

1 **Capturing the spatial variability of algal bloom development in a shallow temperate lake**

2 **Authors:** David Ortiz<sup>1,2\*</sup>, Grace Wilkinson<sup>1,2</sup>

3 <sup>1</sup>Department of Ecology, Evolution, and Organismal Biology, Iowa State University, Ames,  
4 Iowa USA 50010

5 <sup>2</sup>Current Address: Center for Limnology, University of Wisconsin – Madison, Madison,  
6 Wisconsin USA 53706

7 \*Corresponding Author: dortiz4@wisc.edu

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10

11 **Abstract**

12 1. Algal blooms can have profound effects on the structure and function of aquatic  
13 ecosystems and have the potential to interrupt valuable ecosystem services. Despite the  
14 potential ecological and economic consequences of algal blooms, the spatial dynamics of  
15 bloom development in spatially complex ecosystems such as shallow lakes remain poorly  
16 characterized. Our goal was to evaluate the magnitude and drivers of spatial variability of  
17 algal biomass, dissolved oxygen and pH over the course of a season, in a shallow lake in  
18 order to better understand the spatial dynamics of algal blooms in these ecosystems.

19 2. We sampled 98 locations in a small eutrophic lake on a 65m grid for several parameters  
20 (chlorophyll *a*, phycocyanin, dissolved oxygen, pH, and temperature), weekly over 122  
21 days. This was done to estimate the dynamics of variability and spatial autocorrelation  
22 during the course of multiple bloom events. We also compared the spatial measurements  
23 to a high frequency sensor deployed at a fixed station and estimated the optimal spatial  
24 sampling resolution by performing a rarefaction analysis.

- 25 3. Spatial heterogeneity of algal pigments was high, particularly during bloom events, and  
26 this pattern and the overall severity of the bloom was not well captured with the fixed  
27 station monitoring. The pattern of algal pigments and other limnologically important  
28 variables (dissolved oxygen and pH) was related to the direction of prevailing winds 24  
29 hours prior to sampling and likely enhanced by the shallow northern basin of the lake  
30 where the main surface inlet was located. Additionally, a dense bed of floating-leaf  
31 macrophytes contributed to local patchiness in all variables. Finally, from the rarefaction  
32 analysis we found that minimal information about the mean state of the ecosystem was  
33 gained after ~30 locations had been sampled.
- 34 4. This study revealed how spatially heterogeneous shallow lakes are over the course of a  
35 single season, and that the magnitude of variability was highest during biologically -  
36 intensive periods such as algal blooms. As such, continued research is needed across a  
37 range of trophic conditions to better understand the structure of horizontal variability in  
38 lakes. Overall, these data demonstrate the need for spatially-explicit monitoring to better  
39 understand the dynamics and drivers of algal blooms in shallow lakes and to better  
40 manage ecosystem services.

41 **Introduction**

42           Lakes are highly dynamic ecosystems that can undergo rapid physical and chemical  
43 changes at an individual location, throughout their water column, and across the entire lake  
44 surface at the scale of hours, days, seasons, and years (Laas et al., 2012; Read et al., 2011;  
45 Wynne & Stumpf, 2015). Quantifying heterogeneity in aquatic ecosystem structure and function  
46 not only improves our understanding of lake ecology and the underlying mechanisms that drive  
47 spatial and temporal heterogeneity, but also provides insights that improve management of these  
48 ecosystems and the services they provide. With the development of sophisticated sensor  
49 technology, high frequency measurements of variables such as dissolved oxygen and temperature  
50 have helped limnologists grasp the scale of temporal heterogeneity in lakes (Carpenter et al.,  
51 2020; Chaffin et al., 2020; Cotterill et al., 2019). Detailed temporal monitoring has led to  
52 advances in understanding several lake mechanisms such as diel cycles in primary production  
53 (Solomon et al., 2013; Staehr et al., 2012), temperature effects on biogeochemical processes  
54 (Medeiros et al., 2012), and early warnings of the transition to alternative stable states (Carpenter  
55 et al., 2011; Wilkinson et al., 2018). Additionally, high frequency measurements have been used  
56 to better understand heterogeneity over depth (vertical spatial heterogeneity) for important  
57 processes such as stratification (Boehrer & Schultze, 2008; Read et al., 2011). Despite these  
58 advances in understanding temporal and vertical heterogeneity, less is known about the dynamics  
59 of horizontal spatial heterogeneity in the surface waters of lakes.

60           The vast majority of our understanding of lentic ecosystem structure and function comes  
61 from single station sampling, with measurements taken through time over the deepest point in  
62 the lake (Stanley et al., 2019). This location is usually selected to be representative of conditions  
63 in the lake; however, the representativeness of a single location is likely to vary with regards to

64 the variable being measured and with time due to interacting forces such as wind, hydrology,  
65 bathymetry, and biology (Chaffin et al., 2020; Schilder et al., 2013; Wu et al., 2010; Zhou.,  
66 2013). For example, ecosystem metabolism measured at dozens of locations for 10 days in two  
67 north temperate lakes varied 1-2 orders of magnitude, with more than three-quarters of the  
68 variability attributable to the measurement location within the lake (Van de Bogert et al., 2012).  
69 Transect-based studies of reservoirs have revealed gradients in algae pigments, pH, and nutrients  
70 with differences varying between 25%-180% within a waterbody (Moreno-Ostos et al., 2009;  
71 Rychtecky & Znachor, 2011; Smith, 2018). Recently, satellite-based studies have demonstrated  
72 the ability to detect spatial patterns at a high resolution for optical variables in large lakes (Lekki  
73 et al., 2019). Despite these advances, relatively few studies have quantified horizontal spatial  
74 variability over time in lakes (Buttita et al. 2017, Vilas et al. 2017, Loken et al. 2019), hampering  
75 our understanding of the magnitude of heterogeneity in variables important for managing water  
76 quality and ecosystem services.

