

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

2 **Authors:** David Ortiz^{1,2*}, Grace Wilkinson^{1,2}

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

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

7 *Corresponding Author: dortiz4@wisc.edu

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16 **Abstract**

- 17 1. Algal blooms can have profound effects on the structure and function of aquatic
18 ecosystems and have the potential to interrupt valuable ecosystem services. Despite the
19 potential ecological and economic consequences of algal blooms, the spatial dynamics of
20 bloom development in spatially complex ecosystems such as shallow lakes remain poorly
21 characterized. Our goal was to evaluate the magnitude and drivers of spatial variability of
22 algal biomass, dissolved oxygen and pH over the course of a season, in a shallow lake in
23 order to better understand the spatial dynamics of algal blooms in these ecosystems.
- 24 2. We sampled 98 locations in a small eutrophic lake on a 65m grid for several parameters
25 (chlorophyll *a*, phycocyanin, dissolved oxygen, pH, and temperature), weekly over 122
26 days. This was done to estimate the dynamics of variability and spatial autocorrelation
27 during the course of multiple bloom events. We also compared the spatial measurements
28 to a high frequency sensor deployed at a fixed station and estimated the optimal spatial
29 sampling resolution by performing a rarefaction analysis.
- 30 3. Spatial heterogeneity of algal pigments was high, particularly during bloom events, and
31 this pattern and the overall severity of the bloom was not well captured with the fixed
32 station monitoring. The pattern of algal pigments and other limnologically important
33 variables (dissolved oxygen and pH) was related to the direction of prevailing winds 24
34 hours prior to sampling, the shallow northern basin where the main surface inlet is
35 located, and heavy precipitation. Additionally, a dense bed of floating-leaf macrophytes
36 contributed to local patchiness in all variables. Finally, from the rarefaction analysis we
37 found that minimal information about the mean state of the ecosystem was gained after
38 ~30 locations had been sampled.

39 4. This study revealed how spatially heterogeneous shallow lakes are over the course of a
40 single season, and that the magnitude of variability was highest during biologically-
41 intensive periods such as algal blooms. As such, continued research is needed across a
42 range of trophic conditions to better understand the structure of horizontal variability in
43 lakes. Overall, these data demonstrate the need for spatially-explicit monitoring to better
44 understand the dynamics and drivers of algal blooms in shallow lakes and to better
45 manage ecosystem services.

46 **Introduction**

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

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

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

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

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

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

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

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

123

124 **Methods**

125 *Study Site*

126 Swan Lake (42.0396, -94.8454) has an average depth of 2 m, surface area of 40.5
127 hectares, and a shoreline development index value of 1.54 (more irregular shape as compared to
128 a perfect circle with the same surface area). The watershed is 350 hectares with 92% of the land
129 in agricultural use. The estimated water residence time is approximately 1.5 years. During the
130 ice-free period of 2018, Swan Lake had an average total phosphorus concentration of $280 \mu\text{g L}^{-1}$
131 and a total nitrogen concentration of 1.61 mg L^{-1} , making it hypereutrophic (Carlson, 1977).
132 Total nitrogen was measured as the sum of total Kjeldahl nitrogen (method 351.2 v2, US EPA,
133 1993c) and nitrate + nitrite measured using the cadmium reduction method (method 4500-NO₃-
134 F, US EPA, 1993a). Total phosphorus was measured using the ascorbic acid method (method
135 365.1 v2, US EPA, 1993b). The average total alkalinity during the same period was 139 mg
136 $\text{CaCO}_3 \text{ L}^{-1}$ determined through end point titration (APHA, 1998). In addition to seasonal algal
137 blooms, Swan Lake also has non-continuous beds of American lotus (*Nelumbo lutea*) and sago

138 pondweed (*Stuckenia pectinata*) that peak in biomass in the latter half of the summer and then
139 begin senescing. The main surface inlet to the lake enters on the western side and the outlet is at
140 the southern edge of the waterbody (Figure 1). There are no known springs feeding the lake.

141

142 *Field Methods*

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

161 established, permanent macrophyte beds in the lake for that summer. These weekly
162 presence/absence data were used to construct the macrophyte distributions in Figure 1.

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

175

176 *Data analysis*

177 Spatial heterogeneity can be quantified by calculating the spatial variance (e.g.,
178 coefficient of variation; CV) or spatial autocorrelation (Moran's I, Moran, 1950). Increasing
179 spatial variance is indicative of increasing patchiness in the ecosystem, such as areas of high-
180 density algal biomass and areas of low-density biomass within a lake. Spatial autocorrelation
181 accounts for the location of those patches within the ecosystem in relationship to each other.
182 Local Moran's I quantifies how similar the abundance of algae is at one location compared to the
183 density of surrounding neighbors. When measured over time for variables that are indices of

184 algal biomass (e.g., the pigments chlorophyll *a* and phycocyanin), both of these metrics of spatial
185 heterogeneity can provide insight into the dynamics of algal bloom development. In models of
186 algal blooms, both spatial variance and autocorrelation are expected to be high during the bloom
187 period (Buelo et al., 2018).

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

207 order to better visualize the spatial patterns in chlorophyll *a*, phycocyanin, temperature,
208 dissolved oxygen, and pH over the course of the season, the data were interpolated using inverse
209 distance weighting across a 25m grid (Figure 2).

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

225 Finally, we performed a rarefaction analysis to evaluate the optimal spatial sampling
226 frequency to capture the mean value of the biologically-mediated variables. This was done by
227 randomly selecting *n* number of spatial sampling data points ($n=2-97$) during a sampling event,
228 calculating the mean value from that subset, and then calculating the root mean square error
229 (RMSE), comparing the mean estimate from the subset to the mean of all sampling points during

230 that event. This calculation was repeated 1000 times for each value of n , and each iteration was
231 then averaged. The averaged RMSE values for each subset of n were fit using a local polynomial
232 regression with a smoothing factor of 0.1 and each sampling event's RMSE curve was
233 standardized by subtracting the mean of all iterations ("global mean") from the mean at n
234 number of stations, to aid in visual comparison. The spatial data are available through (Ortiz &
235 Wilkinson, 2019) and the fixed station data are available in (Ortiz et al., 2019) and further
236 analyzed in Ortiz et al. (2020). All analyses were performed using R 4.0.2 (R Core Team, 2020)
237 using the gstat (Pebesma, 2004), rstatix (Kassambara, 2020), and sf packages (Pebesma, 2018).

