While moderate disturbance may encourage diversity in many systems (see Smith and Brown 2006). In regulated river reaches below dams, it was reported that sudden increases in flow can cause catastrophic downstream drift (Layzer et al. 1989).

The most frequently reported effect of sedimentation associated with floods is an increase in drift density (e.g., Doeg and Milledge 1991; Suren and Jowett 2001), which may account for the loss of individuals and species in response to a loss of suitable habitat and changes to the food web (Rabeni et al. 2005). In addition, high water flow can also be acute for invertebrate; Doeg and Koehn (1994) identified a reduction in total number of benthic macroinvertebrate taxa and abundance, after a flushing event that increased suspended solid concentration. Pruitt et al. (2001) reported that total suspended solids concentrations greater than 284 mg L⁻¹ resulted in biological impairment of invertebrate communities, while a concentration of 58 mg L⁻¹ or less during storm flow provided an adequate margin of safety and were protective of aquatic invertebrates. Variability in tolerance to suspended solid could be explained by sediment particle characteristics, water temperature, species differences and other stressor that might have synergistic effects (Bash et al. 2001). In addition, the degree of turbidity associated with flood events has been known to affect the response of benthic invertebrate to flood. Present study report that the magnitude of response of macroinvertebrate community to the flood is most severe in the non-turbid, upper main river

and tributaries where benthos community is dominated by the most sensitive Ephemeroptera, Plecoptera and Trichoptera species (Miserendino 2009).

Sediment transport and deposition are processes that are a natural part of the stream environment and play a major role in structuring stream habitats. However, streams are vulnerable to increased sedimentation brought about by altered land uses in the surrounding catchments, with detrimental effects on benthic stream communities. The exacerbation of erosion and sedimentation may be particularly striking in the tropics (Newcombe and MacDonald 1991; Wood and Armitage 1997) where extreme climatic conditions can prevail and aquatic systems are increasingly under threat. However, ways of how sediment affects aquatic ecosystems are various depending on the shape, size and density of the particles; their potential for microbial colonization; the velocity, temperature, flow and turbulence of the water (Hellowell 1986); and the presence of associated factors, such as nutrients (Lemly 1982). Increased levels of sedimentation can bury macroinvertebrates and their habitats (Wood et al. 2001; Wood et al. 2005) leading to shifts in the structure of the habitat and its associated fauna (e.g., Ryder 1989).

Sedimentation has been shown to induce behavioral response of macroinvertebrate, where macroinvertebrates actively avoid substratum coated with excessive fine sediment (McClelland and Brusven 1980; Connolly and Pearson 2007). It is predicted that the upland fauna would be more sensitive to sedimentation because it would naturally experience lower exposure to sedimentation than the lowland fauna; such different response has been demonstrated in both, the mesocosm and in-situ experiments (Connolly and Pearson 2007). Fine sediment deposition can cause shifts in the community structure through the loss of sensitive species, particularly those requiring coarse substrata for attachment or feeding, and through increases in the abundance of burrowing animals, such as some Chironomidae and Oligochaeta (Hellowell 1986). Sedimentation can also affect the filter feeders, scrapers and collector through ingestion of inorganic when feeding, with a negative effect on nutrition and growth (Ryder 1989). Fine silt deposit trapped by periphyton can reduce photosynthesis (Yamada and Nakamura 2002), and thus algal availability to grazers (Donohue and Irvine 2004). There may also be indirect effects of sedimentation transmitted through top-down effects of predators, such as fish and crayfish (Schofield et al. 2004). However, several studies have shown that changes in abundance rather than diversity are commonly associated with sedimentation (e.g., Lenat et al. 1981, Wagener and LaPerriere 1985).

In large lentic ecosystem, water level fluctuation plays a role more importantly than does flow regime. Such effects on ecosystems are very complex, and the biological effects in lakes are greatest in shallow water and littoral areas, where even small changes in water levels can result in the conversion of large areas of a standing-water environment in air exposed habitats (Leira and Cantonati 2008). The potential effects of lake-level changes have been judged by impacts at the physical level, i.e. transparency, sedimentation patterns, erosion; at the species

level, i.e. target species, and by indicators at the ecosystem level, i.e. carrying capacity and biodiversity (Leira and Cantonati 2008). The fluctuation of water level can alter the lake morphometry and transform the characteristics of the sedimentation zone (erosion, transportation, accumulation; Håkanson 1977), thereby water-level drawdown enhances sediment erosion and has the potential to fundamentally change littoral sediment and biogeochemical characteristics (Furey et al. 2004). The water level fluctuation cause changes in the littoral area available for benthic macroinvertebrate. The lost of littoral vegetation due to inundation or the establishment of emergent species from seed during low water is always accompanied by changes in invertebrates and amphibians (Eulis et al. 2004).

The functioning of shallow lakes and floodplains is supposedly very sensitive to water level changes. Moreover, littoral plant communities in shallow lakes located in semi-arid to arid regions appear to be particularly susceptible to water-level fluctuations (Beklioglu et al. 2006). High water level can facilitate the expansion of submerged vegetation that from which some benthic invertebrates take benefits.

The effects of water level fluctuation on benthic macroinvertebrate are very subtle, both directly and indirectly (Leira and Cantonati 2008). Direct effects on invertebrates include changes in the structure and dynamic of taxa that cannot withstand dry periods which subsequently lead to a limiting of their distribution by low water levels (Rossa and Bonecker 2003; Bowers and de Szalay 2004). Meanwhile, indirect effects are generally through alteration of habitats (e.g. substrate composition, periphyton growth, resuspension versus sedimentation). Particularly important are those habitats with cobbles and macrophytes that provide an extensive suitable habitat for periphytic algae which are their major food source, egg-laying and tube building, and also provide a refuge from predation (Scheifhacker et al. 2007). However, different zooplankton groups seem to show different sensibility to water level and are distinctly affected by floods (Ortega-Mayagoitia et al. 2000).