77         The development of algal blooms is expected to be a spatially heterogeneous  
78 phenomenon (Buelo et al., 2018; Butitta, Carpenter et al., 2017; Serizawa et al., 2008) due to  
79 both local heterogeneity in nutrient limitation, zooplankton grazing, and temperature (Davis et  
80 al., 2009; Hansen et al., 1997) and population scale heterogeneity due to wind (George &  
81 Heaney, 1978). Algal blooms can have a negative effect on ecosystem services, and therefore are  
82 often a target for ecosystem monitoring and management. Some bloom-forming taxa,  
83 particularly freshwater cyanobacteria, can produce toxins that rise to dangerous concentrations  
84 for humans, pets, and livestock (Codd et al., 2005; Corbel et al., 2014). Additionally, the  
85 mineralization of settling phytoplankton contributes to anoxic bottom waters, while intense  
86 periods of primary production cause large variation in dissolved oxygen and pH (in poorly

87 buffered ecosystems) over the course of the day, which is stressful for aquatic organisms  
88 (Gilbert, 2017; Landsberg, 2002). Furthermore, the perceived recreational value of lakes declines  
89 when blooms form (Angradi et al., 2018), which in turn can negatively affect local economies  
90 (Dodds et al., 2009). Despite the risk of economic loss, loss in biodiversity, and potential human  
91 harm, the spatial dynamics of bloom development in spatially complex ecosystems such as  
92 shallow lakes remain poorly characterized.

93         Shallow lakes have a large interface between the sediment and water relative to deeper  
94 lakes, making them more susceptible to rapid changes in water residence time and nutrient inputs  
95 (Christensen et al., 2013; Rennella & Quiros, 2006; Romo et al., 2013). Due to the expansive  
96 littoral zones, shallow lakes can have large macrophyte beds which modify the light climate and  
97 turbulence at the sediment-water interface (Andersen et al., 2017; Moller & Rordam, 1985; Vilas  
98 et al., 2017). Many shallow lakes are also polymictic, experiencing multiple periods of  
99 stratification followed by mixing during the ice-free season. During periods of water column  
100 stability, some cyanobacteria taxa thrive, initiating blooms (Carey et al., 2012). Additionally,  
101 episodic nutrient loading from the watershed during storm events (Carpenter et al., 2015; Kelly  
102 et al., 2019), spatial gradients in nutrient availability due to stream inlets and morphology (e.g.  
103 embayments), and wind-driven circulation (Schoen et al., 2014) can all contribute to spatial  
104 heterogeneity of algal blooms over time in shallow lakes.

105         In order to better understand the spatial dynamics of algal blooms in shallow lakes, we  
106 performed intensive spatial sampling on Swan Lake (Iowa, USA), a spatially complex, shallow,  
107 hypereutrophic waterbody with a history of toxic cyanobacteria algal blooms. In addition to  
108 measuring algal pigments throughout the lake over the course of 122 days, we also measured  
109 temperature, dissolved oxygen, and pH. The spatial sampling captured two bloom events and

110 coincided with high frequency monitoring of the same variables using autonomous sensors  
111 deployed at a fixed station (Ortiz et al. 2020). Using these data, we addressed the following  
112 questions: 1) how does spatial variability of algae, dissolved oxygen, and pH change over the  
113 course of a season, 2) are high frequency measurements at a fixed station an adequate  
114 characterization of surface water dynamics in a shallow lake, and 3) what is the optimal spatial  
115 sampling frequency to capture the mean state of a productive waterbody? Evaluating these  
116 questions with data from a spatially complex, hypereutrophic lake will provide valuable  
117 ecological and management-relevant insights into algal bloom dynamics.

118

## 119 **Methods**

### 120 *Study Site*

121 Swan Lake (42.0396, -94.8454) has an average depth of 2 m, surface area of 40.5  
122 hectares, and a shoreline development index value of 1.54. The watershed is 350 hectares with  
123 92% of the land in agricultural use. The estimated water residence time is approximately 1.5  
124 years. During the ice-free period of 2018, Swan Lake had an average total phosphorus  
125 concentration of  $280 \mu\text{g L}^{-1}$  and a total nitrogen concentration of  $1.61 \text{ mg L}^{-1}$ , making it  
126 hypereutrophic (Carlson, 1977). Total nitrogen was measured as the sum of total Kjeldahl  
127 nitrogen (method 351.2 v2, US EPA, 1993c) and nitrate + nitrite measured using the cadmium  
128 reduction method (method 4500-NO<sub>3</sub>-F, US EPA, 1993a). Total phosphorus was measured using  
129 the ascorbic acid method (method 365.1 v2, US EPA, 1993b). The average total alkalinity during  
130 the same period was  $139 \text{ mg CaCO}_3 \text{ L}^{-1}$  determined through end point titration (APHA, 1998). In  
131 addition to seasonal algal blooms, Swan Lake also has non-continuous beds of American lotus  
132 (*Nelumbo lutea*) and sago pondweed (*Stuckenia pectinata*) that peak in biomass in the latter half

133 of the summer and then begin senescing. The main surface inlet to the lake enters on the western  
134 side and the outlet is at the southern edge of the waterbody (Figure 1). There are no known  
135 springs feeding the lake.

