

Extreme Climate Anomalies Enhancing Cyanobacterial Blooms in Eutrophic Lake Taihu, China

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Abstract

Climate warming in combination with nutrient enrichment can greatly promote phytoplankton proliferation and blooms in eutrophic waters. Lake Taihu, China, is a large, shallow and eutrophic system. Since 2007, this lake has experienced extensive nutrient input reductions aimed at controling cyanobacterial blooms. However, intense cyanobacterial blooms have persisted through 2017 with a record-setting bloom occurring in May, 2017. Causal analysis suggested that this bloom was sygenerically driven by high external loading from flooding in 2016 in the Taihu catchment and a notable warmer winter during 2016/2017. High precipitation during 2016 was associated with a strong 2015/2016 El Niño in combination with the joint effects of Atlantic Multi-decadal Oscillation (AMO) and Pacific Decadal Oscillation (PDO), whilst persistent warmth during 2016/2017 was strongly related to warm phases of AMO and PDO. The 2017 blooms elevated water column pH and led to dissolved oxygen depletion near the sediment, both of which mobilized phosphorus from the sediment to overlying water, further promoting cyanobacterial blooms. Our finding indicates that regional climate anomalies exacerbated eutrophication via a positive feedback mechanism, by intensifying internal nutrient cycling and aggravating cyanobacterial blooms. In light of global expansion of eutrophication and blooms, especially in large, shallow and eutrophic lakes, these regional effects of climate anomalies are nested within larger scale global warming predicted to continue in the foreseeable future.

Key words: Lake Taihu; eutrophication; cyanobacterial bloom; winter warmth; heavy rainfall; positive feedback; shallow lakes

Eutrophication and harmful algal blooms (HABs) are a world-wide proliferating environmental issue[*S Wang et al.*, 2018]. Anthropogenic nurient enrichment and climate change are two major drivers, determining the magnitude, extent and persistence of cyanobacterial blooms[*Hans W. Paerl et al.*, 2020]. While nutrient overenrichment is the chief cause of eutrophication and nuisance cyanobacterial blooms[*Brookes and Carey*, 2011], climate warming frequently enhances the dominance of cyanobacteria and further accelerates eutrophication[*Rigosi et al.*, 2014]. Global warming and altered precipitation patterns are projected to aggravate eutrophication at continental scale in the future[*Sinha et al.*, 2017], and is a key management challenge facing eutrophic waters.

Lake eutrophication and harmful algal blooms are severe in China, particularly in the lakes from the middle and lower reaches of the Yangtze River (MLRYR) [*Guan et al.*, 2020; *B. Q. Qin et al.*, 2013]. The number of lakes from MLRYR accounts for ca. 60% of total number of freshwater lakes in China[*S M Wang and dou*, 1998], and all lakes are characterized by shallow depth. Lake eutrophication in this area are mainly induced synergistically by anthropogenic nutrient emission and climate warming[*Guan et al.*, 2020; *B.Q. Qin et al.*, 2010]. In addition, this area experience a monsoonal climate with humid and hot summer, and dry and cold winter, while the climate anomalies of this region are often intimately linked to the variability of global scale atmospheric-oceanic interactions[*Li et al.*, 2019]. Long-term observations suggested the warming trend since the beginning of last century in China is estimated higher than the global mean of temperature increase[*Z Yan et al.*, 2020], implying the influence of climate change tends to be more significant[*Yang et al.*, 2016].

Lake Taihu is a large, shallow and eutrophic lake situated in the delta of the Yangtze River which is one of the most industrialized and urbanized areas in China. This lake has experienced an extensive restoration since the drinking water crisis in 2007. However, a record-setting cyanobacterial bloom occurred in May 2017 and intense bloom persisted throughout 2017 in Lake Taihu [B Qin et al., 2019], which can not be explained simply by a climate warming or nutrient enrichment. As the external loading in 2017 was not the highest, we hypothesize that the severe cyanobacterial bloom in 2017 is resulted from a combination effects of intense internal nutrient loading with climate anomalies caused by global scale, inter-annual, decadal or multi-decadal external climate forcings such as El Niño events, Atlantic multidecadal oscillation (AMO), Pacific decadal oscillation (PDO), North Atlantic Oscillation (NAO) and Arctic Oscillation (AO). There are few studies have addressed such combination effects on the phytoplankton proliferation in inland waters. We tested this hypothesis by documenting: 1) what were the main drivers of severe blooms which occurred in 2017 in Lake Taihu, and 2) whether bloom persistence in 2017 were associated with a combination of internal nutrient loading and and climate anomalies during 2016/2017.

2 Materials and Methods

2.1 Lake Taihu

Lake Taihu, is China's third largest freshwater lake (2338 km²), located in a large, heavily-urbanized (> 40 million inhabitants) catchment (~36500 km²) in the Yangtze River Delta region (Fig. 1). Prevailing southeasterly monsoonal winds during summertime concentrate nutrients discharged from northwestern watershed and cyanobacterial blooms in the northwestern region of lake. Since the 1990's, this shallow (max depth < 3 m) eutrophic lake has experienced accelerated eutrophication accompanied by toxic cyanobacterial blooms (*Microcystis* spp.), especially in the highly eutrophic northern bays or northwestern coastal zone areas that are subjected to prevailing southeasterly, onshore winds in the summer, and receive the most external pollutants from the watershed [*B Qin et al.*, 2007]. In May 2007, a massive bloom overwhelmed the lake's drinking water plants, leaving millions of local residents of Wuxi City without potable water for nearly a week[*Guo*, 2007]. Since the drinking water crisis, Taihu has been the site of intensive restoration efforts. A series of countermeasures, including effluent diversion, flushing and dredging, were implemented, but little improvement of water quality has achieved [*B Qin et al.*, 2019].