Fishes are particularly susceptible to changes in environmental conditions. Flow plays a profound role in the lives of fish with critical life events linked to flow regime (see Bunn et al. 2002; Janá et al. 2010). Numerous studies have shown that changes in stream flow associated with extreme variations in precipitation can alter fish communities and habitats. Many fish species display a preference for particular types of habitat such as pools, riffles, or backwater areas. While habitat structure is generally considered to be a good predictor of fish assemblage structure, so habitat instability associated with variations in stream flow will disturb the fish community residing within it (Gelwick et al. 2001). Therefore, sudden or long term variations in discharge, such as from extended droughts or large storms, can be particularly devastating, causing changes in water chemistry, stream size, water temperature, streambed structure, streambed substrate and stream flow (Medeiros and Maltchik 2001). Extreme discharge associated with storm events can dramatically

alter channel morphology and benthic habitat, which may have significant effects on fish community. Such environmental alteration can dramatically alter the living conditions and aquatic habitats within the water, affecting much of the aquatic fauna inhabiting streams (Moyle and Vondracek 1985; Taylor and Warren 2001).

Crosa et al. (2009) reported that a large volume of sediment associated with reservoir flushing has decreased fish density and biomass; a greater mortality recorded for juveniles will likely result in long-term impairment of the age-structures fish populations. Juveniles' mortality was also reported due to flushing (Garric et al. 1990), while lower effect such as damage of gill epithelium was observed (Petz-Glechner et al. 2003). High level of sediment can cause mortality in fish (Newcombe and MacDonald 1991), in particular sensitive specimen (Lloyd 1987), whilst prolonged lower levels of suspended solids and turbidity is the cause of chronic sublethal effects, such as reduced weight since individuals are not able to feed efficiently (Sigler et al. 1984). Stream fish can become stranded on gravel bars or trapped in off-channel habitats during rapid flow decreases. Susceptibility to stranding is a function of behavioral response to changing flows, and this varies with species, body size, water temperature, time of year and day, substrate characteristics, and the rate of flow reductions (Bradford 1997). However, mature fish may be able to shift into temporarily suitable habitats to compensate for periodic reductions in the quality or availability of habitat (Bunt et al. 1999).

Many variables known to affect fish, including sediment load, pH, dissolved oxygen, and various nutrients, are frequently changed during increased flow associated with storm events (Winemiller et al. 2000, Ostrand and Wilde 2002). Winemiller et al. (2000) showed that diversity and abundance of freshwater fish populations positively correlate with total dissolved nitrogen, nutrient concentration, and food resources in the water. Gelwick et al. (2001) found positive correlations between common measures of assemblage structure (diversity and abundance) and dissolved oxygen and salinity. However, in other case, the degree of change in chemical composition observed was minor (Keaton et al. 2005).

On the other hand, extreme storm events that lead to flooding can introduce new species into assemblages and create new habitats (Winemiller et al. 2000), or increase availability of shelter and allochthonous food sources, and should provide water enrichment with nutrients carried from adjacent areas or present in flooded organic or inorganic material (Agostinho et al. 2004). Nevertheless, floods can dilute the aquatic biota by increasing water depth, reducing the availability of food resources, especially mobile ones. As a result, the hydrological cycle affect interspecific relations, particularly predation and competition. The flooding regime seems to favor piscivores, since floods are associated with the reproductive success of many of their prey species. However, due to their diluting effect, floods also reduce the density of prey species. In addition, increased shelter may also reduce prey availability (Luz-Agostinho et al. 2008; Janáč et al. 2010).

DROUGHT AND FRESHWATER BIOTA

In contrast with the facts on flood effects, there have been relatively few studies of stream faunal dynamics after droughts (see Lake 2000). If floods amplify hydrological connectivity, conversely droughts in streams disrupt hydrological connectivity. With the onset of drought, falling water levels reduce the habitat availability for most aquatic biota, exposing the marginal areas (Stanley et al. 1997), breaking surface water contact between the stream and its riparian zone, and reducing the hydraulic heterogeneity of flow. Changing water levels are another stressor on lake and littoral communities. Water level fluctuations in lakes are dominant forces controlling the functioning of lacustrine ecosystems (Wilcox and Meeker 1992; Poff et al. 1997). It plays an important role in the lake's physical processes (e.g. the geomorphologic processes of erosion and sedimentation) (Leira and Cantonati 2008). With falling water levels, lentic habitats may increase in extent and new types of habitats may be created, that favor some species. As drying proceeds, the threshold of cessation of surface flow is reached.

Droughts can have direct and indirect impacts on stream biota. Direct impacts are those caused by loss of water and flow, and habitat reduction and reconfiguration, whereas indirect impacts are those associated with changes in phenomena such as interspecific interactions, especially predation and competition, and the nature of food resources. The direct and indirect impacts of drought can greatly reduce population densities, species richness and alter life-history schedules, species composition, patterns of abundance, type and strength of biotic interactions (predation and competition), food resources, trophic structure and ecosystem processes. Resh (1992) found that a severe drought has eliminated a population of the caddisfly, *Gumaga nigricula* (McL.).

Following water flow reduction, many aquatic biota cannot move and become trapped and concentrated in lingering pools (Boulton et al. 1992; Matthews 1998; Matthews and Marsh-Matthews 2003). Stream connectivity becomes differentially disrupted by the cessation of upstream-downstream longitudinal links, and the weakening of lateral links between the stream channel and riparian zone (this includes the flood plain) and vertical links between the surface, hyporheic zone and groundwater. In terms of available data on responses to drought in flowing waters, more information is available on invertebrates and fish than on micro-and macroalgae, macrophytes and riparian plants (e.g. Peterson 1996; Matthews 1998; Holmes 1999). During drought, flow may cease and some stretches of river consist of isolated pools, where biota become concentrated in pools and such pools may harbor very high densities of invertebrates (e.g. Boulton and Lake 1992; Miller and Golladay 1996) and of fish (e.g. Matthews 1998; Labbe and Fausch 2000; Matthews and Marsh-Matthews 2003). Different isolated pools may harbor different assemblages of biota and with time, such pools can diverge from each other in their community structure (Power et al. 1985; Meyerhoff and Lind 1987; Stanley et al. 1997). In reservoirs, during

extreme drawdown events, the water quality changes significantly. In particular, drawdown events cause changes to nutrient dynamics and ultimately lead to periods of high algal biomass; in one case leading to the formation of a potentially toxic cyanobacterial bloom (Naselli-Flores 2003).

