Associations between Chlorophyll a and various Microcystin-LR Health Advisory Concentrations

Jeffrey W. Hollister * 1 Betty J. Kreakie 1

¹US Environmental Protection Agency, Office of Research and Development, National Health and Environmental Effects Research Laboratory, Atlantic Ecology Division, 27 Tarzwell Drive Narragansett, RI, 02882, USA

Cyanobacteria harmful algal blooms (cHABs) are associated with a wide range of adverse health effects that stem mostly from the presence of cyanotoxins. To help protect against these impacts, several health advisory levels have been set for some toxins. In particular, one of the more common toxins, microcystin, has several advisory levels set for drinking water and recreational use. However, compared to other water quality measures, field measurements of microcystin are not commonly available due to cost and advanced understanding required to interpret results. Addressing these issues will take time and resources. Thus, there is utility in finding indicators of microcystin that are already widely available, can be estimated quickly and in situ, and used as a first defense against high levels of microcystin. Chlorophyll a is commonly measured, can be estimated in situ, and has been shown to be positively associated with microcystin. In this paper, we use this association to provide estimates of chlorophyll a concentrations that are indicative of a higher probability of exceeding select health advisory concentrations for microcystin-LR. Using the 2007 National Lakes Assessment and a conditional probability approach, we identify chlorophyll a concentrations that are more likely than not to be associated with an exceedance of a microcystin health advisory level. We look at the recent US EPA health advisories for drinking water as well as the World Health Organization levels for drinking water and recreational use and identify a range of chlorophyll a thresholds. A 50% chance of exceeding one of the specific advisory microcystin concentrations of 0.3, 1, 1.6, and 2 μ g/L is associated with chlorophyll a concentration thresholds of 20.88, 69.6, 84.24, and 97.49, respectively. When managing for these various microcystin levels, exceeding these reported chlorophyll a concentrations should be a trigger for further testing and possible management action.

1 Introduction

13

14

15

21

22

23

Over the last decade, numerous events and legislative activities have raised the public awareness of harmful algal blooms (Rinta-Kanto et al. 2009, HABHRCA 2014, Jetoo et al. 2015). In response the US Environmental Protection Agency (USEPA) has recently released suggested microcystin-LR (one of the more common toxins) concentrations that would trigger health advisories (McElhiney and Lawton 2005, Zurawell et al. 2005, USEPA 2015). Additionally, the World Health Organization (WHO) has proposed microcystin advisory levels for drinking water and a range of recreational risk levels (Chorus

^{*} corresponding author: hollister.jeff@epa.gov

and Bartram 1999, Organization 2003). While these levels and associated advisories are likely to help mitigate the impacts from harmful algal blooms, they are not without complications.

One of these complications is that they rely on available measurements of microcystin-LR. While laboratory testing (e.g., chromatography) remains the gold standard for quantifying microcystin-LR concentrations in water samples, several field test kits have been developed. Even though field tests provide a much needed means for rapid assessment, they are not yet widely used and are moderately expensive (approximately \$150-\$200 depending on specific kit) with a limited shelf life (typically one year) (James et al. 2011, Aranda-Rodriguez et al. 2015). Additionally, each technique requires nuanced understanding of the detection method (e.g., limit of detection, specific microcystin variants being measured, and sampling protocol).

Fortunately, microcystin-LR has been shown to be associated with several other, more commonly measured and well understood components of water quality that are readily assessed in the field. For instance, there are small or hand held fluorometers that measure chlorohpyll a. Additionally, chlorophyll a is a very commonly measured component of water quality that is also known to be positively associated with microsystin-LR concentrations (Pip and Bowman 2014, Yuan et al. 2014). Yuan et. al (2014) explore these associations in detail and control for other related variables. In their analysis they find that total nitrogen and chlorophyll a show the strongest association with microcystin. Furthermore, they identify chlorophyll a and total nitrogen concentrations that are associated with exceeding 1 μ g/L of microcystin. Given these facts, it should be possible to identify chlorophyll a concentrations that would be associated with the new USEPA Microcystin-LR health advisory levels for drinking water. Identifying these associations would provide another tool for water resource managers to help manage the threat to public health posed by cHABs and would be especially useful in the absence of microcystin-LR concentrations.