136

### 137 *Field Methods*

138         The spatial sampling occurred approximately weekly from day of year (DOY) 142 to  
139 DOY 264, encompassing the late spring, summer, and early autumn. A total of 16 spatial  
140 sampling events occurred over the course of the 122 days. Measurements of chlorophyll *a*,  
141 phycocyanin, temperature, dissolved oxygen saturation and pH were taken 0.25 m below the  
142 surface at 98 sampling stations using a YSI Pro DSS multiparameter sonde (Yellow Springs  
143 Instrument, Yellow Springs, OH) suspended over the side of a 3-meter long jon boat equipped  
144 with an outboard motor. The sensors, which included the fluorometric Total Algae (chlorophyll *a*  
145 and phycocyanin), optical dissolved oxygen, and Ag/AgCl pH sensors, were calibrated weekly  
146 prior to each sampling event according to manufacturer instructions. The sampling stations were  
147 laid out in a 65 x 65 m grid across the lake (Figure 1) with each location measured in the same  
148 order (north to south) for each sampling event. This spatial resolution was selected to allow for  
149 many sampling locations to be measured in a relatively short window of time, thereby  
150 minimizing the chance that the differences observed between sampling locations was not due to  
151 time of day. Measurements were taken between 10:00 and 14:00, except for the first two and last  
152 three weeks when sampling lasted until 16:00. Beginning on DOY 177 when submerged  
153 macrophytes could be identified from the jon boat, the presence or absence of submerged or  
154 floating leaf macrophytes was noted at each sampling station during each sampling event. These  
155 weekly presence/absence data were used to construct the macrophyte distributions in Figure 1 .

156           The fixed station high frequency monitoring of Swan Lake was performed using a YSI  
157 EXO2 (Yellow Springs Instrument, Yellow Springs, OH) multiparameter sonde equipped with  
158 the same sensors as the YSI ProDSS used for the spatial sampling. The sonde recorded  
159 measurements of chlorophyll *a*, phycocyanin, dissolved oxygen saturation, and pH every 15  
160 minutes. The instrument was deployed on DOY 135 over the deepest point in the lake (3.8 m  
161 deep), hanging approximately 0.5m below the surface, and removed on DOY 264 after the  
162 spatial sampling event on that day. The fixed station sonde was monitored weekly for drift and  
163 calibrated according to manufacturer instructions when indicated by the quality control algorithm  
164 in the KorEXO software. Hourly precipitation, wind speed, and wind direction were collected at  
165 the Arthur N. Neu Airport in Carroll, Iowa, located 4.5 km from the lake, as a part of the  
166 National Oceanic and Atmospheric Automated Surface Observatory System. The meteorological  
167 data were used to aid in the interpretation of spatial dynamics over the course of the summer.

168

#### 169 *Data analysis*

170           Spatial heterogeneity can be quantified by calculating the spatial variance (e.g.,  
171 coefficient of variation; CV) or spatial autocorrelation (Moran's I, Moran, 1950). Increasing  
172 spatial variance is indicative of increasing patchiness in the ecosystem, such as areas of high-  
173 density algal biomass and areas of low-density biomass within a lake. Spatial autocorrelation  
174 accounts for the location of those patches within the ecosystem in relationship to each other.  
175 Local Moran's I quantifies how similar the abundance of algae is at one location compared to the  
176 density of surrounding neighbors. When measured over time for variables that are indices of  
177 algal biomass (e.g., the pigments chlorophyll *a* and phycocyanin), both of these metrics of spatial  
178 heterogeneity can provide insight into the dynamics of algal bloom development. In models of



179 algal blooms, both spatial variance and autocorrelation are expected to be high during the bloom  
180 period (Buelo et al., 2018).

181 Spatial autocorrelation (AC) and the coefficient of variation (CV) were calculated for  
182 each variable on each sampling date in order to evaluate the dynamics of these parameters over  
183 time. Prior to analysis, extreme outliers in the algal pigments were removed from the spatial  
184 dataset as they were well outside the operating range of the Total Algae sensor or there was  
185 known interference with the sensor resulting in an inaccurate measurement. This resulted in five  
186 chlorophyll *a* and three phycocyanin measurements being removed out of 3,136 total pigment  
187 measurements. The spatial CV is the standard deviation of all of the spatial measurements for a  
188 variable on a given sampling date divided by the mean of those measurements, expressed as a  
189 percent. Spatial AC was calculated as the average value of local Moran's I with a queen's  
190 distance neighbor list (92 meters) with equal weight (1/n) on neighbors, as to not impose any  
191 assumptions on possible spatial patterns in the variables. We limited our analysis to surrounding  
192 neighbors because distances beyond this have not shown high spatial autocorrelation of algal  
193 pigments under experimental conditions (Butitta et al. 2017). Local Moran's I values near 1.0  
194 reflect high spatial AC within neighbors, zero indicates a random distribution, whereas spatial  
195 AC values nearing -1.0 indicate a perfectly dispersed distribution (e.g. checkerboard pattern) in  
196 the variable being measured. As the spatial variability in temperature is mediated by physical  
197 processes, we used the dynamics and extent of the spatial AC of temperature as a benchmark to  
198 visually compare the dynamics of spatial AC in the other biological variables. This allowed us to  
199 tease apart the effect of physically- versus biologically-driven spatial patterns. Additionally, in  
200 order to better visualize the spatial patterns in chlorophyll *a*, phycocyanin, temperature,

201 dissolved oxygen, and pH over the course of the season, the data were interpolated using inverse  
202 distance weighting across a 25m grid (Figure 2).