Physicochemical conditions shift rapidly with flow cessation and undoubtedly have adverse effects on the benthos. When flow decreases, the capacity of the stream to transport organic matter decreases and cause an increase in detritus coverage. Further, flow cessation lead to watercourse fragmentation into a series of isolated pools and caused an abrupt change in physicochemical conditions that imposed a threshold on the ecosystem (Acuna 2005). Organic matter or detritus and sediments are accumulated in pools, and reduce physical reaeration causing a decrease in DO concentrations and an increase in nutrient concentrations, typical responses to pool isolation (Stanley et al. 1997; Caruso 2002).

On the onset of drought, tolerant species can grow rapidly leading to a density peak soon after flow cessation, but this density peak dropped rapidly in response to changes caused by flow cessation (Boulton and Lake 1992). The most probable causes of these adverse effects are deoxygenation (Stanley et al. 1997; Labbe and Fausch 2000) and toxicity of certain leachates from leaf decomposition (Townes 1991; Boulton and Lake 1990, 1992; Chergui et al. 1997). As streams dry and the surface water shrinks to unshaded pools, the build-up of nutrients, high temperatures and solar radiation can precipitate blooms of algae (Freeman et al. 1994; Dahm et al. 2003; Winder et al. 2012). The algae may create large diel changes in oxygen concentration (Matthews 1998). As water flow and volumes decrease, water temperatures may start to rise (Acuna 2005) and can become lethal for aquatic biota such as fish (Matthews 1998). Simultaneously, especially in pools, deoxygenation may occur, threatening biota (Stanley et al. 1997; Labbe and Fausch 2000; Golladay et al. 2002).

Low discharge conditions during drought can limit habitat resources and mobility (Lohr and Fausch 1997) and can have marked effects on community composition, diversity, size structure of populations, spawning, and recruitment of fish (see Poff et al. 2001; Lake 2003; Ledger et al. 2012). Droughts also results in intense aggregations of fish and possible competition for food and/or space, because fish are confined to small areas and usually at considerably higher densities, thus potentially increasing competition. Poff and Ward (1989) considered that such biotic interactions contribute relatively little to community structure in rivers. However, during periods of low flow, and the attendant reduction of habitat area or volume, biotic interactions could become temporarily important (Cowx et al. 1984; Matthews 1988). Fish population structure can also be changed by drought (Resh et al. 2013), reducing spawning and recruitment (Cowx et al. 1984; Davies et al. 1988). Pires et al. (1999) note that some species are well adapted to natural droughts, however major native species are considered to be more sensitive to stream fragmentation and hydrological alteration (Parkin et al. 2014). In addition,

habitat degradation and possibly the introduction of exotic species contribute to marked variability in species composition. Meanwhile, droughts may raise water temperature and reduce dissolved oxygen, imposing adverse physical conditions on fish in the pools.

CONCLUDING REMARKS

Research have regarded that tropical areas are expected to suffer more severe impacts of climate change, from heavy floods or extended drought. Such extreme fluctuation of water flow will, to some extent, threat the performance of tropical freshwater biota. Floods and drought can negatively affect the biota, both directly and indirectly, through the changing environment. Research suggested that the magnitude of hydrological changes may be acute to species, disturb reproduction or succession, change the distribution, and lastly restructure the biotic communities in rivers, streams, floodplain, ponds, and lakes.

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REFERENCES

- Acuna V, Munõz I, Giorgi A, Omella M, Sabateri F, Sabateri S. 2005. Drought and postdrought recovery cycles in an intermittent Mediterranean stream: structural and functional aspects. *J N Am Benthol Soc* 24 (4): 919-933.
- Agostinho AA, Gomes LC, Veri'ssimo SV, Okada EK. 2004. Flood regime, dam regulation and fish in the Upper Paraná River: effects on assemblage attributes, reproduction and recruitment. *Rev Fish Biol Fish* 14 (1): 11-19.
- Alderdice DF. 1976. Some concepts and descriptions of physiological tolerance: rate-temperature curves of poikilotherms as transects of response surfaces. *J Fish Res Bd Can* 33: 299-307.
- Allan JD, Palmer M, Poff NL. 2005. Climate change and freshwater ecosystems. In: Lovejoy TE, Hannah L (eds) *Climate Change and Biodiversity*. Yale University Press, New Haven, CT.
- Baranyi C, Hein T, Holarek C, Keckeis S, Schiemer F. 2002. Zooplankton biomass and community structure in a Danube River floodplain system: effects of hydrology. *Freshw Biol* 47: 473-482.
- Bash J, Berman C, Bolton S. 2001. Effects of turbidity and suspended solids on Salmonids. Report by Centre for Streamside Studies, University of Washington, Washington.
- Beklioglu M, Altınayar G, Tan CO. 2006. Water level control over submerged macrophyte development in five shallow lakes of Mediterranean Turkey. *Archiv für Hydrobiologie* 166: 535-556.
- Bertahasa I, Dimitriou E, Karaouzasa I, Laschoua S, Zacharias I. 2006. Climate change and agricultural pollution effects on the trophic status of a Mediterranean lake. *Acta Hydrochim Hydrobiol* 34: 349-359.
- Boulton AJ, Lake PS. 1990. The ecology of two intermittent streams in Victoria, Australia. I. Multivariate analyses of physicochemical features. *Freshw Biol* 24: 123-141.
- Boulton AJ, Lake PS. 1992. The ecology of two intermittent streams in Victoria, Australia. III. Temporal changes in faunal composition. *Freshw Biol* 27: 123-138.