2.2 Cyanobacterial bloom monitoring

Because of the high spatial heterogeneity of cyanobacterial blooms in Lake Taihu, we used remote sensing imagery to map the spatial distribution of the bloom. Due to the inherent problems in atmospheric correction and bio-optical inversion algorithms for inland waters, variable aerosols, cloud cover, and wind speed (turbulent entrainment to make algal bloom migration downward), and other nonliving constituents (e.g., tripton, colored dissolved organic matter (CDOM), suspended sediments and shallow bottom) can significantly affect the remote sensing signal, especially for the case of Lake Taihu . These deviations have been minimized by selecting the bands between red and near-IR and algorithm, and the statistical reliability of long-term monitoring data was evaluated by Hu et al., 2010 [*Hu et al.*, 2010]. Such deviation was evaluated against the concurrent in-situ measurements, and the results showed that relative error (RE) ranged from 0.4% to 64.5% with a mean absolute percent error (MAPE) of 27.1%, and root-mean-square error (RMSE) values between the measured and predicted values were 15.01 μ g/L [*Shi et al.*, 2017].

Bloom area was retrieved from satellite imagery by utilizing Moderate Resolution Imaging Spectroradiometer (MODIS) reflectance data corrected for Rayleigh scattering to establish a cyanobacterial floating algae index (FAI) [*Hu et al.*, 2010]. Index FAI > -0.004 was defined as the threshold value for determining a cyanobacterial bloom. MODIS images with 250 m spatial resolution data were downloaded from the NASA EOS data gateway, which has been available since 2003. The bloom occurrence frequency was defined as the number of images with bloom as a percentage of total number of images. The bloom start dates for each year were calculated as the date when daily chlorophyll *a* concentration (Chl*a*) MODIS-derived reached the background median concentrations +5%[*Shi et al.*, 2019].

2.3 Observations of phytoplankton biomass and water quality parameters as well as riverine external loading

Phytoplankton biomass, indicated as Chl*a* concentration, was monitored monthly along with total nitrogen (TN), total phosphorus (TP) concentrations and pH. There are 14 monthly sampling sites located in the northern lake where cyanobacterial blooms frequently occur due to the summer prevailing southeastern wind (Fig. 1). Monthly observations of Chl*a*, TN, TP concentrations and pH have been conducted by the Taihu Laboratory for Lake Ecosystem Research (TLLER) since 1991. Water samples from the surface, middle and bottom at each site were mixed to represent an integrated water column sample. Dissolved oxygen (DO) concentrations near the bottom were measured every 30 minutes with YSI-6600 sonde which was deployed at the end of pier (ca. 300 m from lakeshore) in TLLER. The measurements were started in 2008 and terminated near the end of 2016 due to equipment failure. Although we have only one site with near-bottom DO observation for exploring the relationship between cyanobacterial

bloom and DO depletion, this location is right in bloom prevailing zone [*Wu et al.*, 2018].

Riverine input of nutrients in Lake Taihu is a dominant fraction of external loading[*Zhai et al.*, 2020]. Riverine loadings of nitrogen and phosphorus have been measured since 2006 by Taihu Basin Authority, Ministry of Water Resources of China, based on observations of flow rate and nutrient concentration of ca. 70 rivers in which 22 large rivers (accounting for ~80% of annual inflow and external nutrient loading)are monitored regularly. Selection of cross-section of river, flow rate measurement instrument and flow rate calculation follow the national standard [*Ministry of Housing and Urban-Rural Development of China (MHURD)*, 2015], which can guarantee the observational deviation of flow rate <10%. Water samples were taken synchronously during flow rate measurement.

Chlorophyll *a* concentration was measured using spectrophotometry at wavelengths of 665 nm and 750 nm, following extraction with hot 90% ethanol[*Lorenzen*, 1967]. TN and TP concentrations were determined by spectrophotometry after digestion with alkaline potassium persulfate[*Ebina et al.*, 1983]. Mean TN, TP and Chl*a* concentrations were obtained by averaging over the northern lake.

2.4 Hydrology and meteorology parameters in association with teleconnective climate indices

External loading is strongly associated with rainfall and runoff [*Daloglu et al.*, 2012; *Williams and King*, 2020], which are intra-annually distributed in three periods, i.e. low precipitation period from January to April (before flood season), the rainy period from May to September (flood season), and the dry period from October to December (dry season). Phytoplankton growth rate is related to the thermal condition

of lake [*Hans W Paerl et al.*, 2011], especially the winter warmth in the subtropical lakes [*J M Deng et al.*, 2014; *Ma et al.*, 2016]. Long-term observations of precipitation and air temperature in the Taihu Basin were collected at the Wuxi Meteorological Station (nearby Lake Taihu).

Local climate anomalies of Taihu basin are strongly influenced by a number of interannual or interdecadal and globally climatological forcings, including El Niño, PDO, AMO, NAO and AO [F Wang and Yang, 2017; Y L Zhu et al., 2011], etc. An El Niño episode is identified by a sea surface temperature (SST) anomaly of 0.5° C above baseline persisting for five months over the equatorial Pacific Ocean (5°N-5°S and 170°W-120°W, known as Niño 3.4 index). The Pacific Decadal Oscillation (PDO) is defined by the leading pattern (EOF) of sea surface temperature (SST) anomalies in the North Pacific basin (typically, polewards of 20°N). The Atlantic Multi-decadal Oscillation (AMO) has been identified as a coherent mode of natural variability occurring in the North Atlantic Ocean. It is based on the average anomalies of sea surface temperatures (SST) in the North Atlantic basin, typically over 0-80°N. The North Atlantic Oscillation (NAO) index is the difference in normalized sea level pressures between Ponta Delgadas (Azores) and Akureyri (Iceland). Arctic oscillation (AO) index is defined as the first leading mode from the EOF analysis of monthly mean height anomalies at 1000-hPa, and normalized by the standard deviation of the monthly index. For statistical telecorrelation analysis, time series of ENSO, PDO, AMO, NAO, and AO indices were collected at https://psl.noaa.gov/data/climateindices/list/.