203 In order to evaluate if high frequency measurements at a fixed station are an adequate  
204 characterization of the surface water dynamics in a shallow lake, we compared the measurements  
205 taken by the fixed station sonde during the same time period as a spatial sampling event. High  
206 frequency data from the fixed station sonde was trimmed to the period that we sampled the lake  
207 spatially. A t-test with a Bonferroni correction, to account for the multiple comparisons, was  
208 performed to compare the distribution from the 98 sampling stations to the fixed station  
209 measurements from the same day for each of the four biologically-mediated variables,  
210 chlorophyll, phycocyanin, dissolved oxygen, and pH. In addition to comparing fixed sonde  
211 values to the spatial sampling, we also used the spatial data to identify locations in the lake that  
212 were consistently representative of mean conditions, and therefore ideal locations for fixed  
213 station monitoring. We identified locations in the lake for each sampling event that had  
214 measurements within the range of  $\pm$  one standard deviation from the mean for each biologically  
215 mediated variable (all variables except temperature). We then collated these locations across all  
216 sampling dates to identify which of the 98 sampling locations had measurements that most  
217 consistently represented the mean conditions of the lake.

218 Finally, we performed a rarefaction analysis to evaluate the optimal spatial sampling  
219 frequency to capture the mean value of the biologically-mediated variables. This was done by  
220 randomly selecting  $n$  number of spatial sampling data points ( $n=2-97$ ) during a sampling event,  
221 calculating the mean value from that subset, and then calculating the root mean square error  
222 (RMSE), comparing the mean estimate from the subset to the mean of all sampling points during  
223 that event. This calculation was repeated 1000 times for each value of  $n$ , and each iteration was

224 then averaged. The averaged RMSE values for each subset of  $n$  were fit using a local polynomial  
225 regression with a smoothing factor of 0.1 and each sampling event's RMSE curve was  
226 standardized by subtracting the mean of all iterations ("global mean") from the mean at  $n$   
227 number of stations, to aid in visual comparison. The spatial data are available through (Ortiz &  
228 Wilkinson, 2019) and the fixed station data are available in (Ortiz et al., 2019) and further  
229 analyzed in Ortiz et al. (2020). All analyses were performed using R 4.0.2 (R Core Team, 2020)  
230 using the gstat (Pebesma, 2004), rstatix (Kassambara, 2020), and sf packages (Pebesma, 2018).

231

## 232 **Results**

233         There were two bloom events during the summer of 2018 in Swan Lake. The first bloom  
234 occurred from DOY 156 – 184 and was dominated by the diatom *Aulacoseira spp.* based on a  
235 sample taken on DOY 177 examined under a compound microscope at 400x magnification. The  
236 phycocyanin concentrations on DOY 177 were the lowest during this first bloom period (Figure  
237 2), and no cyanobacteria were identified in the sample. The second bloom, peaking on DOY 236,  
238 was dominated by the cyanobacterium *Microcystis spp.* There were also two large precipitation  
239 events during the summer, occurring after sampling on DOY 170 and lasting through DOY 171,  
240 and on DOY 232 (Figure 2; Supplemental Figure 1). The maximum wind speed recorded during  
241 the first precipitation event was 10.8 m s<sup>-1</sup> coming from the southwest and 11.8 m s<sup>-1</sup> during the  
242 second precipitation event coming from the southeast. During the first half of the summer (DOY  
243 142 – 191) the prevailing winds 24 hours prior to the sampling events were from the south,  
244 switched to being predominantly from the north from DOY 198 – 219, and then varied in  
245 direction for the rest of the season (Figure 2). The median wind speed for the first period when  
246 winds were out of the south was 3.6 m s<sup>-1</sup> (Figure 3b). When the winds switched to being

247 predominantly from the north between DOY 198 – 219, the median wind speed was lower at 2.5  
248 m s<sup>-1</sup> (Figure S1).

#### 249 *Spatial dynamics*

250 During the two bloom periods there was not a latitudinal or longitudinal trend in  
251 chlorophyll *a* concentrations; instead, there were patches of high chlorophyll *a* concentration on  
252 otherwise low-concentration dates (Figure 2). Unlike chlorophyll *a*, phycocyanin had a strong  
253 latitudinal trend with higher concentrations in the northern portion (sample sites A1-G4) of the  
254 lake during the first bloom. This spatial pattern is readily observed on DOY 184 but is also  
255 noticeable for many of the sampling events during the first bloom (Figure 2). During sampling  
256 events with a strong latitudinal gradient in phycocyanin (DOY 166 – 184, and 236) the mean  
257 concentration in the northern portion of the lake was nearly double the concentration in the  
258 southern portion of the lake (7.29 and 3.76 µg L<sup>-1</sup>, respectively). On these dates, the prevailing  
259 winds 24 hours prior to the sampling event were out of the north (Supplemental Figure S1), yet  
260 the lowest concentrations of phycocyanin were found in the southern portion of the lake. Even  
261 when the lake was not blooming, there were patches of high concentrations of phycocyanin in  
262 the northern portion of the lake (e.g., DOY 212), located among the densest patch of American  
263 Lotus (Figure 1). The average phycocyanin concentrations at the sampling locations within the  
264 American Lotus patch was higher than the average concentration in the rest of the lake for 14 of  
265 the 16 sampling events (Figure S2).