238

239 **Results**

240 There were two bloom events during the summer of 2018 in Swan Lake. The first bloom
241 occurred from DOY 156 – 184 and was dominated by the diatom *Aulacoseira spp.* based on a
242 sample taken on DOY 177 examined under a compound microscope at 400x magnification. The
243 phycocyanin concentrations on DOY 177 were the lowest during this first bloom period (Figure
244 2), and no cyanobacteria were identified in the sample. The second bloom, peaking on DOY 236,
245 was dominated by the cyanobacterium *Microcystis spp.* There were also two large precipitation
246 events during the summer, occurring after sampling on DOY 170 and lasting through DOY 171,
247 and on DOY 232 (Figure 2; Supplemental Figure 1). The maximum wind speed recorded during
248 the first precipitation event was 10.8 m s^{-1} coming from the southwest and 11.8 m s^{-1} during the
249 second precipitation event coming from the southeast. During the first half of the summer (DOY
250 142 – 191) the prevailing winds 24 hours prior to the sampling events were from the south,
251 switched to being predominantly from the north from DOY 198 – 219, and then varied in
252 direction for the rest of the season (Figure 2). The median wind speed for the first period when

253 winds were out of the south was 3.6 m s^{-1} (Figure 3b). When the winds switched to being
254 predominantly from the north between DOY 198 – 219, the median wind speed was lower at 2.5
255 m s^{-1} (Figure S1).

256 *Spatial dynamics*

257 During the two bloom periods there was not a latitudinal or longitudinal trend in
258 chlorophyll *a* concentrations; instead, there were patches of high chlorophyll *a* concentration on
259 otherwise low-concentration dates (Figure 2). Unlike chlorophyll *a*, phycocyanin had a strong
260 latitudinal trend with higher concentrations in the northern portion (sample sites A1-G4) of the
261 lake during the first bloom. This spatial pattern is readily observed on DOY 184 but is also
262 noticeable for many of the sampling events during the first bloom (Figure 2). During sampling
263 events with a strong latitudinal gradient in phycocyanin (DOY 166 – 184, and 236) the mean
264 concentration in the northern portion of the lake was nearly double the concentration in the
265 southern portion of the lake (7.29 and $3.76 \mu\text{g L}^{-1}$, respectively). On these dates, the prevailing
266 winds 24 hours prior to the sampling event were out of the north (Supplemental Figure S1), yet
267 the lowest concentrations of phycocyanin were found in the southern portion of the lake. Even
268 when the lake was not blooming, there were patches of high concentrations of phycocyanin in
269 the northern portion of the lake (e.g., DOY 212), located among the densest, permanent patch of
270 American Lotus (Figure 1, Figure S3). The average phycocyanin concentrations at the sampling
271 locations within the American Lotus patch was higher than the average concentration in the rest
272 of the lake for 14 of the 16 sampling events (Figure S2).

273 The daytime saturation of dissolved oxygen varied the most out of the five variables
274 monitored, ranging from borderline hypoxic (30% saturation) to supersaturated (up to 350%)
275 (Figure 2). While the dissolved oxygen saturation increased near the peak of the bloom, the

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

290 Spatial variability in algal pigments during the first bloom event was low, with two
291 exceptions. There was an increase in the CV of chlorophyll *a* on the last day of the bloom (DOY
292 184; Figure 3a) that continued to increase as the bloom collapsed. There was also a temporary
293 increase in phycocyanin CV during the first bloom on DOY 177 (Figure 3b), coinciding with a
294 temporary decline in phycocyanin concentration across the lake. The CV of both algal pigments
295 was higher than the CV of temperature over the course of the entire sampling period.

296 Conversely, the CV of pH and dissolved oxygen were elevated during the first bloom
297 period, with pH CV declining and remaining low after the first bloom (Figure 3c) and dissolved
298 oxygen CV only temporarily declining after the first bloom (Figure 3d). Temperature had low

309 variability throughout the first bloom until DOY 177, when the lake began to heat up, peaking in
300 both temperature and spatial variability on DOY 191 (Figure 3e). Between the first and second
301 bloom, DOY 191-226, there was a decrease in spatial variability among the algal pigments and
302 pH as the bloom collapsed, while temperature and dissolved oxygen CV remained relatively high
303 and variable. During the second bloom period, CV was low for all variables except for
304 chlorophyll *a*. In general, the CV, of temperature and pH, expressed as a percentage, was an
305 order of magnitude lower than the other variables.

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

317

318 *Fixed station versus spatial sampling*

319 There were a greater number of days with a significant difference between the spatial and
320 fixed station measurements than days in which the data sets were not significantly different
321 (Figure 4). Among all 64 comparisons (4 variables \times 16 sampling events), the spatial and fixed

322 station data sets had a means that were not significantly different 37.5% of the time. However,
323 the direction of change from week to week was generally consistent between the spatial and
324 fixed station data sets. Phycocyanin had the greatest number of events with similar values, with 7
325 of the 16 sampling events having non-statistically different mean values measured spatially and
326 at the fixed station (Figure 4b). These occurrences were mainly during non-bloom periods.
327 However, even when the mean phycocyanin values were similar between the sampling methods
328 on a given day, the range of values captured by the fixed station was five times less than the
329 variability captured in the spatial data. This pattern of infrequent occurrences of similar mean
330 values between the two methods during non-bloom periods and a diminished range in the fixed
331 station data, was shared to a degree, among the other three variables as well. Interestingly,
332 dissolved oxygen saturation only had 5 out of the 16 events with means that were not
333 significantly different, all of which occurred when the lake was above 100% saturation (Figure
334 4c).