- Bowers RW, de Szalay FA. 2004. Effects of hydrology on unionids (Unionidae) and zebra mussels (Dreissenidae) in a Lake Erie coastal wetland. *Am Midl Nat* 151: 286-300.
- Boyd C, Tucker C. 1998. Pond aquaculture water quality management. Kluwer Academic Publishers, Norwell, MA.
- Bradford MJ. 1997. An experimental study of stranding of juvenile salmonids on gravel bars and inside channels during rapid flow fluctuations. *Regul Rivers Res Manag* 13: 395-401.
- Brett JR. 1971. Energetic responses of salmon to temperature. A study of some thermal relations in the physiology and freshwater ecology of sockeye salmon (*Oncorhynchus nerka*). *Am Zool* 11: 99-113.
- Brittain JE, Eikeland TJ. 1988. Invertebrate drift. *Hydrobiologia* 166: 77-93.
- Bunt CM, Cooke SJ, Katopodis C, McKinley RS. 1999. Movement and summer habitat of brown trout (*Salmo trutta*) below a pulsed discharge hydroelectric generating station. *Regul Rivers Res Manag* 15: 395-403.
- Bunn SE, Arthington AH. 2002. Basic principles and ecological consequences of altered flow regimes for aquatic biodiversity. *Environ Manag* 30: 492-507.
- Campagna CG, Cech Jr JJ. 1981. Gill ventilation and respiratory efficiency of Sacramento blackfish, *Orthodon microlepidotus*, in hypoxic environments. *J Fish Biol* 19: 581-591.
- Caruso BS. 2002. Temporal and spatial patterns of extreme low flows and effects on stream ecosystems in Otago. *N Z J Hydrol* 257: 115-133.
- Chellappa S, Câmara MR, Chellappa NT. 2004. Ecology of *Cichla monoculus* (Osteichthyes: Cichlidae) from a reservoir in the semi-arid region of Brazil. *Hydrobiologia* 504: 267-273.
- Chergui H, Haddy L, Markaoui M, Pattee E. 1997. Impact of dead leaves leaching products on water oxygen content and on the survival of a gastropod. *Acta Oecologica* 18: 531-542.
- Christensen JH, Christensen OB. 2003. Severe summertime flooding in Europe. *Nature* 421: 805-806.
- Connolly NM, Pearson RG. 2007. The effect of fine sedimentation on tropical stream macroinvertebrate assemblages: a comparison using flow through artificial stream channels and recirculating mesocosms. *Hydrobiologia* 592: 423-438.
- Cowx IG, Young WO, Hellawell JM. 1984. The influence of drought on the fish and invertebrate populations of an upland stream in Wales. *Freshw Biol* 14: 165-177.
- Crosa G, Castelli E, Gentili G, Espa P. 2009. Effects of suspended sediments from reservoir flushing on fish and macroinvertebrates in an alpine stream. *Aquat Sci* 72 (1): 85-95.
- Dahm C, Baker MA, Moore DI, Thibault JR. 2003. Coupled biogeochemical and hydrological responses of streams and rivers to drought. *Freshw Biol* 48: 1219-1231.
- Dai AG, Trenberth KE, Qian T. 2004. A global dataset of Palmer Drought severity index for 1870-2002: relationship with soil moisture and effects of surface warming. *J Hydrometeorol* 5: 1117-1130.
- Davies PE, Sloane RD, Andrew J. 1988. Effects of hydrological change and the cessation of stocking on a stream population of *Salmo trutta* L. *Aust J Mar Freshw Res* 39: 337-354.
- Deutsch C., Tewksbury A., Huey JJ., Sheldon RB., Ghalambor CK., Haak D C., Martin PR. 2008. Impacts of climate warming on terrestrial ectotherms across latitude. *Proc Natl Acad Sci USA* 105: 6668-6672.
- Dirnberger JM, Threlkeld ST. 1986. Advective effects of a reservoir flood on zooplankton abundance and dispersion. *Freshw Biol* 16: 387-396.
- Doeg TJ, Milledge GA. 1991. The effects of experimentally increasing suspended sediment concentrations on macroinvertebrate drift. *Aust J Mar Freshw Res* 42: 519-526.
- Doeg TJ, Koehn JD. 1994. Effects of draining and desilting a small weir on downstream fish and macroinvertebrates. *Regul Rivers Res Manag* 9: 263-277.
- Dokulil MT. 1984. Assessment of components controlling phytoplankton photosynthesis and bacterioplankton production in a shallow, alkaline, turbid lake (Neusiedler See, Austria). *Intl Revue Ges Hydrobiol* 69: 679-727.
- Dokulil MT. 1994. Environmental control of phytoplankton productivity in turbulent turbid Systems. *Hydrobiologia* 289: 65-72.
- Donohue I, Irvine K. 2004. Size-specific effects of increased sediment loads on gastropod communities in Lake Tanganyika, Africa. *Hydrobiologia* 522: 337-342.
- Dudgeon D. 2003. The contribution of scientific information to the conservation and management of freshwater biodiversity in tropical Asia. *Hydrobiologia* 500: 295-314.
- Elber F, Schanz F. 1990. The influence of a flood event on phytoplankton succession. *Aquatic Sciences* 52 (4): 330-344.
- Euliss NH, Labaugh JW, Fredrickson LH, Mushet DM, Laubhan MRK, Swanson GA, Winter TC, Rosenberry DO, Nelson RD. 2004. The wetland continuum: a conceptual framework for interpreting biological studies. *Wetlands* 24: 448-458.
- Ficke AD, Myrick CA, Hansen LJ. 2007. Potential impacts of global climate change on freshwater fisheries. *Rev Fish Biol Fish* 17: 581-613.
- Flecker AS, Feifarek B. 1994. Disturbance and the temporal variability of invertebrate assemblages in two Andean streams. *Freshw Biol* 31: 131-142.
- Franklin CE, Johnston IA, Crockford T, Kamunde C. 1995. Scaling of oxygen consumption of Lake Magadi tilapia, a fish living at 37°C. *J Fish Biol* 46: 829-834.
- Freeman C, Gresswell R, Guasch H, Hudson JA, Lock MA, Reynold B, Sabater S, Sabater F. 1994. The Role of drought in the impact of climatic-change on the Microbiota of peatland streams. *Freshw Biol* 32: 223-230.
- Fry FEJ. 1971. The Effect of Environmental Factors on the Physiology of Fish. In: Hoar WS, Randall DJ (eds) *Fish Physiology: Environmental Relations and Behavior*. Academic Press, New York.
- Furey PC, Nordin RN, Mazumder A. 2004. Water level drawdown affects physical and biogeochemical properties of littoral sediments of a reservoir and a natural lake. *Lake Reserv Manag* 20: 280-295.
- Furey PC., Nordin RN., Mazumder A. 2006. Littoral benthic macroinvertebrates under contrasting drawdown in a reservoir and a natural lake. *J N Am Benthol Soc* 25: 19-31.
- Garric J, Migeon B, Vindimian E. 1990. Lethal effects of draining on brown trout. A predictive model based on field and laboratory studies. *Water Res* 24 (1): 59-65.
- Gelwick FP, Akin S, Arrington DA, Winemiller KO. 2001. Fish assemblage structure in relation to environmental variation in a Texas gulf coastal wetland. *Estuaries* 24: 285-296.
- Gibbons JW. 1976. Thermal alteration and the enhancement of species populations. In: Esch GW, McFarlane RW (eds) *Thermal Ecology II Conference 750425*. ERDA Symposium Series. Technical Information Center, United States Atomic Energy Commission, Oakridge, TN.
- Gliwicz ZM. 1986. Predation and the evolution of vertical migration in zooplankton. *Nature* 320: 746-748.
- Godlewska M, Mazurkiewicz-Boron G, Pocięcha A, Wilk-Wozniak E, Jelonek M. 2003. Effects of flood on the functioning of the Dobczyce reservoir ecosystem. *Hydrobiologia* 504: 305-313.
- Grobbelaar JU. 1992. Nutrients versus physical factors in determining the primary productivity of waters with high inorganic turbidity. *Hydrobiologia* 238: 177.
- Groisman PY, Karl TR, Easterling DR, Knight RW, Jamason PF, Hennessy KJ, Suppiah R, Page CM, Wibig J, Fortuniak K, Razuvaev VN, Douglas A, Forland E, Zhai PM. 1999. Changes in the probability of heavy precipitation: important indicators of climate change. *Climatic Change* 42: 243-283.
- Groisman PY, Knight RW, Karl TR, Easterling DR, Sun B, Lawrimore J. 2004. Contemporary changes of the hydrological cycle over the contiguous United States: trends. *J Hydrometeorol* 5: 64-85.
- Groisman PY, Knight RW, Easterling DR, Karl TR, Hegerl GC, Razuvaev VN. 2005. Trends in intense precipitation in the climate record. *J Clim* 18: 1326-1350.
- Håkanson L. 1977. Influence of wind, fetch, and water depth on distribution of sediments in lake Vanern, Sweden. *Can J Earth Sci* 14: 397-412.
- Harem A, Kucklentz V. 1981. Effects of hydraulic load changes on the eutrophication of an alpine lake. *Verb Internat Verein Limnol* 21: 466-472.
- Havel JE, Eisenbacher EM, Black AA. 2000. Diversity of crustacean zooplankton in riparian wetlands: colonization and egg banks. *Aquat Ecol* 34: 63-76.
- Hellawell J. 1986. *Biological Indicators of Freshwater Pollution and Environmental Management*. Elsevier, London.
- Holmes NTH. 1999. Recovery of headwater stream flora following the 1989-1992 groundwater drought. *Hydrol Process* 13: 341-354.
- Holst I, Dokulil M. 1987. Die steuemden Faktoren der planktischen Primtrproduktion im Stauraum Altenworth an der Donau in Osterreich. 26. Arbeitstagung der IAD, Passau/Deutschland, Wiss. Kurzreferate: 133-137.