2.5 Statistical analysis

Long-term trends of MODIS-derived blooms events (bloom areas, frequency, start dates) were evaluated by generalized additive models (GAMs) according to Harding *et al.*, 2016]. GAMs were also used in order to explore the

relationships between climate forcings, nutrients, Chl*a* concentrations and bloom intensities, based on the daily, monthly or yearly observations. The significance level (p) of the statistical tests were examined at two levels: p < 0.05 and p < 0.01. Occurrence probability of yearly mean bloom area, frequency and start date were determined by fitting probability distribution whose density functions (kernel density estimates) were first developed by *density* function in R 3.6.1[*R Core Team*, 2019], and then cumulative probability were calculated based on the probability density functions.

3 Results

3.1 Variation of cyanobacterial blooms during 2006-2018

The bloom extent and frequency in Lake Taihu have increased recently (Fig. 2a, b). The unprecedented intensive blooms occurred in 2017, evidenced by the largest annual mean area and highest frequency of blooms (Fig. 2a, b). A record-setting bloom area was observed as large as 1582 km² on May 16, 2017, covering nearly 70% of the lake surface. In addition, the bloom start date in 2017 was the earliest observed during the 2006 to 2018 period (Fig. 2c). According to cumulative distribution curves (Fig. 2, right panel), the probability of bloom area, frequency and start date in 2017 were 0.002, 0.04 and 0.04, respectively, indicating that the severe blooms in 2017 were extremely rare in this lake during 2006 to 2018. Large bloom areas persisted throughout the summer and autumn, rendering the mean algal blooms in 2017 to be the largest over 2006-2018.

3.2 Changes in nutrient concentrations and external loading during 2006-2018

Based on monthly monitoring of Chla, total nitrogen (TN) and total phosphorus (TP) in the northern lake, Chla was not correlated with TN ($r^2 = 0.005$, p > 0.05)

(Supplementary Fig. 1(a)) but significantly correlated with TP (r^2 =0.3, p < 0.01) (Supplementary Fig. 1(b)). Averaged TN to TP mass ratios of each month over the period 1992-2018 indicated that TN:TP mass ratio decreased from ca. 50 in April to 12.4 in September (Supplementary Fig. 2), which further illustrated that phytoplankton production in Lake Taihu was P limited in spring and P and N colimited during summer and autumn [*Guildford and Hecky*, 2000; *Xu et al.*, 2010]. In addition, TN has declined significantly from 4.82 mg/L to 2.50 mg/L since 2006 (p < 0.01, Fig. 3a), while TP has fluctuated without a trend during 2006 to 2010 and increased after 2010 with a peak in 2017 (Fig. 3b). It is noteworthy that the highest and the second highest riverine loadings of both TN and TP were in 2010 and 2016, respectively (Fig. 3a, 3b), while the yearly mean TP concentration peaked in 2017 at 0.20 mg/L (Fig. 3b), which was much higher than mean concentration 0.14 mg/L over 2006-2018. This suggested that both TN and TP were not responsive to external loading due to the complex influence of in-lake biogeochemical processes such as denitrification, nutrient regeneration and mobilization from sediment (i.e. internal loading).

3.3 Rainfall and temperature anomalies in association with phytoplankton biomass and cyanobacterial blooms in Lake Taihu

Anthropogenic climate change exerts bottom-up influence on phytoplankton community through changes in precipitation that affect riverine nutrient inputs, and increases in temperature that affect cyanobacterial growth rates. Long-term observations of rainfall have shown an increase of annual precipitation since the 1950's in the Taihu basin (Supplementary Fig. S3). A heavy flood occurred in 2016 (Supplementary Fig. S3), resulting in the water level in Lake Taihu reaching its second highest value since the 1950s. Meanwhile, long-term observations indicated that winter and spring surface air temperatures have risen since 1950's (Supplementary Fig. S3). The average winter air temperature of 2016/2017 has been the highest since 1950's (Supplementary Fig. S3), suggesting notable warming during 2016/2017 winter following the flood of 2016.

Correlation analyses suggest that mean TP concentrations during growing seasons (Mar-Aug) were positively related to annual rainfall of the previous year (GAM, r_{adj}^2 = 0.19, p < 0.05, Fig. 4a), and winter/spring temperature anomalies were significantly correlated with annual mean bloom areas (GAM, r_{adj}^2 = 0.47, p < 0.01, Fig. 4b).

4 Discussion

Our observations indicate that Lake Taihu had experienced unprecedented intense cyanobacterial blooms in 2017, illustrated by the large mean area, high frequency and early start date. Intense algal blooms occurred following high external nutrient loading that resulted from highly anomal precipitation and runoff in the previous year (2016) (Supplementary Fig. S4 (b)) and a notable warm preceding winter (2016/2017) (Supplementary Fig. S4 (c)).

4.1 Drivers of the extraordinary blooms in Lake Taihu in 2017

Cyanobacterial blooms in Lake Taihu are mainly driven by nutrient availability and water warmth [*B Qin et al.*, 2019]. After a decadal intensive effluent reduction effort, annual mean TN decreased significantly (Fig. 3(a)), while annual mean TP concentrations in Lake Taihu sustained at high level (Fig. 3(b)), resulting the yearly mean TN:TP ratio continued decline [*B Qin et al.*, 2020b]. Water warmth limits the cyanobacterial growth in Lake Taihu during the winter because of the low water temperature (~5-6° C, supplementray Fig. S5) which is close to the lower limitation for algal growth [*Butterwick et al.*, 2004].

Extreme climatic conditions have increasingly exerted influences on the phytoplankton community and cyanobacterial blooms in Lake Taihu recently[*Yang et al.*, 2016]. Heavy rainfall can induce intense cyanobacterial blooms due to an increase in external loading[*Yang et al.*, 2016]. However, compared to precipitation in the previous year, the preceding winter temperature plays an important role in supporting the severe cyanobacterial blooms in 2017 (Fig. 4). Correlation analysis demonstrated that winter temperature deviation from average significantly affected spring and summer bloom intensity (Fig. 4b) [*Ma et al.*, 2016]. The very warm winter of 2016/2017 resulted in an extensive cyanobacterial bloom (718 km²) on December 31, 2016 [*B Qin et al.*, 2019], which even for Taihu is a rare occasion. An increase in winter water temperature frequently led to an early initiation and proliferation of cyanobacterial blooms [*Duan et al.*, 2009; *Shi et al.*, 2019]. Extremely warm winter months combined with the water diversion from Yangtze River has been blamed for a severe bloom that caused a drinking water crisis in spring 2007 in Wuxi [*B.Q. Qin et al.*, 2010].