266 The daytime saturation of dissolved oxygen varied the most out of the five variables  
267 monitored, ranging from borderline hypoxic (30% saturation) to supersaturated (up to 350%)  
268 (Figure 2). While the dissolved oxygen saturation increased near the peak of the bloom, the  
269 highest average saturation was on DOY 191, after the first bloom had collapsed. There was a

270 weak pattern over the course of the season of higher saturation in the northern portion of the  
271 lake, similar to the distribution of higher phycocyanin concentrations. However, within the  
272 northern portion of the lake, regions of low dissolved oxygen saturation formed in the surface  
273 waters, particularly later in the summer (Figure 1). Beginning on DOY 198, the mean dissolved  
274 oxygen concentration in the American lotus patch was consistently lower than the average for the  
275 rest of the lake until DOY 250 (Figure S2). The distribution of pH also had a weak spatial pattern  
276 during the summer, with slightly elevated values in the northern portion of the lake during the  
277 first bloom (e.g. DOY 177; Figure 2). While pH was elevated at the onset of the first bloom  
278 period from DOY 149 – 170, it was highest overall on DOY 191 and 198 after the first bloom  
279 had collapsed. Unlike the other variables, temperature had a subtle south to north latitudinal  
280 gradient with warmer temperatures in the southern portion of the lake and colder in the north  
281 during the latter half of the summer (Figure 2). On average this difference between the northern  
282 portion of the lake and the southern was 0.5°C. The warmest day of sampling was DOY 191.

283         Spatial variability in algal pigments during the first bloom event was low, with two  
284 exceptions. There was an increase in the CV of chlorophyll *a* on the last day of the bloom (DOY  
285 184; Figure 3a) that continued to increase as the bloom collapsed. There was also a temporary  
286 increase in phycocyanin CV during the first bloom on DOY 177 (Figure 3b), coinciding with a  
287 temporary decline in phycocyanin concentration across the lake. The CV of both algal pigments  
288 was higher than the CV of temperature over the course of the entire sampling period.

289         Conversely, the CV of pH and dissolved oxygen were elevated during the first bloom  
290 period, with pH CV declining and remaining low after the first bloom (Figure 3c) and dissolved  
291 oxygen CV only temporarily declining after the first bloom (Figure 3d). Temperature had low  
292 variability throughout the first bloom until DOY 177, when the lake began to heat up, peaking in

293 both temperature and spatial variability on DOY 191 (Figure 3e). Between the first and second  
294 bloom, DOY 191-226, there was a decrease in spatial variability among the algal pigments and  
295 pH as the bloom collapsed, while temperature and dissolved oxygen CV remained relatively high  
296 and variable. During the second bloom period, CV was low for all variables except for  
297 chlorophyll *a*. In general, the CV, of temperature and pH, expressed as a percentage, was an  
298 order of magnitude lower than the other variables.

299         Spatial autocorrelation (AC), quantified as local Moran's I, did not fall substantially  
300 below 0 for any of the variables and peaked at 0.79 among all variables (Figure 3). The highest  
301 AC value for chlorophyll *a* and phycocyanin was during the first bloom event (Figure 3f, g);  
302 however, phycocyanin AC also increased substantially during the second bloom. During the first  
303 bloom, the AC of temperature varied similarly to both pigments' AC, particularly phycocyanin,  
304 but became decoupled after the bloom collapsed. While the AC of temperature remained high  
305 during the inter-bloom period, the AC of the pigments was substantially lower. Conversely, the  
306 dynamics of AC of temperature, dissolved oxygen and pH remained coupled throughout the  
307 summer (Figure 3h, i). Dissolved oxygen saturation and pH both increased in AC during the first  
308 bloom and then declined throughout the rest of the season with the exception of a minor increase  
309 in AC during the second bloom event.

310

### 311 *Fixed station versus spatial sampling*

312         There were a greater number of days with a significant difference between the spatial and  
313 fixed station measurements than days in which the data sets were not significantly different  
314 (Figure 4). Among all 64 comparisons (4 variables  $\times$  16 sampling events), the spatial and fixed  
315 station data sets had a means that were not significantly different 37.5% of the time.

316 Phycocyanin had the greatest number of events with similar values, with 7 of the 16 sampling  
317 events having non-statistically different mean values measured spatially and at the fixed station  
318 (Figure 4b). These occurrences were mainly during non-bloom periods. However, even when the  
319 mean phycocyanin values were similar between the sampling methods on a given day, the range  
320 of values captured by the fixed station was five times less than the variability captured in the  
321 spatial data. This pattern of infrequent occurrences of similar mean values between the two  
322 methods during non-bloom periods and a diminished range in the fixed station data, was shared  
323 to a degree, among the other three variables as well. Interestingly, dissolved oxygen saturation  
324 only had 5 out of the 16 events with means that were not significantly different, all of which  
325 occurred when the lake was above 100% saturation (Figure 4c).

326         While a majority of the comparisons between the fixed station and spatial data indicate  
327 that the algal pigments had a larger range of values in the spatial data, there were a handful of  
328 instances where the opposite was true. During the first bloom, the fixed station sonde measured a  
329 wide range of chlorophyll *a* concentrations and had a higher mean chlorophyll *a* for all dates  
330 (Figure 4a). Similarly, we observed higher mean phycocyanin at the fixed station sonde on DOY  
331 156, 166, 177, 191, and 219 (Figure 4b). However, this pattern did not hold true for dissolved  
332 oxygen or pH (Figure 5c, d).