- Huntington TG. 2006. Evidence for intensification of the global water cycle: review and synthesis. *J Hydrol* 319: 83-95.
- Hynes HBN. 1970. *The Ecology of Running Waters*. Liverpool University Press, Liverpool, UK.
- Intergovernmental Panel on Climate Change (IPCC). 2001. *Climate change 2001: the scientific basis*. Cambridge University Press, Oxford, UK.
- Irion G, Junk WJ. 1997. The large Central Amazonian River floodplains near Manaus. In: Junk WJ (ed) *The Central Amazon Foodplain: Ecology of a Pulsing System*. Springer, Berlin.
- Jacobsen D, Encalada A. 1998. The macroinvertebrate fauna of Ecuadorian Highland streams in wet and dry seasons. *Archive Für Hydrobiologie* 142 (1): 53-70.
- Janá M, Ondra ková M, Jurajda P, Valová Z, Reichard M. 2010. Flood duration determines the reproduction success of fish in artificial oxbows in a floodplain of a Potamal river. *Ecol Freshw Fish* 19: 644-655.
- Jenkins DG. 1995. Dispersal-limited zooplankton distribution and community composition in new ponds. *Hydrobiologia* 313/ 314: 15-20.
- Keaton M, Haney D, Andersen CB. 2005. Impact of drought upon fish assemblage structure in two South Carolina Piedmont streams. *Hydrobiologia* 545: 209-223.
- Keckeis S, Baranyi C, Hein T, Holarek C, Riedler P, Schiemer F. 2003. The significance of zooplankton grazing in a floodplain system of the River Danube. *J Plankton Res* 25: 243-253.
- Kingsolver JG. 2009. The well-temperated biologist. *Am Nat* 174:755-768.
- Kitchell JF, Stewart DJ, Weininger D. 1977. Applications of a bioenergetics model to yellow perch (*Perca flavescens*) and walleye (*Stizostedion vireum vireum*). *J Fish Res Bd Can* 34: 1922-1935.
- Labbe TR, Fausch KD. 2000. Dynamics of intermittent stream habitat regulate persistence of a threatened fish at multiple scales. *Ecol Appl* 10: 1774-1791.
- Lake PS. 2000. Disturbance patchiness and diversity in streams. *J N Am Benthol Soc* 19: 573-592.
- Lake PS. 2003. Ecological effects of perturbation by drought in flowing waters. *Freshw Biol* 48: 1161-1172.
- Layzer JB, Nehus TJ, Pennington W, Gore JA, Nestler JM. 1989. Seasonal variation in the composition of drift below a peaking hydroelectric project. *Regul Rivers Res Manag* 3: 305-317
- Ledger ME, Brown LE, Edwards FK, Milner AM, Woodward G. 2012. Drought alters the structure and functioning of complex food webs. *Nature Climate Change. Global Change Biol* 17: 2288-2297.
- Leibold MA, Holyoak M, Mouquet N, Amarasekare P, Chase JM, Hoopes MF, Holt RD, Shurin JB, Law R, Tilman D, Loreau M, Gonzalez A. 2004. The metacommunity concept: a framework for multi-scale community ecology. *Ecol Lett* 7: 601-613.
- Laira M, Cantonati M. 2008. Effects of water-level fluctuations on lakes: an annotated bibliography. *Hydrobiologia* 613: 171-184.
- Lemly AD. 1982. Modification of benthic insect communities in polluted streams: combined effects of sedimentation and nutrient enrichment. *Hydrobiologia* 87: 229-245.
- Lenat DR, Penrose DL, Eagleson KW. 1981. Variable effects of sediment addition on stream benthos. *Hydrobiologia* 79: 187-194.
- Lloyd DS, Koenings JP, LaPerriere JD. 1987. Effects of turbidity in fresh waters of Alaska. *N Am J Fish Manag* 7: 18-33.
- Lloyd D. 1987. Turbidity as a water quality standard for salmonid habitats in Alaska. *North Am J Fish Manag* 7: 34-45.
- Lohr SC, Fausch KD. 1997. Multiscale analysis of natural variability in stream fish assemblages of a western Great Plains watershed. *Copeia* 1997: 706-724.
- Lowe-McConnell RA. 1987. *Ecological Studies in Tropical Fish Communities*. Cambridge University Press, Cambridge.
- Luz-Agostinho KDG, Agostinho AA, Gomes LC, Jílio-Jr HF. 2008. Influence of flood pulses on diet composition and trophic relationships among piscivorous fish in the upper Parana River floodplain. *Hydrobiologia* 607: 187-198.
- Magnuson JJ. 2002. Future of adapting to climate change and variability. In: McGinn NA (ed) *Fisheries in a Changing Climate*. American Fisheries Society, Bethesda, MD.
- Matthews WJ. 1998. *Patterns in Freshwater Fish Ecology*. Chapman & Hall, New York.
- Matthews WJ, Marsh-Matthews E. 2003. Effects of drought on fish across axes of space, time and ecological complexity. *Freshw Biol* 48: 1232-1253.