As one of Pacific Rim nations, China's climate anomalies are strongly influenced by global-scale interannual variation of climate cycles such as El Niño and Southern Oscillation (ENSO) which are modulated by the Pacific Decadal Oscillation (PDO) [*Heng et al.*, 2020; *L Wang et al.*, 2008], especially rainfall anomalies[*Heng et al.*, 2020], associated with the Atlantic Multidecadal Oscillation (AMO)[*L H Gao et al.*, 2015; *Lyu and Yu*, 2017], North Atlantic Oscillation (NAO) and Arctic Oscillation (AO). These teleconnection patterns are often accompanied by a warm winter, excessive rainfall and flooding throughout Eastern Asia[*L Wang et al.*, 2008]. Climate

anomalies during 2016-2107 in the Taihu basin (Supplementary Fig S4) were an outcome of combined effects imposed by ENSO, AMO and PDO (Fig. 5). High precipitation during 2016 (Supplementary S4 (b)) was likely associated with the strong 2015/2016 El Niño (Fig. 5f) which terminated in spring 2016[WMO, 2017] but its influences extended more than three months later (Supplementary Table S1), and positive (warm) phases of AMO (Fig. 5a) and PDO (Fig. 5b) (Supplementary Table S1), while the persistent warming during 2016/2017, including the notable warm winter 2016/2017 (Supplementary S4 (c)), likely resulted from a number of global scale climate devitations in association with AMO, PDO, as well as NAO and AO (Fig. 5a, 5c, 5d) (Supplementary Table S1). Surprisingly, both AMO and NAO can distantly affect the local climate of Taihu basin, especially the AMO which can affect both surface air temperature (SAT) and precipitation anomalies in the Taihu basin (Fig. 5a, 5b, Supplementary Table S1). Influence from AMO is more significant relative to PDO in the Taihu basin (Fig. 5) (Supplementary Table S1). Consequently, a combination of climate anomalies resulted from the strong 2015/2016 El Niño in association with joint effects of warm phases of AMO and PDO were responsible for the highly frequent and long persistent intense cyanobacterial blooms in 2017.

4.2 Response of Lake Taihu ecosystem to extreme climate anomalies 2016/2017

Ecological responses of aquatic ecosystems to climate changes are increasingly evident [*O'Reilly et al.*, 2015; *van de Waal et al.*, 2010]. However, few studies have addressed ecosystem responses to a combination of extreme climate anomalies, particularly during the 2-year, prolonged warm surface air temperature in the Taihu basin (Supplementary Fig. S4). Such combination of climate anomalies resulted in an increasing amplitude of ecosystem response. P concentrations peaked in 2017, while

external P loading to Lake Taihu was low (Fig. 3), which was likely associated with high internal loading. Because most external P inputs (60-70% of riverine loading) is retained in the sediments [B Qin et al., 2019], legacy P in sediment can be mobilized by positive feedback effects induced by the intense cyanobacterial blooms. The warm winter of 2016/2017 stimulated intense cyanobacterial blooms, which occurred early and persisted through the summer and autumn. Thus, extensive blooms in turn altered physical-chemical conditions, including elevated pH [Y. Gao et al., 2012; Krausfeldt et al., 2019] and decreased dissolved oxygen (DO) [X Yan et al., 2017]. Monthly water quality monitoring revealed that water column pH in the northern lake was significantly related to the Chla concentration when pH > 8.0 (Supplementary Fig. S6, p < 0.001). High-frequency *in-situ* monitoring of near-bottom dissolved oxygen (DO) concentration at the TLLER pier suggested that the DO had decreased since 2008 (Supplementary Fig. S7), and hypoxia frequently took place during low wind velocity periods (Supplementary Fig. S8). Both high pH [Christophoridis and Fytianos, 2006; NiemistÖ et al., 2011] and low DO [Ding et al., 2018; L Zhu et al., 2020] would promote nutrients mineralizing and degrading into dissolved forms in the sediments. Occasional sediment resuspension under windy conditions can liberate significant amounts of soluble P into overlying water [J Deng et al., 2018]. Coupling nutrient mobilization induced by high pH in the water column to DO decline at the lake bottom supports additional cyanobacterial blooms by enhancing internal nutrient cycling [Yonghui Gao et al., 2014] in response to a combination of extreme climate anomalies (Fig. 6). This scenario represents a positive feedback to ensure proliferation and persistence of cyanobacterial blooms in 2017 (Fig. 6). This can also explain why TP in 2017 was 30.6% above predicted value based on temporal trend regression (TP = $0.003 \times \text{Year}$ – 5.813, $r^2 = 0.54$, p < 0.001), while the Chla concentration deviated 79.4% over the

prediction of temporal trend regression (Chla = $1.13 \times \text{Year} - 2236.2$, $r^2 = 0.54$, p < 0.001).

4.3 Implication for eutrophication and bloom management

Eutrophication and cyanobacterial blooms favor shallow lakes and estuaries [B Qin et al., 2020a]. Global increases in excessive nutrient input due to extreme climate events driven by climate change will be likely intensified in magnitude and frequency in the future [Sinha et al., 2017], which will boost cyanobacterial blooms. This is one of the reasons why rehabilitation efforts on many large eutrophic waters have resulted in minimal improvement worldwide [Ho et al., 2019], including Lakes Okeechobee (USA) [Rosen et al., 2017], Winnebago (USA) [Miller and Beversdorf, 2017], Pontchartrain (USA) [Mishra and Mishra, 2010], Kasumigura (Japan) [Salem et al., 2017], Champlain (shallow northern part, USA/Canada) [Fortin et al., 2015], Winnipeg (Canada) [Ulrich et al., 2016], Erie (shallow western part, USA) [Bullerjahn et al., 2016], Peipsi (Eastonia) [Alikas et al., 2015], Lake Victoria (Africa) [Mbonde et al., 2015] and Chaohu (China) [Zhang et al., 2016]. More intense cyanobacterial blooms will, in turn, further deteriorate water quality by decreasing clarity and transparency, reducing submersed macrophyte coverage, increasing hypoxic area. This scenario leads to impairment of aquatic ecosystem functions and services, including drinking water supplies, loss of fisheries and recreational/aesthetic uses, etc.