335 While a majority of the comparisons between the fixed station and spatial data indicate
336 that the algal pigments had a larger range of values in the spatial data, there were a handful of
337 instances where the opposite was true. During the first bloom, the fixed station sonde measured a
338 wide range of chlorophyll *a* concentrations and had a higher mean chlorophyll *a* for all dates
339 (Figure 4a). Similarly, we observed higher mean phycocyanin at the fixed station sonde on DOY
340 156, 166, 177, 191, and 219 (Figure 4b). However, this pattern did not hold true for dissolved
341 oxygen or pH (Figure 5c, d).

342 The spatial sampling sites that most consistently captured the mean values in the lake on
343 a given sampling date were in the northwest portion of the lake, near the inlet. The best
344 performing site for all variables was site E3, just west of the American lotus patch and adjacent

345 to a bed of sago pondweed (Figure 1). The four biologically-mediated variables from sample site
346 E3 were within the mean (\pm standard deviation) range of all of the spatial measurements 95% of
347 the time. The second best performing location was in the middle of the American lotus patch, site
348 D4, with the values from this site being within the mean (\pm standard deviation) range 92% of the
349 time. The site where the fixed station was located, site H2, was only within the mean (\pm standard
350 deviation) range 58% of the time.

351

352 *Optimal Spatial Resolution*

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

361 Mean values were underestimated for all variables on all sampling dates when there were
362 less than 10 sampling stations (Figure 5). However, the severity of the underestimation differed
363 among the variables. The rarefaction analysis for chlorophyll *a* indicated that 10 – 30 sampling
364 locations was sufficient for capturing the mean chlorophyll *a* in Swan Lake, otherwise the mean
365 concentration would be under estimated (Figure 5a). When an algal bloom was occurring it took
366 more sampling locations to near the mean chlorophyll *a* concentration on that date. However,
367 when the bloom was particularly patchy during development (DOY 226) or collapse (DOY 191),

368 including a larger number of sampling locations led to overestimating the mean chlorophyll *a*
369 concentration as locations with high concentrations were over-represented in the data set. There
370 were similar patterns in phycocyanin RMSE with most sampling dates plateauing between 20 –
371 30 sampling locations with a few exceptions (Figure 5b). For DOYs 156-170 (rise of the first
372 bloom) and 212, at least 60 sampling locations were needed to capture the overall mean in
373 phycocyanin for that sampling date. Dissolved oxygen saturation and pH were generally well
374 characterized by approximately 10 – 15 sampling locations as both had a majority of dates in
375 which the RMSE curves plateaued at that spatial sampling resolution (Figure 5c, d). However, at
376 the beginning (DOY 154), peak (DOY 184), and end (DOY 205) of the first bloom, twice as
377 many sampling locations were needed to capture the mean dissolved oxygen. Only two dates
378 required more sampling locations for pH to capture the mean, DOY 177 and 198, which
379 plateaued at approximately 40 sampling locations. The largest RMSE were observed during
380 bloom conditions for all variables: DOY 177 had the largest error for phycocyanin and pH, while
381 the largest RMSE was on DOY 184 for dissolved oxygen and on DOY 236 for chlorophyll *a*
382 (Figure 5).

383

384 **Discussion**

385 The spatial heterogeneity of water quality parameters was highly dynamic in Swan Lake,
386 a shallow, hypereutrophic, temperate waterbody. The temporal dynamics in heterogeneity were
387 driven in part by the two blooms, the peaks of which were preceded by large precipitation events.
388 These rain events could have delivered nutrients from the agriculturally dominated watershed
389 into the lake from the northern inlet, helping to fuel the subsequent algal blooms and the spatial
390 patterns observed during blooms (Stockwell et al. 2020). However, there are also a number of

391 other factors that likely contributed to the spatial variability and pattern during and following
392 these bloom events, including the prevailing wind direction prior to sampling, the bathymetry of
393 the basin and location of the surface inlet, and the potential for macrophyte beds to contribute to
394 local patchiness.

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

412 The sampling dates with a strong gradient of phytoplankton concentrations from north to
413 south also resulted in north-south gradients in water chemistry. On these dates, both dissolved

414 oxygen and pH formed a gradient of high values in the northern portion of the lake and lower
415 values in the south, which would be expected with greater primary production where
416 phytoplankton concentrations were highest. The spatial patterns in the surface water chemistry
417 demonstrate how phytoplankton spatial distribution, driven by wind, can create hot spots and
418 moments of biogeochemical activity within lakes (McClain et al. 2003) that may be missed with
419 traditional, single-station sampling. The dense, permanent patch of floating leaf American lotus
420 macrophytes also created a hot spot of biogeochemical activity.

421 Macrophyte beds can have a large local influence on water chemistry by inducing
422 stratification, decreasing flow and trapping particles, and modifying the light environment
423 (Green, 2006; Vilas et al., 2017). For 14 of the 16 weeks (87.5%) of the season the phycocyanin
424 concentrations were higher in the bed of American lotus than concentrations elsewhere in the
425 lake. In fact, even on sampling dates when phycocyanin concentrations were otherwise low (e.g.,
426 DOY 212), the American lotus patch can be identified based on the phycocyanin concentrations
427 that are nearly twice as high as the rest of the lake. We hypothesize that the macrophyte patch
428 allowed for microstratification in the water column and reduced wind-driven flow. These
429 physical conditions are likely to favor cyanobacteria dominance and the formation of surface
430 scums. Similarly, the dissolved oxygen concentrations in the American lotus patch became
431 consistently lower than the rest of the lake later in the summer, likely due to the plants beginning
432 to senesce, creating a hot spot of decomposition, decreasing both dissolved oxygen and pH
433 (Vilas et al., 2017). While there is not strong evidence in the data that the other submerged
434 macrophyte beds had a similarly strong effect on water chemistry, the data from the American
435 lotus patch illustrates how macrophytes can contribute to local patchiness and overall spatial
436 heterogeneity.