- McClelland WT, Brusven MA. 1980. Effects of sedimentation on the behaviour and distribution of riffle insects in a laboratory stream. *Aquat Insects* 2: 161-169.
- Medeiros ESF, Maltchik L. 2001. Fish assemblage stability in an intermittently flowing stream from the Brazilian semiarid region. *Austr Ecol* 26: 156-164.
- Medley KA, Havel JE. 2007. Hydrology and local environment factors influencing zooplankton communities in floodplain ponds. *Wetlands* 27 (4): 864-872.
- Meisner JD. 1992. Assessing potential effects of global climate change on tropical freshwater fishes. *Geojournal* 28: 21-27.
- Meyerhoff RD, Lind OT. 1987. Factors affecting the benthic community structure of a discontinuous stream in Guadalupe Mountains National Park, Texas. *Internationale Revue der gesamten Hydrobiologie* 72: 283-296.
- Mihaljević M, Stević F, Horvatić J, Kutuzovic BH. 2009. Dual impact of the flood pulses on the phytoplankton assemblages in a Danubian floodplain lake (Kopa ki Rit Nature Park, Croatia). *Hydrobiologia* 618: 77-88.
- Miller AM, Golladay SW. 1996. Effects of spates and drying on macroinvertebrate assemblages of an intermittent and perennial prairie stream. *J N Am Benthol Soc* 15: 670-689.
- Miquelis A, Rougier C, Pourriot R. 1998. Impact of turbulence and turbidity on the grazing rate of the rotifer *Brachionus calyciflorus* (Pallas). *Hydrobiologia* 386: 203-211.
- Miserendino ML. 2009. Effects of flow regulation, basin characteristics and land-use on macroinvertebrate communities in a large arid Patagonian river. *Biodivers Conserv* 18: 1921-1943.
- Mitchell JFB, Manabe S, Meleshko V, Tokioka T. 1990. Equilibrium climate change-and its implications for the future. In: Houghton JT, Jenkins GJ, Ephraums JJ (eds) *Climate Change. The IPCC Scientific Assessment*. Cambridge University Press, Cambridge.
- Mol J, Vandenberghe J, Kasse C. 2000. River response to variations of periglacial climate in mid-latitude Europe. *Geomorphology* 33: 131-148.
- Moyle PB, Vondracek B. 1985. Persistence and structure of the fish assemblages in a small California stream. *Ecology* 66: 1-13
- Moyle PB, Cech JJ. 2004. *Fishes: An Introduction to Ichthyology*. 5th ed. Prentice Hall, Englewood Cliffs, NJ
- Muyllaert K, Van Wichelen J, Sabbe K, Vyverman W. 2001. Effects of freshets on phytoplankton dynamics in a freshwater tidal estuary (Schelde, Belgium). *Arch Hydrobiol* 150 (2): 269-288.
- Naselli-Flores L. 2003. Man-made lakes in Mediterranean semiarid climate: The strange case of Dr Deep Lake and Mr Shallow Lake. *Hydrobiologia* 506/509: 13-21.
- Newbury R, Gaboury M. 1993. Exploration and rehabilitation of hydraulic habitats in streams using principles of fluvial behaviour. *Freshw Biol* 29: 195-210.
- Newcombe CP, MacDonald DD. 1991. Effect of suspended sediments on aquatic ecosystems. *J N Am Benthol Soc* 11: 72-82.
- Nicholls KH. 1998. El Nino, ice cover, and Great Lakes phosphorus: implications for climate warming. *Limnol Oceanogr* 43: 715-719.
- Ortega-Mayagoitia E, Armengol X, Rojo C. 2000. Structure and dynamics of zooplankton in a semi-arid wetland, the National Park Las Tablas de Daimiel (Spain). *Wetlands* 20: 629-638.
- Ostrand KG, Wilde GR. 2002. Seasonal and spatial variation in a prairie stream-fish assemblage. *Ecol Freshw Fish* 11 (3): 137-149.
- Palmer TN, Räisänen J. 2002. Quantifying the risk of extreme seasonal precipitation events in a changing climate. *Nature* 415: 512-514.
- Parmesan C. 2006. Ecological and evolutionary responses to recent climate change. *Ann Rev Ecol Evol Syst* 37: 637-669.
- Perkin JS, Gido KB, Costigan KH, Daniel MD, Johnson ER. 2014. Fragmentation and drying ratchet down Great Plains streamfish diversity. *Aquat Conserv Mar Freshw Ecosyst*. DOI: 10.1002/aqc.2501
- Peterson CG. 1996. Response of benthic algal communities to natural physical disturbance. Pp. 375-402 In: Stevenson JR, Bothwell ML, Lowe RL (eds) *Algal Ecology: Freshwater Benthic Ecosystems*. Academic Press, San Diego.
- Petz-Glechner R, Petz W, Kainz E, Lapuch O. 2003. Die Auswirkungen von Stauraumpülungen auf Fische. *Natur in Tirol-Ökologie und Wasserkraftnutzung*. Amt der Tiroler Landesregierung-Abteilung Umweltschutz 12: 74-93.
- Pires AM, Cowx IG, Coelho MM. 1999. Seasonal changes in fish community structure of intermittent streams in the middle reaches of the Guadiana basin, Portugal. *J Fish Biol* 54: 235-249.