The current warming trend has been documented on a global scale and is a known manifestation of the greenhouse effect, which is reflected in rising global temperatures since the 1950s[*IPCC*, 2013]. In addition to global warming, both the amplitude and the frequency of regional interannual climate variations in association with global scale external forcings such as El Niño, PDO and AMO have likely increased with

widespread climatic (e.g., changes in precipitation and surface air temperature) consequences[*Cai et al.*, 2015]. Lake Taihu is a mirror for global accelerated eutrophication and cyanobacterial proliferation[*Huisman et al.*, 2018], resulting from anthropogenic nutrient discharge and the combined regional warming and heavy precipitation exerted by a changing climate[*Hans W. Paerl et al.*, 2020]. Extensive lake restoration initiated after the drinking water crisis in 2007 is likely challenged by these climatic changes. A combination of extreme climate anomalies cause magified responses of highly productive aquatic ecosystem, which ultimately mobilizes phosphrous from the sediment and promotes algal blooms by forming a positive feedback to exacerbate the water quality. Such ecosystem responses may dampen the effect of external loading reduction and lengthen the restoration time. Considering the profound influences exerted by the extreme climate anomalies during 2016 to 2017, new targets for nutrient reduction control strategies are urgently needed, especially for shallow and eutrophic systems.

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References

Alikas, K., K. Kangro, R. Randoja, P. Philipson, E. Asuküll, J. Pisek, and A. Noorma (2015), Satellite-based products for monitoring optically complex inland waters in support of EU Water Framework Directive, *Int J Remote Sens*, *36*, 4446-4468.

Brookes, J. D., and C. C. Carey (2011), Resilience to blooms, Science, 334(6052), 46-47.

Bullerjahn, G. S., et al. (2016), Global solutions to regional problems: Collecting global expertise to address the problem of harmful cyanobacterial blooms. A Lake Erie case study, *Harmful Algae*, *54*, 223-238.

Butterwick, C., S. I. Heaney, and J. F. Talling (2004), Diversity in the influence of temperature on the growth rates of freshwater algae, and its ecological relevance, *Freshwater Biol*, *50*(2), 291-300.

Cai, W., et al. (2015), ENSO and greenhouse warming, Nature Climate Change, 5(9), 849-859.

Christophoridis, C., and K. Fytianos (2006), Conditions affecting the release of phosphorus from surface lake sediments, *Journal of Environmental Quality*, *35*(4), 1181-1192.

Daloglu, I., K. H. Cho, and D. Scavia (2012), Evaluating Causes of Trends in Long-Term Dissolved Reactive Phosphorus Loads to Lake Erie, *Environ Sci Technol*, *46*(19), 10660-10666.

Deng, J., H. W. Paerl, B. Qin, Y. Zhang, G. Zhu, E. Jeppesen, Y. Cai, and H. Xu (2018), Climaticallymodulated decline in wind speed may strongly affect eutrophication in shallow lakes, *Sci Total Environ*, *645*, 1361-1370.

Deng, J. M., B. Q. Qin, H. W. Paerl, Y. L. Zhang, J. R. Ma, and Y. W. Chen (2014), Earlier and warmer springs increase cyanobacterial (Microcystis spp.) blooms in subtropical Lake Taihu, China, *Freshwater Biol*, *59*(5), 1076-1085.

Ding, S., M. Chen, M. Gong, X. Fan, B. Qin, H. Xu, S. Gao, Z. Jin, D. C. W. Tsang, and C. Zhang (2018), Internal phosphorus loading from sediments causes seasonal nitrogen limitation for harmful algal blooms, *Sci Total Environ*, *625*, 872-884.

Duan, H., R. Ma, X. Xu, F. Kong, S. Zhang, W. Kong, J. Hao, and L. Shang (2009), Two-decade reconstruction of algal blooms in China's Lake Taihu, *Environ Sci Technol*, 43(10), 3522-3528.

Ebina, J., T. Tsutsui, and T. Shirai (1983), Simultaneous determination of total nitrogen and total phosphorus in water using peroxodisulfate oxidation, *Water Res*, *17*(12), 1721-1726.

Fortin, N., V. Munoz-Ramos, D. Bird, B. Lévesque, L. G. Whyte, and C. W. Greer (2015), Toxic cyanobacterial bloom triggers in missisquoi bay, lake champlain, as determined by next-generation sequencing and quantitative PCR, *Life (Basel, Switzerland)*, *5*(2), 1346-1380.

Gao, L. H., Z. W. Yan, and X. W. Quan (2015), Observed and SST-forced multidecadal variability in global land surface air temperature, *Climate Dynamics*, *44*(1), 359-369.

Gao, Y., J. C. Cornwell, D. K. Stoecker, and M. S. Owens (2012), Effects of cyanobacterial-driven pH increases on sediment nutrient fluxes and coupled nitrification-denitrification in a shallow fresh water estuary, *Biogeosciences*, *9*(7), 2697-2710.

Gao, Y., J. C. Cornwell, Stoecker, Diane K., and M. S. Owens (2014), Influence of cyanobacteria blooms on sediment biogeochemistry and nutrient fluxes, *Limnol Oceanogr*, *59*(3), 959-971.

Guan, Q., L. Feng, X. Hou, G. Schurgers, Y. Zheng, and J. Tang (2020), Eutrophication changes in fifty large lakes on the Yangtze Plain of China derived from MERIS and OLCI observations, *Remote Sens*

Environ, 246, 111890.