437

438 *Considerations for Monitoring*

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

457 Despite the high degree of horizontal spatial variability that has been documented in this
458 study and others (Loken et al. 2019, Van de Bogert et al. 2012, Buttita et al. 2017), fixed station
459 designs are widely used in water quality monitoring programs. In Swan Lake, we determined that

460 the historical location for water quality monitoring, where the fixed station sensors were
461 deployed, was one of the least-representative locations for mean conditions in the lake. Given the
462 hypereutrophic state of the lake, the most immediate management concerns are toxic
463 cyanobacteria blooms and summer fish kills due to low dissolved oxygen. Yet, the mean value of
464 these variables (phycocyanin and dissolved oxygen) across the lake were only captured by the
465 fixed station sensors 58% of the time. While selecting a fixed station site for high frequency
466 sensor deployment includes many considerations including the location of previous data
467 collection and management needs, based on our analysis we would advise performing a spatial
468 survey to identify if and when the fixed station site is representative of mean conditions in the
469 lake. A complementary spatial survey will help contextualize the fixed station dynamics and
470 provide additional, management-relevant information about the lake.

471 It's also important to consider the trade-offs between high frequency fixed station
472 monitoring and higher resolution, but less frequent spatial monitoring. High frequency
473 monitoring at a single station provides insight into ecosystem function such as metabolism
474 (Staehr et al., 2012), early warnings of impending regime shifts (Carpenter et al., 2011;
475 Wilkinson et al., 2018), and crucial information on diel variability in limnological conditions
476 (Andersen et al., 2017). However, as we observed in Swan Lake, the spatial variability within a
477 given day often exceeds the temporal variability at a single point in a shallow lake. Without the
478 spatial sampling snapshots, we would have underestimated the magnitude of the algal blooms,
479 hampering our limnological understanding of the ecosystem's functioning and impeding our
480 ability to accurately estimate rates such as methane emissions on a global scale (DelSontro et al.
481 2018).

482 From a practical stand point, the understanding gleaned from the spatial sampling could
483 help managers design targeted algal toxin monitoring or management interventions to help
484 control fish habitat quality in persistently hypoxic areas (Bardshaw et al., 2015). However, the
485 time and cost investment in repeated spatial sampling at the resolution performed in this study
486 may not be feasible for both research and management programs. The rarefaction analysis we
487 performed for all four of the key water quality monitoring variables revealed that minimal
488 information was gained after ~30 locations were sampled across many conditions and variables.
489 Often 12-20 sample locations across the 40.5 ha lake (or a 1-2 samples per hectare) was
490 sufficient to capture the spatial variability within the lake, with a few exceptions. These
491 exceptions occurred during times of higher variability such as when the blooms were just starting
492 or when the bloom began to collapse. The need for a higher spatial resolution during bloom
493 events to fully capture their variability has also been found using remote sensing techniques in
494 other, larger lakes (Lekki et al., 2019). As the spatial resolution of remote sensing technologies
495 continues to improve, it may become more cost effective to capture the spatial heterogeneity of
496 algal pigments in small lakes over time. However, one of the benefits of manual spatial sampling
497 is being able to pair other measurements such as dissolved oxygen, pH, and nutrients (e.g.,
498 nitrate; Loken et al., 2018; Pellerin et al., 2016) with information on the distribution of algal
499 biomass.

500 Our intensive spatial monitoring of a shallow, hypereutrophic lake revealed how spatially
501 heterogeneous shallow lakes are over the course of a single season and allowed us to tease apart
502 the drivers of that spatial heterogeneity. We found that variability was greatest during
503 biologically-intensive periods, such as during algal blooms and in dense floating-leaf macrophyte
504 beds, and that failure to capture this variability would have hampered our understanding of the

505 ecosystem's functioning and overall mean state. Small lakes such as Swan Lake dominate the
506 global distribution of waterbodies (Verpoorter et al. 2014). Adequately capturing and
507 characterizing the magnitude of variability in production of these waterbodies is important given
508 their role in mediating global nutrient cycles (Downing et al. 2010, Biddanda et al. 2017),
509 especially methane emissions (DeSontro et al. 2018, Loken et al. 2019). Our data provided an
510 estimate of the spatial resolution needed to capture the dynamics in ecosystems similar to Swan
511 Lake and a method which could be readily adapted to other ecosystems. While our results
512 provide new understanding of the magnitude and temporal dynamics of spatial heterogeneity in
513 shallow lakes, continued investigation of horizontal spatial heterogeneity in a range of aquatic
514 ecosystems, from oligotrophic to eutrophic, is needed to better understand the structure and
515 drivers of horizontal spatial variability in lakes.

516

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524

525 **Data Availability Statement**

526 The data that support the findings of this study are available from Environmental Data
527 Initiative repository: <https://portal.edirepository.org/nis/mapbrowse?packageid=edi.420.1>