- Poff NL, Ward JV. 1989. Implications of stream flow variability and predictability for lotic community structure: a regional analysis of stream flow patterns. *Can J Fish Aquat Sci* 46: 1805-1818
- Poff NL, Angermeier PL, Cooper SD. 2001. Fish diversity in streams and rivers. In: Sala OE, Chapin F, Huber-Sannwald E (eds) *Global Biodiversity in a Changing Environment: Scenarios for the 21st Century*. Springer, New York.
- Poff NL, Allan JD, Bain MB. 1997. The natural flow regime: a paradigm for river conservation and restoration. *Bioscience* 47: 769-784.
- Power ME, Matthews WJ, Stewart AJ. 1985. Grazing minnows, piscivorous bass and stream algae. *Ecology* 66: 1448-1456.
- Pruitt AB, Melgaard DL, Flexner MC, Able AS. 2001. Chattooga River Watershed Ecological/Sedimentation Project. In: Proceedings of the Federal Interagency Sedimentation Conference, held 26-30 March in Reno, NV.
- Rabeni CF, Doisy KE, Zweig LD. 2005. Stream invertebrate community functional responses to deposited sediment. *Aquat Sci* 6 (7): 395-402
- Regier HA, Meisner JD. 1990. Anticipated effects of climate change on freshwater fishes and their habitat. *Fisheries* 15 (6): 10-15.
- Resh VH. 1992. Year-to-year changes in the age structure of a caddisfly population following loss and recovery of a Springbrook habitat. *Ecography* 15: 314-317.
- Resh VH, Brown AV, Covich AP, Gurtz ME, Li HW, Minshall GW, Reice SR, Sheldon AL, Wallace JB, Wissmar RC. 1988. The role of disturbance in stream ecology. *J N Am Benthol Soc* 7: 433-455.
- Resh VH, Bêche LA, Lawrence JE, Mazor RD, McElravy EP, O'Dowd AP, Rudnick D, Carlson SP. 2013. Long-term population and community patterns of benthic macroinvertebrates and fishes in Northern California Mediterranean-climate streams. *Hydrobiologia* 719: 93-118.
- Reyjol Y, Lim P, Dauba F, Baran P, Belaud A. 2001. Role of temperature and flow regulation on the Salmoniform-Cypriniform transition. *Arch Hydrobiol* 152 (4): 567-582.
- Robinson CT, Aebischer S, Uehlinger U. 2004. Immediate and habitat-specific responses of macroinvertebrates to sequential, experimental floods. *J N Am Benthol Soc* 23: 853-867.
- Root TL, Price JT, Hall KR, Schneider SH, Rosenzweig C, Pounds JA. 2003. Fingerprints of global warming on wild animals and plants. *Nature* 421: 57-60.
- Rossa DC, Bonecker CC. 2003. Abundance of planktonic and non-planktonic rotifers in floodplain lakes of the Upper Paraná River floodplain. *Amazoniana* 17 (3-4): 567-581.
- Ryder GI. 1989. Experimental Studies on the Effects of Fine Sediment on Lotic Invertebrates. [Ph.D. Dissertation]. Department of Zoology, University of Otago, Dunedin, New Zealand.
- Sala OE, Chapin FS, Armesto JJ. 2000. Biodiversity-global biodiversity scenarios for the year 2100. *Science* 287: 1770-1774.
- Sampl H. 1986. Einfluß von Nährstoffabschwemmung und Bodenerosion auf die Gewässereutrophierung. Bundesministerium für Land- und Forstwirtschaft, Wien.
- Scheiffhacken N, Fiek C, Rothhaupt KO. 2007. Complex spatial and temporal patterns of littoral benthic communities interacting with water level fluctuations and wind exposure in the littoral zone of a large lake. *Fund Appl Limnol* 169 (2): 115-129.
- Schofield KA, Pringle CM, Meyer JL. 2004. Effects of increased bedload on algal-and detrital-based stream food webs: Experimental manipulation of sediment and macroconsumers. *Limnol Oceanogr* 49 (4): 900-909.
- Sigler JW, Bjorn TC, Everest FH. 1984. Effects of chronic turbidity on density and growth of steelheads and co salmon. *T Am Fish Soc* 113: 142-150.
- Smith, MAK. 1991. Models of seasonal growth of the equatorial carp *Labeo dussumieri* in response to the river flood cycle. *Environ Biol Fishes* 31: 157-170.
- Smith F, Brown AV. 2006. Effects of flow on meiofauna colonization in artificial streams and reference sites within the Illinois River, Arkansas. *Hydrobiologia* 571: 169-180.
- Somero GN, Hofmann GE. 1997. Temperature thresholds for protein adaptation: when does temperature start to 'hurt'? In: Wood CM, McDonald DG (eds) *Global Warming: Implications for Freshwater and Marine Fish*. Cambridge University Press, Cambridge, UK.
- Sorensen DL, McCarthy MM, Middlebrook EJ, Porcella DB. 1977. Suspended and Dissolved Solids Effects on Freshwater Biota: A Review. United States Environmental Protection Agency. 600/3-77-042.
- Spooner DE, Vaughn CC. 2008. A trait-based approach to species' roles in stream ecosystems: climate change, community structure, and material cycling. *Oecologia* 158: 307-317.
- Stanley EH, Fisher SG, Grimm NB. 1997. Ecosystem expansion and contraction in streams. *BioScience* 47: 427-435.