Guildford, S. J., and R. E. Hecky (2000), Total nitrogen, total phosphorus, and nutrient limitation in lakes and oceans: Is there a common relationship?, *Limnol Oceanogr*, *45*(6), 1213-1223.

Guo, L. (2007), Doing Battle with the Green Monster of Taihu Lake, Science, 317(5842), 1166.

Harding, L. W., C. L. Gallegos, E. S. Perry, W. D. Miller, J. E. Adolf, M. E. Mallonee, and H. W. Paerl (2016), Long-term trends of nutrients and phytoplankton in Chesapeake Bay, *Estuaries and Coasts, 39*(3), 664-681.

Heng, C., T. Lee, J.-S. Kim, and L. Xiong (2020), Influence analysis of central and Eastern Pacific El Niños to seasonal rainfall patterns over China using the intentional statistical simulations, *Atmospheric Research*, 233, 104706.

Ho, J. C., A. M. Michalak, and N. Pahlevan (2019), Widespread global increase in intense lake phytoplankton blooms since the 1980s, *Nature*.

Hu, C., L. Zhongping, R. Ma, K. Yu, D. Li, and S. Shang (2010), Moderate Resolution Imaging Spectroradiometer (MODIS) observations of cyanobacteria blooms in Taihu Lake, China, *Journal of Geophysical Research Oceans*, *115*(C4), -.

Huisman, J., G. A. Codd, H. W. Paerl, B. W. Ibelings, J. M. H. Verspagen, and P. M. Visser (2018), Cyanobacterial blooms, *Nature Reviews Microbiology*, *16*(8), 471-483.

IPCC (2013), Summary for Policymakers, in *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*, edited by T. F. Stocker, D. Qin, G. K. Plattner, M. Tignor, S. K. Allen, J. Boschung, A. Nauels, Y. Xia, V. Bex and P. M. Midgley, Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA.

Krausfeldt, L. E., A. T. Farmer, H. F. Castro Gonzalez, B. N. Zepernick, S. R. Campagna, and S. W. Wilhelm (2019), Urea Is Both a Carbon and Nitrogen Source for Microcystis aeruginosa: Tracking (13)C Incorporation at Bloom pH Conditions, *Front Microbiol*, *10*, 1064.

Li, J., F. Zheng, C. Sun, J. Feng, and J. Wang (2019), Pathways of Influence of the Northern Hemisphere Mid-high Latitudes on East Asian Climate: A Review, *Adv Atmos Sci*, *36*(9), 902-921.

Lorenzen, C. J. (1967), Determination Of Chlorophyll And Pheo-Pigments: Spectrophotometric Equations, *Limnol Oceanogr*, *12*(2), 343-346.

Lyu, K., and J.-Y. Yu (2017), Climate impacts of the Atlantic Multidecadal Oscillation simulated in the CMIP5 models: A re-evaluation based on a revised index, *Geophysical Research Letters*, 44(8), 3867-3876.

Ma, J. R., et al. (2016), The persistence of cyanobacterial (Microcystis spp.) blooms throughout winter in Lake Taihu, China, *Limnol Oceanogr*, *61*(2), 711-722.

Mbonde, A., L. Sitoki, and R. Kurmayer (2015), Phytoplankton composition and microcystin concentrations in open and closed bays of Lake Victoria, Tanzania, *Aquatic Ecosystem Health & Management*, 18.

Miller, T., and L. Beversdorf (2017), Variable Cyanobacterial Toxin and Metabolite Profiles across Six Eutrophic Lakes of Differing Physiochemical Characteristics, *Toxins*, *9*, 62.

Ministry of Housing and Urban-Rural Development of China (MHURD) (2015), Code for liquid flow measurement in open channels, http://www.mohurd.gov.cn/wjfb/201510/t20151021_225294.html, Beijing.

Mishra, D., and S. Mishra (2010), Plume and bloom: effect of the Mississippi River diversion on the water quality of Lake Pontchartrain, *Geocarto International*, *25*, 555-568.

NiemistÖ, J., H. Holmroos, and J. Horppila (2011), Water pH and sediment resuspension regulating internal phosphorus loading in a shallow lake – field experiment on diurnal variation, *J Limnol*, *70*(1), 3. O'Reilly, C. M., et al. (2015), Rapid and highly variable warming of lake surface waters around the globe, *Geophysical Research Letters*, *42*(24), 10,773-710,781.

Paerl, H. W., N. S. Hall, and E. S. Calandrino (2011), Controlling harmful cyanobacterial blooms in a world experiencing anthropogenic and climatic-induced change, *Sci Total Environ*, *409*(10), 1739-1745.

Paerl, H. W., K. E. Havens, N. S. Hall, T. G. Otten, M. Zhu, H. Xu, G. Zhu, and B. Qin (2020), Mitigating a global expansion of toxic cyanobacterial blooms: confounding effects and challenges posed by climate change, *Marine and Freshwater Research*, *71*(5), 579-592.

Qin, B., P. Xu, Q. Wu, L. Luo, and Y. Zhang (2007), Environmental issues of lake Taihu, China, *Hydrobiologia*, *581*(1), 3-14.

Qin, B., J. Zhou, J. J. Elser, W. S. Gardner, J. Deng, and J. D. Brookes (2020a), Water Depth Underpins the Relative Roles and Fates of Nitrogen and Phosphorus in Lakes, *Environ Sci Technol*, *54*(6), 3191-3198. Qin, B., H. W. Paerl, J. D. Brookes, J. Liu, E. Jeppesen, G. Zhu, Y. Zhang, H. Xu, K. Shi, and J. Deng (2019), Why Lake Taihu continues to be plagued with cyanobacterial blooms through 10 years (2007–2017) efforts, *Science Bulletin*.

Qin, B., et al. (2020b), Are nitrogen-to-phosphorus ratios of Chinese lakes actually increasing?, *Proceedings of the National Academy of Sciences*, *117*(35), 21000.