528

529 **Conflict of Interest Statement**

530 There are no conflicts of interest to declare

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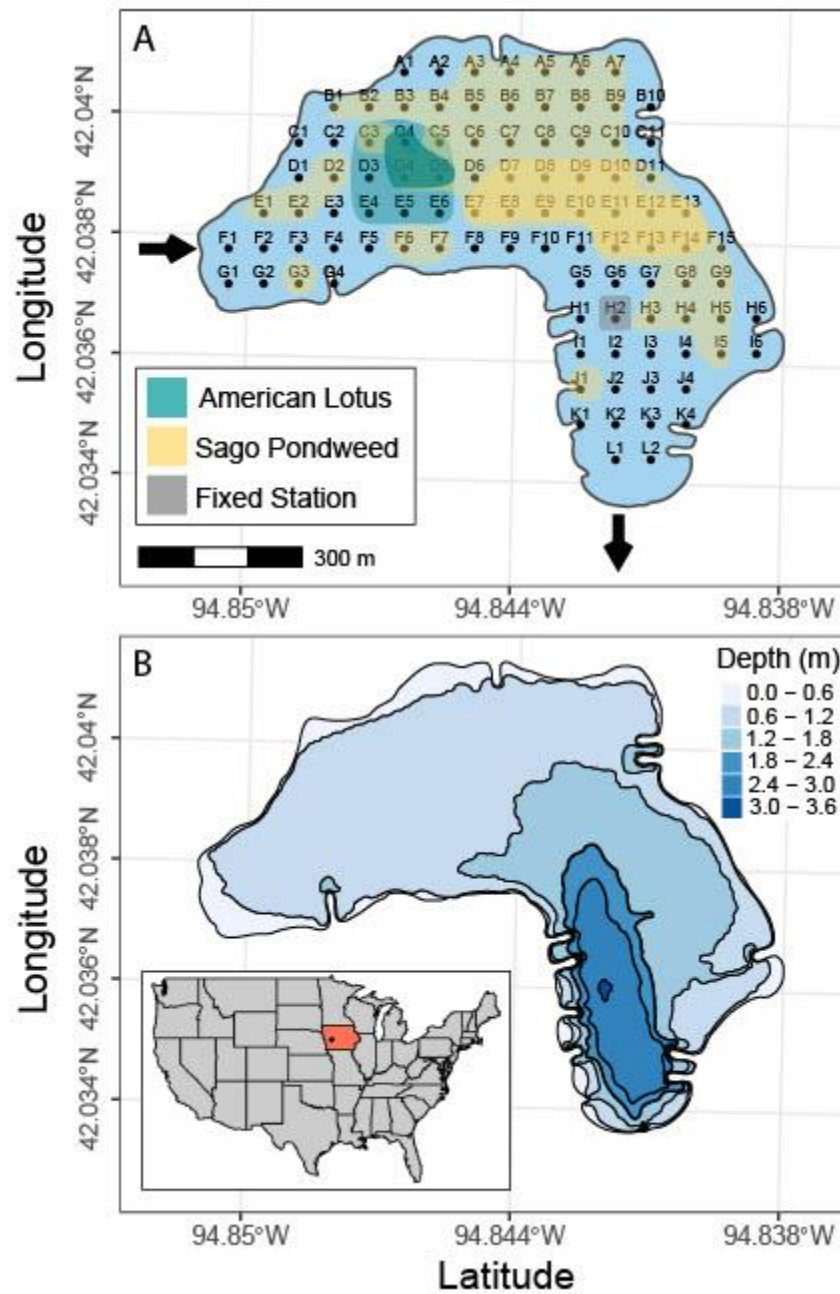
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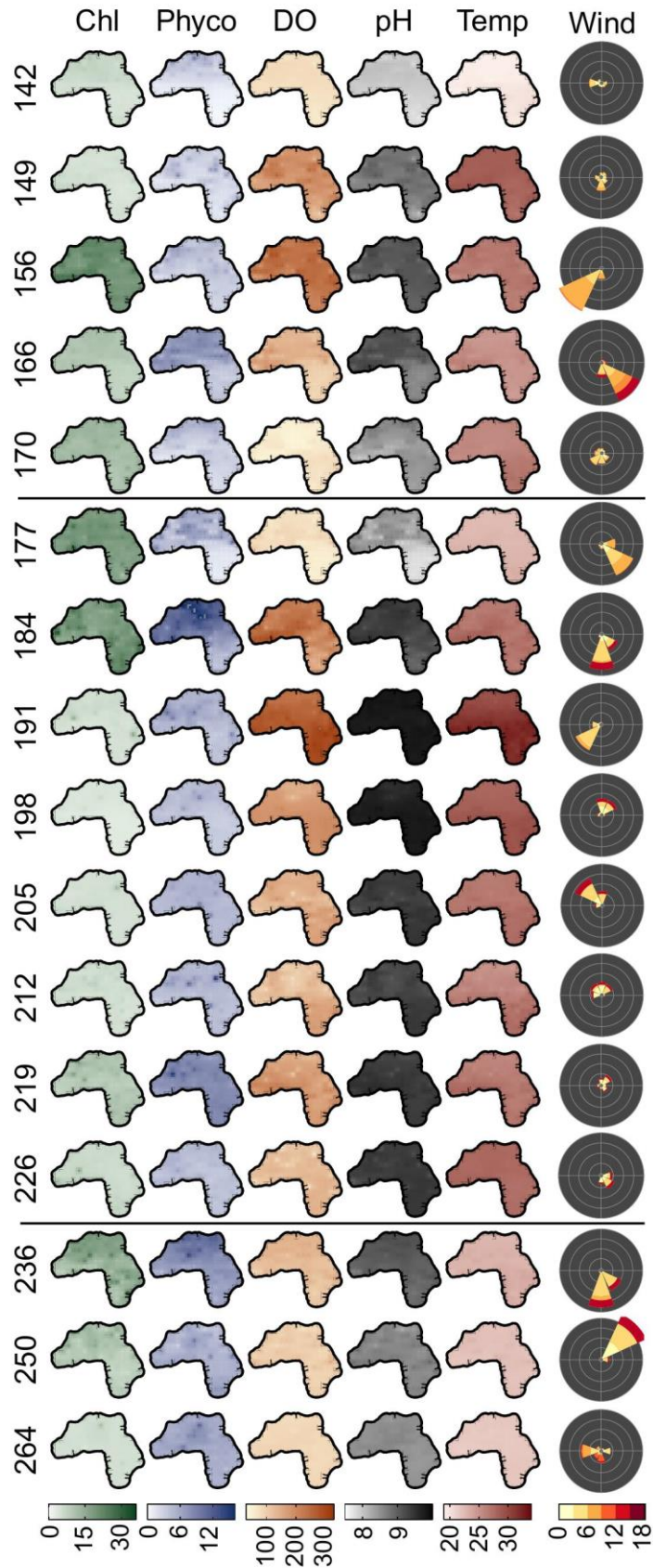
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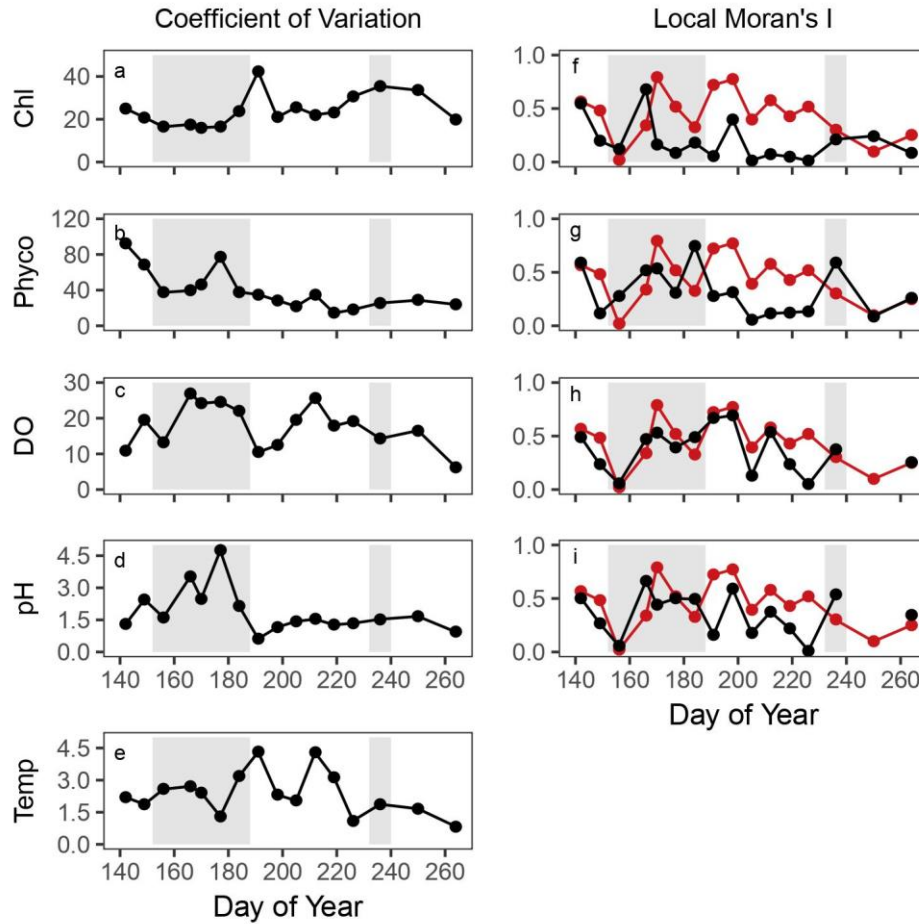
738 **Figure 1.** Sampling locations on a 65 m square grid of Swan Lake, a 40.5 hectare waterbody in
 739 western Iowa, USA. The main inlet to the lake and only outlet indicated with arrows. a) The
 740 location of the macrophyte beds of the two dominant species within the lake are shown on the
 741 map, with darker shading indicating the regions with the vegetation was always observed,