In this paper we build on past efforts and utilize the National Lakes Assessment (NLA) data and identify chlorophyll a concentrations that are associated with higher probabilities of exceeding several microcystin-LR health advisory concentrations (Chorus and Bartram 1999, USEPA 2009, 2015). We add to past studies by exploring associations with newly announced advisory levels and by also applying a different method, conditional probability analysis. Utilizing different methods strengthens

the evidence for suggested chlorophyll *a* levels that are associated with increased risk of exceeding the health advisory levels as those levels are not predicated on a single analytical method. So that others may repeat or adjust this analysis, the data, code, and this manuscript are freely available via https://github.com/USAPE/microcystinchla.

68 2 Methods

69 **2.1** Data

We used the 2007 NLA chlorophyll *a* and microcystin-LR concentration data (USEPA 2009). These data represent a snapshot of water quality from the summer of 2007 for the conterminous United States and were collected as part of an ongoing probabilistic monitoring program (USEPA 2009). Data on chlorophyll *a* and microcystin-LR concentrations are available for lakes.

2.2 Conditional Probability Analysis

We used a conditional probability analysis (CPA) approach to explore associations between chlorophyll a concentrations and World Health Organization (WHO) and USEPA microcystin-LR health advisory levels (Paul and Munns 2011). Many levels have been suggested (Table 1), but lakes with higher microcystin-LR concentrations in the NLA were rare. Only 1.16 % of lakes sampled had a concentration greater than 10. Thus, for this analysis we focus on the microcystin concentrations that are better represented in the NLA data. These were the USEPA children's drinking water advisory level of 0.3 μ g/L (USEPA Child), the WHO drinking water advisory level of 1 μ g/L (WHO Drinking), the USEPA adult drinking water advisory level of 1.6 μ g/L (USEPA Adult), and the WHO recreational, low probability of effect advisory level of 2 μ g/L (WHO Recreational).

Conditional probability analysis provides information about the probability of observing one event given another event has also occured. For this analysis, we used CPA to examine how the conditional probability of exceeding one of the health advisories changes as chlorophyll a increases in a lake. We expect to find higher chlorophyll a concentrations to be associated with higher probabilities of exceeding

the microcystin-LR health advisory levels. We also caclulated bootstrapped 95% confidence intervals (CI) using 1000 bootstrapped samples. Thus, to identify chlorophyll a concentrations of concern we identify the value of the upper 95% CI across a range of conditional probabilities of exceeding each health advisory level. As both microcystin-LR and chlorophyll a values were highely skewed right, a log base 10 transformation was used. Additional details of the specific implementation are available at https://github.com/USEPA/microcystinchla. A more detailed discussion of CPA is beyond the scope of this paper, but see Paul et al. (2005) and Hollister et al. (2008) for greater detail. Lastly, all analyses were conducted using R version 3.2.2 and code and data from this analysis are freely available as an R package at https://github.com/USAPE/microcystinchla.

$_{\scriptscriptstyle 97}$ $\, \, 3 \quad { m Results}$

In the 2007 NLA, microcystin-LR concentrations ranged from 0.05 to 225 µg/L. Microcystin-LR concentrations of 0.05 μ g/L represent the detection limits. Any value greater than that indicates the presence of microcystin-LR. Of those lakes with microcystin, the median concentration was 0.51 and 100 the mean was 3.17. Of all lakes sampled, 21% of lakes exceeded the USEPA Child level, 8.8% of lakes 101 exceeded the USEPA Adult level, 11.7% of lakes exceeded the WHO Drinking level, and 7.3% of lakes 102 exceeded the WHO Recreational level. For chlorophyll a, the range was 0.07 to 936 μ g/L. All lakes had 103 reported chlorophyll a concentration that exceeded detection limits. The median concentration was 7.79 104 $\mu g/L$ and the mean was 29.63 $\mu g/L$. The associations between chlorophyll a and the upper confidence 105 interval across a range of conditional probability values is shown in Table 2. Specific chlorophyll a that 106 are associated with greater than even odds of exceeding the advisory levels were 0.07, 0.07, 2.89, and 107 10.27 for 0.3, 1.0, 1.6 and 2.0 µg/L advisory levels, respectively (Table 2 & Figure 2).