333         The spatial sampling sites that most consistently captured the mean values in the lake on  
334 a given sampling date were in the northwest portion of the lake, near the inlet. The best  
335 performing site for all variables was site E3, just west of the American lotus patch and adjacent  
336 to a bed of sago pondweed (Figure 1). The four biologically-mediated variables from sample site  
337 E3 were within the mean ( $\pm$  standard deviation) range of all of the spatial measurements 95% of  
338 the time. The second best performing location was in the middle of the American lotus patch, site

339 D4, with the values from this site being within the mean ( $\pm$  standard deviation) range 92% of the  
340 time. The site where the fixed station was located, site H2, was only within the mean ( $\pm$  standard  
341 deviation) range 58% of the time.

342

### 343 *Optimal Spatial Resolution*

344 In order to evaluate the spatial sampling resolution needed to capture the mean state of  
345 the surface water on a given day, we performed a rarefaction analysis for each variable and each  
346 sampling event, calculating the root mean squared error (RMSE) of a subset of sampling  
347 locations compared to the mean value of all 98 measurements that day. The plateaus of the  
348 RMSE curves from the rarefaction analysis were used to evaluate the smallest number of spatial  
349 sampling locations needed to capture the mean across the lake during that sampling event (Figure  
350 5). Additionally, we also evaluated the temporal pattern of the minimum number of sampling  
351 locations needed to capture the mean.

352 Mean values were underestimated for all variables on all sampling dates when there were  
353 less than 10 sampling stations (Figure 5). However, the severity of the underestimation differed  
354 among the variables. The rarefaction analysis for chlorophyll *a* indicated that 10 – 30 sampling  
355 locations was sufficient for capturing the mean chlorophyll *a* in Swan Lake, otherwise the mean  
356 concentration would be under estimated (Figure 5a). When an algal bloom was occurring it took  
357 more sampling locations to near the mean chlorophyll *a* concentration on that date. However,  
358 when the bloom was particularly patchy during development (DOY 226) or collapse (DOY 191),  
359 including a larger number of sampling locations led to overestimating the mean chlorophyll *a*  
360 concentration as locations with high concentrations were over-represented in the data set. There  
361 were similar patterns in phycocyanin RMSE with most sampling dates plateauing between 20 –



362 30 sampling locations with a few exceptions (Figure 5b). For DOYs 156-170 (rise of the first  
363 bloom) and 212, at least 60 sampling locations were needed to capture the overall mean in  
364 phycocyanin for that sampling date. Dissolved oxygen saturation and pH were generally well  
365 characterized by approximately 10 – 15 sampling locations as both had a majority of dates in  
366 which the RMSE curves plateaued at that spatial sampling resolution (Figure 5c, d). However, at  
367 the beginning (DOY 154), peak (DOY 184), and end (DOY 205) of the first bloom, twice as  
368 many sampling locations were needed to capture the mean dissolved oxygen. Only two dates  
369 required more sampling locations for pH to capture the mean, DOY 177 and 198, which  
370 plateaued at approximately 40 sampling locations. The largest RMSE were observed during  
371 bloom conditions for all variables: DOY 177 had the largest error for phycocyanin and pH, while  
372 the largest RMSE was on DOY 184 for dissolved oxygen and on DOY 236 for chlorophyll *a*  
373 (Figure 5).

374

## 375 **Discussion**

376 The spatial heterogeneity of water quality parameters was highly dynamic in Swan Lake,  
377 a shallow, hypereutrophic, temperate waterbody. The temporal dynamics in heterogeneity were  
378 driven in part by the two blooms, the peaks of which were preceded by large precipitation events.  
379 These rain events could have delivered nutrients from the agriculturally dominated watershed  
380 into the lake from the northern inlet, helping to fuel the subsequent algal blooms and the spatial  
381 patterns observed during blooms (Stockwell et al. 2020). However, there are also a number of  
382 other factors that likely contributed to the spatial variability and pattern during and following  
383 these bloom events, including the prevailing wind direction prior to sampling, the bathymetry of

384 the basin and location of the surface inlet, and the potential for dense macrophyte beds to  
385 contribute to local patchiness.

386         The spatial patterns that the algal blooms created were consistent with the expectations  
387 from previous modeling and experimental work that spatial AC increases as algal blooms  
388 develop (Buelo et al., 2018; Butitta et al., 2017; Serizawa et al., 2008). This pattern was the  
389 strongest for phycocyanin, evident by the strong latitudinal gradient in concentrations during the  
390 bloom periods. The sampling dates with phycocyanin concentration gradients (e.g., DOY 166,  
391 177, 184, 236) coincided with persistent winds from the south 24 hours prior to the sampling  
392 event, which likely resulted in the higher concentration of algal cells in the northern portion of  
393 the lake. The effect of persistent wind directions influencing the distribution of cyanobacteria has  
394 also been documented in other shallow eutrophic lakes (Wu et al. 2010). The shallow sediments  
395 of the northern basin were also likely a source of akinete recruitment (Karlsson-Elfgren and  
396 Brunburg, 2004), further contributing to the higher concentrations of phycocyanin in the northern  
397 portion of the lake during the first bloom. Augmented nutrient availability in the northern part of  
398 the lake due to external loading from the watershed through the surface inlet and internal loading  
399 from the sediments overlain by an unstratified water column (Song and Burgin 2017) may have  
400 further amplified the phytoplankton gradient, particularly following precipitation events. Finally,  
401 the tendency of the dominant cyanobacteria taxa *Microcystis spp.* to form surface scums likely  
402 enhanced the spatial patterns observed with our surface sampling approach.