742 indicating permanent macrophyte beds, and the location of the high frequency sensor, b) the
743 bathymetry of the lake and location of the lake in the state of Iowa, in reference to the United
744 States of America.

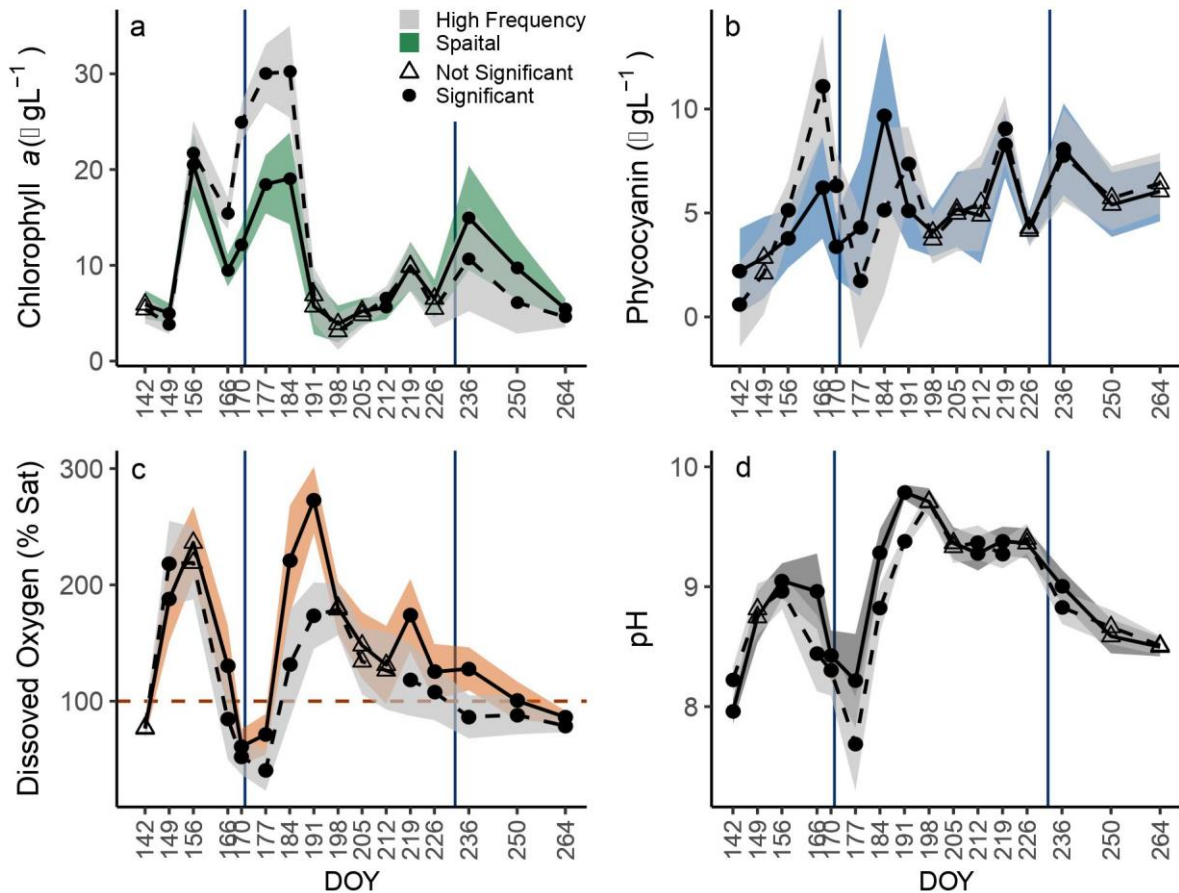


745 **Figure 2.** The spatial pattern of each of the variables chlorophyll *a* (Chl, $\mu\text{g L}^{-1}$), phycocyanin