9 4 Discussion

The association between Log10 microcystin-LR and Log10 chlorophyll a shows a wedge pattern (Figure 1). This indicates that, in general, higher concentrations of microcystin-LR almost always co-occur with higher concentrations of chlorophyll a yet the inverse is not true. Higher chlorophyll a is not necessarily

predictive of higher microcystin-LR concentrations; however, chlorophyll *a* may be predictive of the probability of exceeding a certain threshold.

This is the case as the probability of exceeding each of the four tested health advisory levels increases as a function of chlorophyll a concentration (Figure 2). We use this association to identify chlorophyll a concentrations that are associated with a range of probabilities of exceeding a given health advisory level (Table 2). For the purposes of this discussion we focus on a conditional probability of 50% or greater (i.e. greater than even odds to exceed a health advisory level). The 50% conditional probability chlorophyll a thresholds represents 30.1%, 10.6%, 8.8%, and 7.9% of sample lakes for the USEPA Child, the WHO Drinking, the USEPA Adult, and the WHO recreational levels, respectively.

Furthermore, the chlorophyll *a* cutoffs may be used to predict whether or not a lake exceeds the microcystin-LR health advisories. Doing so allows us to compare the accuracy of the prediction as well as evaluate false negatives. Total accuracy of the four cutoffs predicting microcystin-LR exceedances were 74% for the USEPA children's advisory, 86% for the WHO drinking water advisory, 89% for the USEPA adult advisory and, 90% for the WHO regreational advisory (Tables 3, 4, 5, & 6). However, total accuracy is only one part of the prediction performace with which we are concerned.

When using the chlorophyll *a* cutoffs as an indicator of microcystin-LR exceedances, the error that should be avoided is predicting that no exceedance has occurred when in fact it has. In other words, we would like to avoid Type II errors and minimize the proportion of false negatives. For the four chlorophyll *a* cut-offs we had a proportion of false negatives of 9%, 8%, 6% and , 5% for the U.S EPA childrens drinking water, the WHO drinking water, the USEPA adult drinking water, and the WHO recreational advisories, respectively. In each case we miss less than 10% of the lakes that are in fact exceeding the microcystin-LR advisory.

There are numerous possible uses for the chlorophyll a and microcystin-LR advisory cut-off values.

First, in the absence of microcystin-LR measurements, exceedence of the chlorophyll a concentrations

could be a trigger for further actions. Given that there is uncertainty around these chlorophyll a cutoffs

the best case scenario would be to monitor for chlorophyll a and in the event of exceeding a target

concentration take water samples and have those samples tested in a lab for microcystin-LR.

A second potential use is to identify past bloom events from historical data. As harmful algal blooms are made up of many species and have various mechanisms responsible for adverse impacts (e.g. toxins, hypoxia, odors), there is no single definition of a bloom. For cHABs one approach has been to identify an increase over a baseline concentration of phycocyanin (Miller et al. 2013). This is a useful approach for targeted studies, but phycocyanin is also not always available and measures the predominance of cyanobacterial pigments and not toxins. Using our chlorophyll *a* cutoffs provides a value that is more directly associated with microcystin-LR and can be used to classify lakes, from past surveys, as having bloomed.

Lastly, using chlorophyll a is not meant as a replacement for testing of microcystin-LR or other toxins.

It should be used when other, direct measurements of cyanotoxins are not available. In those cases,
which are likely to be common at least in the near future, using a more ubiquitous measurement, such as
chlorophyll a will provide a reasonable proxy for the probability of exceeding a microcystin-LR health
advisory level and provide better protection against adverse effects in both drinking and recreational
use cases.