403         The sampling dates with a strong gradient of phytoplankton concentrations from north to  
404 south also resulted in north-south gradients in water chemistry. On these dates, both dissolved  
405 oxygen and pH formed a gradient of high values in the northern portion of the lake and lower  
406 values in the south, which would be expected with greater primary production where

407 phytoplankton concentrations were highest. The spatial patterns in the surface water chemistry  
408 demonstrate how phytoplankton spatial distribution, driven by wind, can create hot spots and  
409 moments of biogeochemical activity within lakes (McClain et al. 2003) that may be missed with  
410 traditional, single-station sampling. The dense patch of floating leaf American lotus macrophytes  
411 also created a hot spot of biogeochemical activity.

412         Macrophyte beds can have a large local influence on water chemistry by inducing  
413 stratification, decreasing flow and trapping particles, and modifying the light environment  
414 (Green, 2006; Vilas et al., 2017). For 87.5% of the season the phycocyanin concentrations were  
415 higher in the bed of American lotus than concentrations elsewhere in the lake. In fact, even on  
416 sampling dates when phycocyanin concentrations were otherwise low (e.g., DOY 212), the  
417 American lotus patch can be identified based on the phycocyanin concentrations that are nearly  
418 twice as high as the rest of the lake. We hypothesize that the macrophyte patch allowed for  
419 microstratification in the water column and reduced wind-driven flow. These physical conditions  
420 are likely to favor cyanobacteria dominance and the formation of surface scums. Similarly, the  
421 dissolved oxygen concentrations in the American lotus patch became consistently lower than the  
422 rest of the lake later in the summer, likely due to the plants beginning to senesce, creating a hot  
423 spot of decomposition, decreasing both dissolved oxygen and pH (Vilas et al., 2017). While there  
424 isn't strong evidence in the data that the other submerged macrophyte beds had a similarly strong  
425 effect on water chemistry, the data from the American lotus patch illustrates how macrophytes  
426 can contribute to local patchiness and overall spatial heterogeneity.

427

428 *Considerations for Monitoring*

429           The variables that we measured in this study are often the target of water quality  
430 monitoring as the dynamics of these variables coincide with changes in ecosystem function and  
431 services. Monitoring is often performed at a fixed station over time to capture the dynamics of  
432 the ecosystem, but this strategy could potentially result in missed information about the  
433 ecosystem's behavior. While the temporal dynamics of all the variables were synchronous  
434 between the fixed station and spatial sampling data sets in Swan Lake, our conclusions regarding  
435 the magnitude of the blooms and variability in the lake's structure would have been substantially  
436 different relying solely on the fixed station data. Among the four biologically-mediated  
437 variables, only 37.5% of the fixed station estimates of the mean state of the lake statistically  
438 matched the estimate from the spatial sampling. The vast majority of those instances (96%)  
439 occurred during non-bloom periods, which also coincided with lower wind speed conditions, no  
440 prevailing wind direction, and no major precipitation events. The large difference between the  
441 spatial sampling and fixed station measurements of algal pigments during blooms was likely  
442 driven, in part, by the depth of the sensors at the fixed station and the variable accumulation of  
443 cyanobacteria at the surface of the lake dependent upon environmental conditions and the  
444 dominant taxa (Chaffin et al., 2020). It is clear from our data that during periods of heightened  
445 biological activity such as blooms, fixed station monitoring is unlikely to be representative of the  
446 mean ecosystem state in shallow lakes.

447           Despite the high degree of horizontal spatial variability that has been documented in this  
448 study and others (Loken et al. 2019, Van de Bogert et al. 2012, Buttita et al. 2017), fixed station  
449 designs are widely used in water quality monitoring programs. In Swan Lake, we determined that  
450 the historical location for water quality monitoring, where the fixed station sensors were  
451 deployed, was one of the least-representative locations for mean conditions in the lake. Given the

452 hypereutrophic state of the lake, the most immediate management concerns are toxic  
453 cyanobacteria blooms and summer fish kills due to low dissolved oxygen. Yet, the mean value of  
454 these variables (phycocyanin and dissolved oxygen) across the lake were only captured by the  
455 fixed station sensors 58% of the time. While selecting a fixed station site for high frequency  
456 sensor deployment includes many considerations including the location of previous data  
457 collection and management needs, based on our analysis we would advise performing a spatial  
458 survey to identify if and when the fixed station site is representative of mean conditions in the  
459 lake. A complementary spatial survey will help contextualize the fixed station dynamics and  
460 provide additional, management-relevant information about the lake.

461         It's also important to consider the trade-offs between high frequency fixed station  
462 monitoring and higher resolution, but less frequent spatial monitoring. High frequency  
463 monitoring at a single station provides insight into ecosystem function such as metabolism  
464 (Staeher et al., 2012), early warnings of impending regime shifts (Carpenter et al., 2011;  
465 Wilkinson et al., 2018), and crucial information on diel variability in limnological conditions  
466 (Andersen et al., 2017). However, as we observed in Swan Lake, the spatial variability within a  
467 given day often exceeds the temporal variability at a single point in a shallow lake. Without the  
468 spatial sampling snapshots, we would have underestimated the magnitude of the algal blooms,  
469 hampering our limnological understanding of the ecosystem's functioning and impeding our  
470 ability to accurately estimate rates such as methane emissions on a global scale (DelSontro et al.  
471 2018).