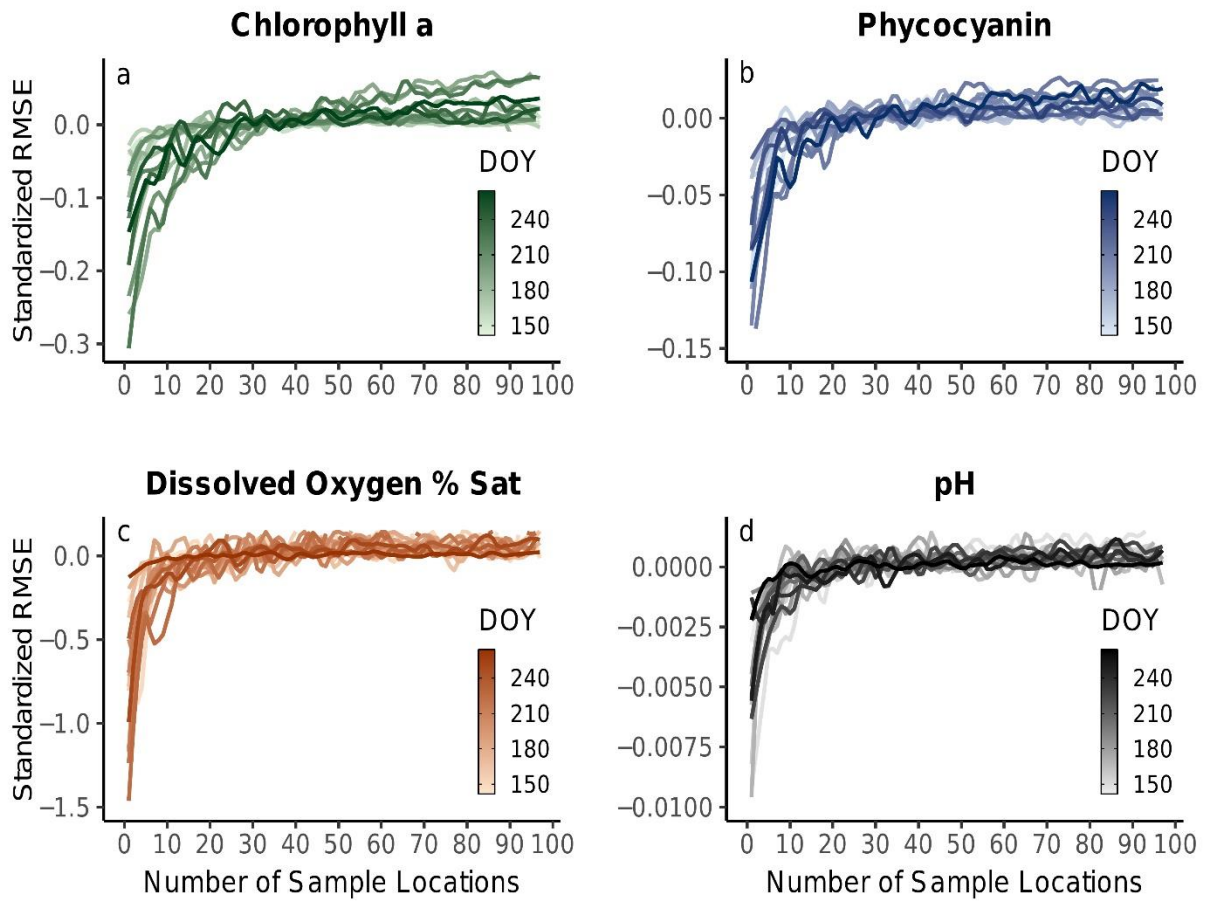
746 (Phyco, $\mu\text{g L}^{-1}$), dissolved oxygen (DO, percent saturation), pH, and temperature (Temp, $^{\circ}\text{C}$) for
747 each sampling event. The 98 sampling locations taken in a 65m grid (Figure 1) were interpolated
748 to a 25m grid using spatial inverse distance interpolation for visualization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



756 **Figure 3.** Time series of the spatial coefficient of variation (CV) and spatial autocorrelation (AC;
 757 local Moran's I) of the biologically-mediated variables in Swan Lake (same variable
 758 abbreviations as Figure 2). The gray polygons indicate periods of algal bloom. The red line is the
 759 time series of temperature local Moran's I for comparison.



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



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

1

Supplementary material for

2 **Title:** Capturing the spatial variability of algal bloom development in a shallow temperate lake

3 **Authors:** Ortiz and Wilkinson

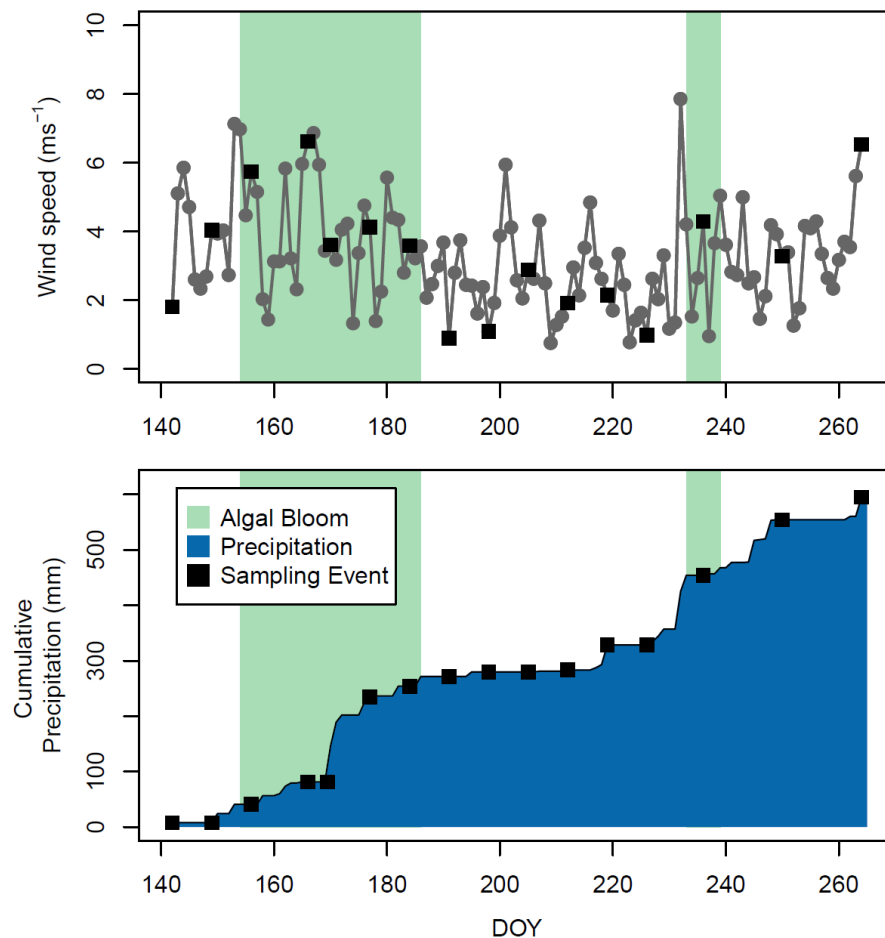
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Hourly Weather Data