$_{54}$ 5 Acknowledgements

¹⁵⁵ 6 Figures

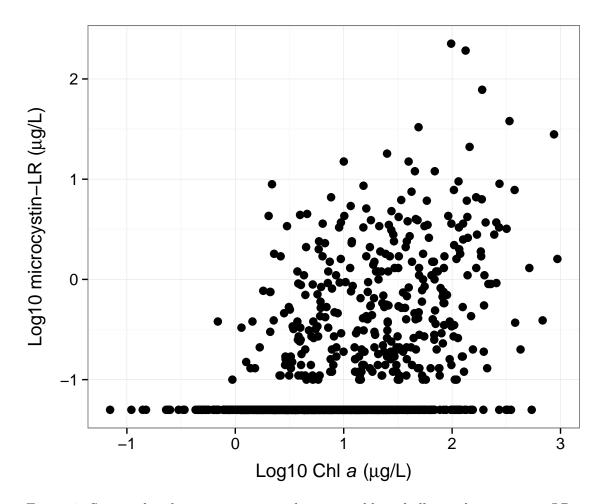


Figure 1: Scatterplot showing association between chlorophyll a and microcystin-LR.

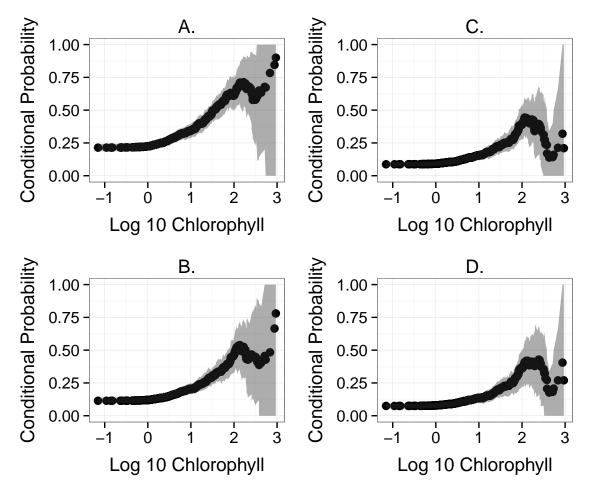


Figure 2: Conditional probability plots showing association between the probability of exceeding various microcystin-LR (MLR) health advisory Levels. A.) Plot for USEPA Child (0.3 μ g/L). B.) Plot for WHO Drinking (1 μ g/L). C.) Plot for USEPA Adult (1.6 μ g/L). D.) Plot for WHO Recreational (2 μ g/L).

Tables

Table 1: Various suggested microcystin-LR health advisory concentrations.

Source	Type	Concentration
USEPA	Adult Drinking Water Advisory	$1.6 \ \mu \mathrm{g/L}$
USEPA	Child Drinking Water Advisory	$0.3~\mu\mathrm{g/L}$
WHO	Drinking Water	$1~\mu\mathrm{g/L}$
WHO	Recreational: High Prob. of Effect	20-2000 $\mu \mathrm{g/L}$
WHO	Recreational: Low Prob. of Effect	$2\text{-}4~\mu\mathrm{g/L}$
WHO	Recreational: Moderate Prob. of Effect	$10\text{-}20~\mu\mathrm{g/L}$
WHO	Recreational: Very High Prob. of Effect	$>$ 2000 $\mu g/L$

Table 2: Chlorophyll a concentrations that are associated with a 50% probability of exceeding a microcystin-LR health advisory concentration.

WHO Recreational	USEPA Adult	WHO Drink	USEPA Child	Cond. Probability
1.17	0.07	0.07	0.07	0.1
16.80	11.84	4.38	0.07	0.2
45.22	32.11	17.49	2.89	0.3
76.80	66.96	38.30	10.27	0.4
97.49	84.24	69.60	20.88	0.5
125.40	114.62	100.51	38.20	0.6
871.20	871.20	128.74	65.60	0.7
871.20	871.20	271.44	114.62	0.8
871.20	871.20	338.40	166.63	0.9

Table 3: Confusion matrix comparing chlorophyll a predicted exceedences (rows) versus real exceedances (columns) for the USEPA childrens drinking water advisory.

	FALSE	TRUE
FALSE	632	90
TRUE	179	127

Table 4: Confusion matrix comparing chlorophyll a predicted exceedences (rows) versus real exceedances (columns) for the WHO drinking water advisory.