Qin, B. Q., G. W. Zhu, G. Gao, Y. L. Zhang, W. Li, H. W. Paerl, and W. W. Carmichael (2010), A drinking water crisis in Lake Taihu, China: linkage to climatic variability and lake management, *Environ Manage*, *45*(1), 105-112.

Qin, B. Q., G. Gao, G. W. Zhu, Y. L. Zhang, Y. Z. Song, X. M. Tang, H. Xu, and J. M. Deng (2013), Lake eutrophication and its ecosystem response, *Chinese Sci Bull*, *58*(9), 961-970.

R Core Team (2019), R: A language and environment for statistical computing.

Rigosi, A., C. C. Carey, B. W. Ibelings, and J. D. Brookes (2014), The interaction between climate warming and eutrophication to promote cyanobacteria is dependent on trophic state and varies among taxa, *Limnol Oceanogr*, *59*(1), 99-114.

Rosen, B. H., T. W. Davis, C. J. Gobler, B. J. Kramer, and K. A. Loftin (2017), Cyanobacteria of the 2016 Lake Okeechobee and Okeechobee Waterway harmful algal bloom, *Report Rep. 2017-1054*, Reston, VA. Salem, S., M. Strand, H. Higa, H. Kim, K. Kazuhiro, K. Oki, and T. Oki (2017), Evaluation of MERIS Chlorophyll-a Retrieval Processors in a Complex Turbid Lake Kasumigaura over a 10-Year Mission, *Remote Sens-Basel*, *9*, 1022.

Shi, K., Y. Zhang, Y. Zhou, X. Liu, G. Zhu, B. Qin, and G. Gao (2017), Long-term MODIS observations of cyanobacterial dynamics in Lake Taihu: Responses to nutrient enrichment and meteorological factors, *Sci Rep-Uk*, *7*, 40326.

Shi, K., Y. Zhang, Y. Zhang, N. Li, B. Qin, G. Zhu, and Y. Zhou (2019), Phenology of Phytoplankton Blooms in a Trophic Lake Observed from Long-Term MODIS Data, *Environ Sci Technol*, *53*(5), 2324-2331.

Sinha, E., A. Michalak, and V. Balaji (2017), Eutrophication will increase during the 21st century as a result of precipitation changes, *Science*, *357*(6349), 405-408.

Ulrich, A. E., D. F. Malley, and P. D. Watts (2016), Lake Winnipeg Basin: Advocacy, challenges and

progress for sustainable phosphorus and eutrophication control, *Sci Total Environ*, *542*, 1030-1039. van de Waal, D. B., A. M. Verschoor, J. M. Verspagen, E. van Donk, and J. Huisman (2010), Climatedriven changes in the ecological stoichiometry of aquatic ecosystems, *Frontiers in Ecology and the Environment*, *8*(3), 145-152.

Wang, F., and S. Yang (2017), Regional characteristics of long-term changes in total and extreme precipitations over China and their links to atmospheric-oceanic features, *Int J Climatol*, *37*(2), 751-769. Wang, L., W. Chen, and R. Huang (2008), Interdecadal modulation of PDO on the impact of ENSO on the east Asian winter monsoon, *Geophysical Research Letters*, *35*(20).

Wang, S., et al. (2018), Trophic state assessment of global inland waters using a MODIS-derived Forel-Ule index, *Remote Sens Environ*, *217*, 444-460.

Wang, S. M., and H. S. dou (1998), Annals of Lakes in China Science Press, Beijing.

Williams, M. R., and K. W. King (2020), Changing Rainfall Patterns Over the Western Lake Erie Basin (1975–2017): Effects on Tributary Discharge and Phosphorus Load, *Water Resour Res*, *56*(3), e2019WR025985.

WMO, W. M. O. (2017), WMO Statement on the State of the Global Climate in 2016, Geneva, Switzerland.

Wu, T. F., B. Q. Qin, W. H. Ding, G. W. Zhu, Y. L. Zhang, G. Gao, H. Xu, W. Li, B. L. Dong, and L. C. Luo (2018), Field Observation of Different Wind-Induced Basin-Scale Current Field Dynamics in a Large, Polymictic, Eutrophic Lake, *J Geophys Res-Oceans*, *123*(9), 6945-6961.

Xu, H., H. W. Paerl, B. Qin, G. Zhu, and G. Gaoa (2010), Nitrogen and phosphorus inputs control phytoplankton growth in eutrophic Lake Taihu, China, *Limnol Oceanogr*, *55*(1), 420-432.

Yan, X., X. Xu, M. Wang, G. Wang, S. Wu, Z. Li, H. Sun, A. Shi, and Y. Yang (2017), Climate warming and cyanobacteria blooms: Looks at their relationships from a new perspective, *Water Res*, *125*, 449-457.

Yan, Z., Y. Ding, P. Zhai, L. Song, L. Cao, and Z. Li (2020), Re-Assessing Climatic Warming in China since 1900, *Journal of Meteorological Research*, *34*(2), 243-251.

Yang, Z., M. Zhang, X. Shi, F. Kong, R. Ma, and Y. Yu (2016), Nutrient reduction magnifies the impact of extreme weather on cyanobacterial bloom formation in large shallow Lake Taihu (China), *Water Res*, *103*, 302-310.

Zhai, S. H., Y. Zhou, Y. H. Cheng, J. Cai, and Y. W. Hu (2020), Calculation of Total Phosphorus Loads from Rivers around Lake Taihu and Analysis of Total Phosphorus Fluctuation in the Lake in 2015-2016, *Journal of Lake Science*, *32*(1), 48-57.

Zhang, M., Y. Zhang, Z. Yang, L. Wei, W. Yang, C. Chen, and F. Kong (2016), Spatial and seasonal shifts in bloom-forming cyanobacteria in Lake Chaohu: Patterns and driving factors, *Phycological Research*, *64*(1), 44-55.

Zhu, L., W. Shi, B. Van Dam, L. Kong, J. Yu, and B. Qin (2020), Algal Accumulation Decreases Sediment Nitrogen Removal by Uncoupling Nitrification-Denitrification in Shallow Eutrophic Lakes, *Environ Sci Technol*, *54*(10), 6194-6201.