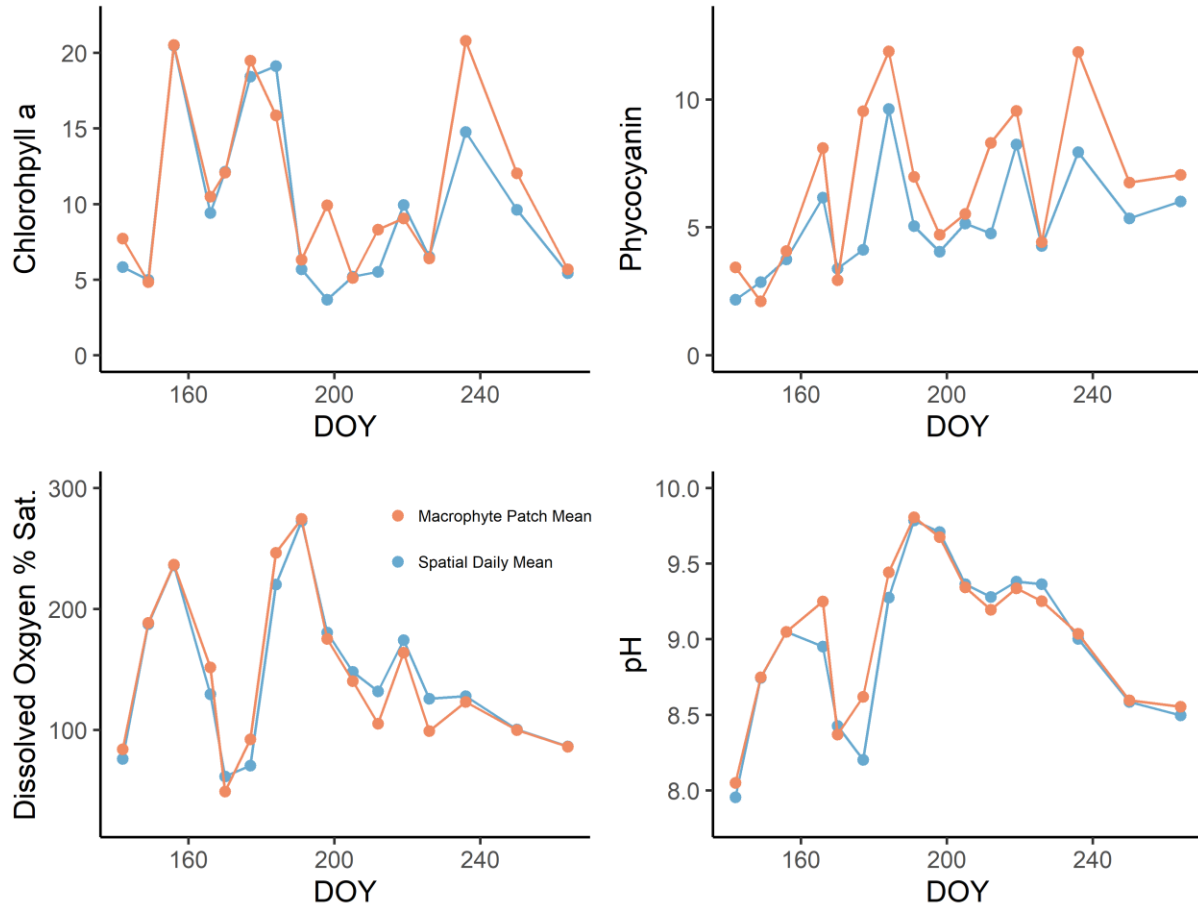
6 Hourly wind and precipitation data were downloaded from the National Oceanic and
7 Atmospheric Automated Surface Observing System (NOAA ASOS) for Arthur N. Neu Airport
8 Carroll, Iowa, USA, less than 5km from the lake, through the Iowa State University Iowa
9 Environmental Mesonet (<https://mesonet.agron.iastate.edu/>). The data were summarized to daily
10 means and plotted as mean wind speed and cumulative daily precipitation.

11 **Figures**



12

13 **Supplement Figure 1.** Daily meteorological data for the duration of the study. A) Daily mean
14 wind speed during the study period. Periods of algal bloom in the lake are denoted by the green
15 polygons. B) The cumulative precipitation from the first to the last day of the study period. There
16 were two large precipitation events from DOY 170-171 and on DOY 232.



17

18 **Supplement Figure 2.** A comparison of the mean value for each variable in the American lotus
 19 macrophyte patch (sites C4, D4, D5) and the rest of the lake.



20
21 **Supplement Figure 3.** Photographs from DOY 212 highlight the macrophyte densities. Photo A
22 is of the American Lotus patch and photo B shows the density of the Sag Pondweed. In photo
23 B, you can see behind the boat/rake that there are Sag Pondweed patches that are growing to the
24 water surface and creating mats.