- Starks E, Cooper R, Leavitt PR, Wissel B. 2014. Effects of drought and pluvial periods on fish and zooplankton communities in prairie lakes: systematic and asystematic responses. *Global Change Biology* 20: 1032-1042.
- Strayer DL, Downing JA, Haag WR, King TL, Layzer JB, Newton TJ, Nichols S. 2004. Changing perspectives on pearly mussels, North America's most imperiled animals. *BioScience* 54: 429-439
- Suren AM, Jowett IA. 2001. Effects of deposited sediment on invertebrate drift: an experimental study. *New Zeal J Mar Fresh Res* 35: 725-737
- Taylor EW, Egginton G, Taylor SE, Butler PJ. 1997. Factors which may limit swimming performance at different temperatures. In: Wood CM, McDonald DG (eds) *Global Warming: Implications for Freshwater and Marine Fish*. Cambridge University Press, Cambridge, UK.
- Taylor CM, Warren ML. 2001. Dynamics in species composition of stream fish assemblages: environmental variability and nested subsets. *Ecology* 82: 2320-2330
- Taylor CM, Winston MR, Matthews WJ. 1996. Temporal variation in tributary and mainstem fish assemblages in a Great Plains stream system. *Copeia* 1996: 280-289
- Thomas EA. 1973. Phosphorus and eutrophication. In: Griffith EJ, Beeton A, Spencer JM, Mitchell DT (eds) *Environmental Phosphorus Handbook*. John Wiley & Sons, New York.
- Thomas CD, Cameron A, Green RE. 2004. Extinction risk from climate change. *Nature* 427: 145-148
- Towns DR. 1991. Ecology of leptocecid caddisfly larvae in an intermittent South Australian stream receiving Eucalyptus litter. *Freshw Biol* 25: 117-129.
- Trenberth KE. 1998. Atmospheric moisture residence times and cycling: Implication for rainfall rates with climate change. *Climate Change* 36: 667-694.
- Trenberth KE. 2005. Uncertainty in hurricanes and global warming. *Science* 308: 1753-1754.
- Val AL, Almeida-Val VMF. 1995. Fishes of the Amazon and their environment: physiological and biochemical aspect. Springer, Berlin.
- Van Der Kraak G, Pankhurst NW. 1997. Temperature effects on the reproductive performance of fish. In: Wood CM, McDonald DG (eds) *Global Warming: Implications for Freshwater and Marine Fish*. University Press, Cambridge, UK.
- Vanderploeg HA. 1994. Zooplankton particle selection and feeding mechanisms. In: Wotton RS (ed) *The Biology of Particles in Aquatic Systems*. 2nd ed. Lewis Publishers, Boca Raton.
- Wagener SM, LaPerriere JD. 1985. Effects of placer mining on the invertebrate communities of interior Alaska streams. *Freshw Invert Biol* 4: 208-214.
- Wang J, Tsai C. 2000. Effects of temperature on the deformity and sex differentiation of tilapia, *Oreochromis mossambicus*. *J Exp Zool* 286: 534-537.
- Webb MAH, Van Eenennaam JP, Feist GW, Linares-Casenave J, Fitzpatrick MS, Schreck CB, Doroshov SI. 2001. Effects of thermal regime on ovarian maturation and plasma sex steroids in farmed white sturgeon, *Acipenser transmontanus*. *Aquaculture* 201: 137-151.
- Wilcox DA, Meeker JE. 1992. Implications for faunal habitat related to altered macrophyte structure in regulated lakes in northern Minnesota. *Wetlands* 12: 192-203.
- Winder M, Berger SA, Lewandowska A, Aberle N, Lengfellner K, Sommer U, Diehl S. 2012. Spring phenological responses of marine and freshwater plankton to changing temperature and light conditions. *Mar Biol*. DOI 10.1007/s00227-012-1664-z
- Winemiller KO, Tarim S, Shormann D, Cotner J. 2000. Fish assemblage structure in relation to environmental variation among Brazos River oxbow lakes. *T Am Fish Soc* 129: 451-468.
- Wohlschlag DE, Cameron JN, Cech JJ. 1968. Seasonal changes in the respiratory metabolism of the pinfish (*Lagodon rhomboids*). *Contr Mar Sci* 13: 89-104.
- Wood PJ Armitage PD. 1997. Biological effects of fine sediment in the lotic environment. *Environ Manag* 21: 203-217
- Wood PJ, Vann AR, Wanless PJ. 2001. The response of *Melampophylax mucoreus* (Hagen) (Trichoptera: Limnephilidae) to rapid sedimentation. *Hydrobiologia* 455: 183-188

- Wood PJ, Toone J, Greenwood MT, Armitage PD. 2005. The response of four lotic macroinvertebrate taxa to burial by sediments. *Archiv für Hydrobiologie* 163: 145-162
- Wotton RS. 1994. Particulate and dissolved organic matter as food. In: Wotton RS (ed) *The Biology of Particles in Aquatic Systems*. 2nd ed. Lewis Publishers, Boca Raton.
- Wotton RS. 1995. Temperature and lake-outlet communities. *J Thermal Biol* 20: 121-125
- Xenopoulos MA, Lodge DM, Alcamo J, Märker M, Schulze K, van Vuuren DP. 2005) Scenarios of freshwater fish extinctions from climate change and water withdrawal. *Glob Change Biol* 11: 1557-1564
- Yamada H., Nakamura F. 2002. Effect of fine sediment deposition and channel works on periphyton biomass in the Makomanai River, Northern Japan. *River Res Appl* 18: 481-493.