Zhu, Y. L., H. J. Wang, W. Zhou, and J. H. Ma (2011), Recent changes in the summer precipitation pattern in East China and the background circulation, *Climate Dynamics*, *36*(7-8), 1463-1473.

Figure caption

Fig. 1 Location of Lake Taihu (main part of map), monthly sampling sites in the northern lake (red dot, low left corner) and location of Taihu Laboratory for Lake Ecosystem Research (TLLER), Chinese Academy of Science, and location of Lake Taihu and defined area of Nino 3.4 at equatorial Pacific (upper left coner)

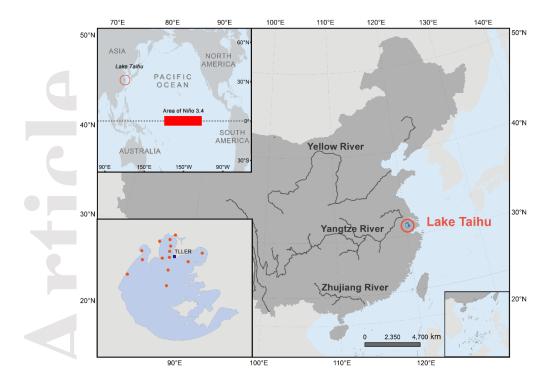
Fig. 2 Bloom events in Lake Taihu from 2006 to 2018. (a) mean surface areas of cyanobacterial blooms derived from MODIS satellite imagery; (b) annual occurrence frequency (the number of bloom images in percentage of total detectable images); (c) annual bloom start dates (Julian day). Solid lines in the left panels indicate long-term trends estimated by generalized additive models (GAMs, N = 13), and shading indicates the standard error of the estimates. The cumulative probability for bloom area, frequency and start date are presented in the right panels. The shadows indicated cumulative probability range from 0.05 to 0.95. The red dash lines are the 2017 value.

Fig. 3 Annual mean TN and TP concentrations in the water column (blue line) and annual riverine external loading (orange line) to Lake Taihu from 2006 to 2018. (a) TN concentrations and riverine loading N; (b) TP concentrations and riverine loading TP.

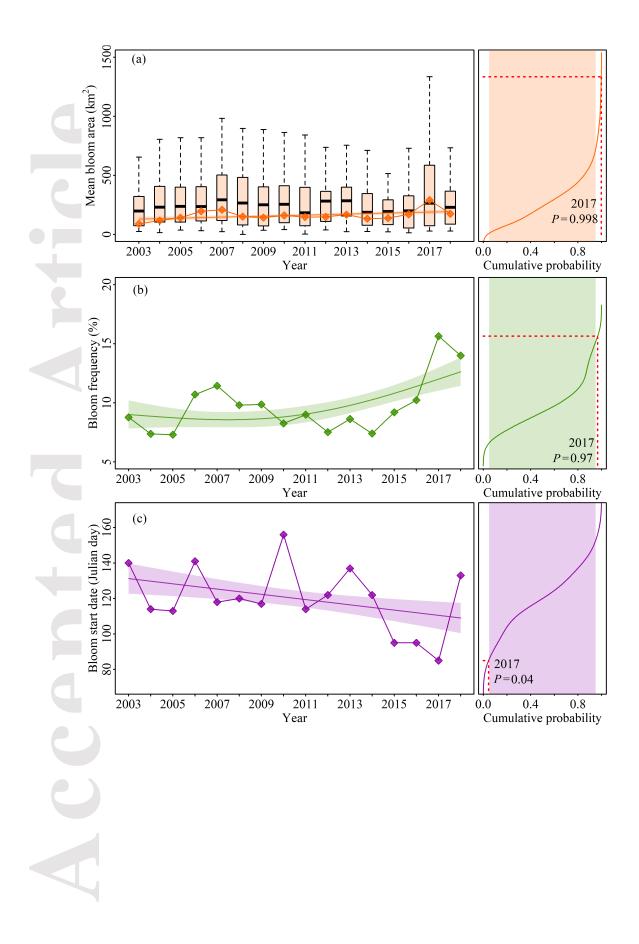
Fig. 4 Correlation between annual rainfall of previous year and phosphorus availability of growing season (Mar-Aug) in Lake Taihu (r^2 =0.19, p < 0.05) (a), and winter/spring temperature deviation from average vs annual mean bloom areas retrieved from satellite imagery (r^2 =0.47, p < 0.01) (b). Some extreme responses of TP concentrations and bloom areas were marked by individuals years. Relationships were estimated by generalized additive models, and shading indicates the standard error of the estimates.

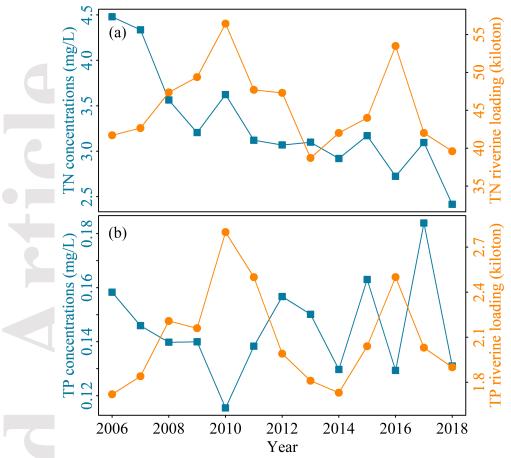
Fig. 5 Pearson correlation between surface air temperature (SAT, Wuxi) with AMO (a), PDO (c) and ENSO (e), and precipitation (Wuxi) with AMO (b), PDO (d) and ENSO (f), based on 12 month moving average over period from 1955 to 2018.

Fig. 6 Conceptual diagram of a combination of climate (high precipitation and warm winter) anomalies induced internal phosphorus cycling through boosting cyanobacterial blooms to increase pH and decrease DO, which led to a positive feedback to ensure cyanobacterial bloom persistence



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