472         From a practical stand point, the understanding gleaned from the spatial sampling could  
473 help managers design targeted algal toxin monitoring or management interventions to help  
474 control fish habitat quality in persistently hypoxic areas (Bardshaw et al., 2015). However, the

475 time and cost investment in repeated spatial sampling at the resolution performed in this study  
476 may not be feasible for both research and management programs. The rarefaction analysis we  
477 performed for all four of the key water quality monitoring variables revealed that minimal  
478 information was gained after ~30 locations were sampled across many conditions and variables.  
479 Often 12-20 sample locations across the 40.5 ha lake (or a 1-2 samples per hectare) was  
480 sufficient to capture the spatial variability within the lake, with a few exceptions. These  
481 exceptions occurred during times of higher variability such as when the blooms were just starting  
482 or when the bloom began to collapse. The need for a higher spatial resolution during bloom  
483 events to fully capture their variability has also been found using remote sensing techniques in  
484 other, larger lakes (Lekki et al., 2019). As the spatial resolution of remote sensing technologies  
485 continues to improve, it may become more cost effective to capture the spatial heterogeneity of  
486 algal pigments in small lakes over time. However, one of the benefits of manual spatial sampling  
487 is being able to pair other measurements such as dissolved oxygen, pH, and nutrients (e.g.,  
488 nitrate; Loken et al., 2018; Pellerin et al., 2016) with information on the distribution of algal  
489 biomass.

490         Our intensive spatial monitoring of a shallow, hypereutrophic lake revealed how spatially  
491 heterogeneous shallow lakes are over the course of a single season and allowed us to tease apart  
492 the drivers of that spatial heterogeneity. We found that variability was greatest during  
493 biologically-intensive periods, such as during algal blooms and in dense floating-leaf macrophyte  
494 beds, and that failure to capture this variability would have hampered our understanding of the  
495 ecosystem's functioning and overall mean state. Small lakes such as Swan Lake dominate the  
496 global distribution of waterbodies (Verpoorter et al. 2014). Adequately capturing and  
497 characterizing the magnitude of variability in production of these waterbodies is important given

498 their role in mediating global nutrient cycles (Downing et al. 2010, Biddanda et al. 2017),  
499 especially methane emissions (DelSontro et al. 2018, Loken et al. 2019). Our data provided an  
500 estimate of the spatial resolution needed to capture the dynamics in ecosystems similar to Swan  
501 Lake and a method which could be readily adapted to other ecosystems. While our results  
502 provide new understanding of the magnitude and temporal dynamics of spatial heterogeneity in  
503 shallow lakes, continued investigation of horizontal spatial heterogeneity in a range of aquatic  
504 ecosystems, from oligotrophic to eutrophic, is needed to better understand the structure and  
505 drivers of horizontal spatial variability in lakes.

506

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514

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720

721 **Figures Captions**

722 **Figure 1.** Sampling locations on a 65 m square grid of Swan Lake, a 40.5 hectare waterbody in  
723 western Iowa, USA. The main inlet to the lake and only outlet indicated with arrows. a) The  
724 location of the macrophyte beds of the two dominant species within the lake are shown on the  
725 map, with darker shading indicating the regions with the densest vegetation and the location of  
726 the high frequency sensor, b) the bathymetry of the lake and location of the lake in the state of  
727 Iowa, in reference to the United States of America.

728

729 **Figure 2.** The spatial pattern of each of the variables chlorophyll *a* (Chl,  $\mu\text{g L}^{-1}$ ), phycocyanin  
730 (Phyco,  $\mu\text{g L}^{-1}$ ), dissolved oxygen (DO, percent saturation), pH, and temperature (Temp,  $^{\circ}\text{C}$ ) for  
731 each sampling event. The 98 sampling locations taken in a 65 m grid (Figure 1) were interpolated  
732 to a 25m grid using spatial inverse distance interpolation for visualization here. The color ramps  
733 for each variable are scaled from the lowest to the highest value observed over the course of the  
734 season across all sampling locations. The wind roses are the wind speeds ( $\text{m s}^{-1}$ ; color ramp) and  
735 direction the wind came from for the 24 hours prior to a sampling event. The concentric circles  
736 are the frequency of winds from that direction for the 24 hour period (expressed as a percentage,  
737 largest circle is 80% of the time). In the case of a longer “spoke”, the greater amount of time the  
738 wind was from that direction. The horizontal lines between DOY 170 and 177, and DOY 226  
739 and 236 mark the two large precipitation events that occurred between those sampling dates.

740

741 **Figure 3.** Time series of the spatial coefficient of variation (CV) and spatial autocorrelation (AC;  
742 local Moran’s I) of the biologically-mediated variables in Swan Lake (same variable

743 abbreviations as Figure 2). The gray polygons indicate periods of algal bloom. The red line is the  
744 time series of temperature local Moran's I for comparison.

745  
746 **Figure 4.** Comparison of the mean (lines and points) and range (shaded polygon) of  
747 measurements from the spatial sampling and fixed station measurements. The fixed station data  
748 were trimmed to the period that spatial sampling occurred. A filled circle is used for the  
749 sampling dates when the means from the two sampling approaches were significantly different  
750 ( $p < 0.05$ ), and an open triangle is used for the sampling dates when the mean of the two  
751 approaches were not significantly different. The dark blue vertical lines indicate the dates of the  
752 two major precipitation events and the red dashed line in panel c is at 100% dissolved oxygen  
753 saturation.

754  
755 **Figure 5.** Standardized root mean squared errors (RMSE) of rarefaction analysis. Fit lines  
756 represent each sampling dates standardized RMSE (16 in total) and the gradient from light to  
757 dark indicates first sampling event to last.