	FALSE	TRUE
FALSE	844	80
TRUE	63	41

Table 5: Confusion matrix comparing chlorophyll a predicted exceedences (rows) versus real exceedances (columns) for the USEPA adult drinking water advisory.

FALSE	TRUE
885	57
52	34
	885

Table 6: Confusion matrix comparing chlorophyll a predicted exceedences (rows) versus real exceedances (columns) for the WHO recreational water advisory.

	FALSE	TRUE
FALSE	903	48
TRUE	50	27

157 References

- ¹⁵⁸ Aranda-Rodriguez, R., Z. Jin, J. Harvie, and A. Cabecinha. 2015. Evaluation of three field test kits to
- detect microcystins from a public health perspective. Harmful Algae 42:34–42.
- 160 Chorus, E. I., and J. Bartram. 1999. Toxic cyanobacteria in water: A guide to their public health
- 161 consequences, monitoring and management.
- 162 HABHRCA. 2014. Harmful algal bloom and hypoxia research and control amendments act of 2014.
- Hollister, J. W., H. A. Walker, and J. F. Paul. 2008. CProb: A computational tool for conducting
- conditional probability analysis. Journal of environmental quality 37:2392–2396.
- James, R., A. Gregg, A. Dindal, and J. McKernan. 2011. Environmental technology verification report:
- Abraxis microcystin test kits. Online document. Accessed online: June 22.
- Jetoo, S., V. I. Grover, and G. Krantzberg. 2015. The toledo drinking water advisory: Suggested
- application of the water safety planning approach. Sustainability 7:9787–9808.
- McElhiney, J., and L. A. Lawton. 2005. Detection of the cyanobacterial hepatotoxins microcystins.
- Toxicology and Applied Pharmacology 203:219–230.
- Miller, T. R., L. Beversdorf, S. D. Chaston, and K. D. McMahon. 2013. Spatiotemporal molecular
- analysis of cyanobacteria blooms reveals microcystis-aphanizomenon interactions. PloS one 8:e74933.
- Organization, W. H. 2003. Cyanobacterial toxins: Microcystin-lR in drinking-water. background
- document for development of wHO guidelines for drinking-water quality, geneva, switzerland. World
- 175 Health Organization, 2nd ed. Geneva.
- Paul, J. F., and M. E. McDonald. 2005. Development of empirical, geographically specific water quality
- criteria: A conditional probability analysis approach 41:1211–1223.
- Paul, J. F., and W. R. Munns. 2011. Probability surveys, conditional probability, and ecological risk
- assessment. Environmental Toxicology and Chemistry 30:1488–1495.

- Pip, E., and L. Bowman. 2014. Microcystin and algal chlorophyll in relation to nearshore nutrient concentrations in lake winnipeg, canada. Environment and Pollution 3:p36.
- Rinta-Kanto, J. M., E. A. Konopko, J. M. DeBruyn, R. A. Bourbonniere, G. L. Boyer, and S. W.
- Wilhelm. 2009. Lake erie microcystis: Relationship between microcystin production, dynamics of
- genotypes and environmental parameters in a large lake. Harmful Algae 8:665–673.
- USEPA. 2009. National lakes assessment: A collaborative survey of the nation's lakes. ePA 841-r-09-001.
- Office of Water; Office of Research; Development, US Environmental Protection Agency Washington,
- 187 DC.
- USEPA. 2015. Drinking water health advisory for the cyanobacterial microcystin toxins. ePA-820R15100.
- Office of Water, US Environmental Protection Agency Washington, DC.
- Yuan, L. L., A. I. Pollard, S. Pather, J. L. Oliver, and L. D'Anglada. 2014. Managing microcystin:
- 191 Identifying national-scale thresholds for total nitrogen and chlorophyll a. Freshwater Biology 59:1970—
- 192 1981.
- ¹⁹³ Zurawell, R. W., H. Chen, J. M. Burke, and E. E. Prepas. 2005. Hepatotoxic cyanobacteria: A review
- of the biological importance of microcystins in freshwater environments. Journal of Toxicology and
- Environmental Health, Part B 8:1–37.