POTENTIAL TOXIC CYANOBACTERIA IN THREE LAKES OF JAKARTA-BOGOR AREA, INDONESIA

Nining Betawati Prihantini*, Arya Widyawan, Ronny Rianto, Yuni Ariyani, Wisnu Wardhana, Dian Hendrayanti

Department of Biology
Faculty of Mathematics and Natural Science, University of Indonesia
Depok 16424 Indonesia
*e-mail: nprihantini@hotmail.com

ABSTRACT

The research was aimed to understand the composition and dominancy of aquatic microalgae, and to know the potential toxicity of Cyanobacteria occuring in three lakes of Jakarta-Bogor area. The study was done in 2006. The sampling sites were Lake Sunter 2 (North Jakarta), Lake Babakan (South Jakarta), and Lake Lido (Bogor). The water samples were taken using plankton-net (20 µm mesh) with horizontal tow. The results showed that the number of microalgal genera in Sunter 2, Babakan, and Lido were different, i.e. 10 genera, 40 genera, and 14 genera, respectively. Cyanobacteria were dominant in Sunter 2 (96%) and Babakan (90.01%). The dominant Cyanobacteria in Sunter 2 was *Planktothrix agardhii* (53.48%), whereas in Babakan was *Chroococcus dispersus* (68.52%). The second abundant Cyanobacteria in Sunter 2 was *Arthrospira* (42.54%). *Planktothrix agardhii* and *Arthrospira* are known to produce harmful toxin.

Keywords: Sunter 2, Babakan, Lido, Planktothrix, Arthrospira.

INTRODUCTION

Cyanobacteria or Cyanophyta is one of the microalgal divisions that often occurs abundantly in the fresh, brackish, or marine waters. It becomes a part of Harmful Algal Bloom (HAB) phenomenon. These microorganisms have been known to have wide range of habitat condition compared to other aquatic microalgae. This character causes Cyanobacteria to grow in extreme environmental condition. Usually optimal growth occurs if the water temperature is warm enough (25-35°C), acquire enough light intensity, the water is alkaline (pH 7.5-9), and nutrient concentration increase during eutrophication (Nicholas, 1980).

Almost all members of Cyanobacteria can produce toxin and are usually called toxic Cyanobacteria, for examples, *Microcystis*, *Anabaena*, *Planktothrix / Oscillatoria*, *Nostoc*, *Aphanizomenon*, and *Cylindrospermopsis* (Carmichael, 1995). These toxins occur in the waters because dead microalgae release chemical

substance (toxins) into the waters. Toxins can be hepatotoxic (microcystin and nodularin) and neurotoxic (anatoxin). Microcystin that can damage liver is produced by *Microcystis aeruginosa*, whereas anatoxin, which can damage the nervous system, is produced by *Anabaena flos-aquea* and *Aphanizomenon flos-aquea* (Carmichael, 1995; Hoek *et al.*, 1995).

Cyanobacteria occur abundantly in several lakes and ponds in and around Jakarta. Bluishgreen scums cover many water surfaces which indicates that Cyanobacteria is blooming in the waters. In Depok (near Jakarta) for example, Cyanobacteria were dominated in Baru Pond in 1994 (Budiman, 1995), and Rawa Besar and Rawa Kalong Pond in 1998 (Junwinanto, 1998). A preliminary research in Lakes Sunter 2 and Sunter Barat showed that *Plantothrix agardhii* was dominant in 2003. This species also occurred in the ponds of University of Indonesia (Prihantini *et al.*, 2006; Rianto *et al.*, 2006).

Although Cyanobacteria seem to be dominant in several lakes and river, the abundance of the scum does not spread everywhere. The explorations on aquatic microalgae in three lakes of Jakarta-Bogor area were done to understand the composition and dominancy of aquatic microalgae, and to know the potential of toxic Cyanobacteria, in those three lakes.

MATERIAL AND METHODS

The samples were taken from Lake Babakan (South Jakarta) in August 2006, and from Lake Sunter 2 (North Jakarta) and Lake Lido (Bogor) in September 2006. All samples were collected between 10.00 am to 12.00 am (Western Indonesia Time). Sampling site was decided with purposive sampling, and conducted at three stations, i.e. *inlet, midlet*, and *outlet* of the lakes. The water was sampled using plankton net with horizontal tow in triplicate. The samples were fixed with 6%. Fresh samples were also taken in order to isolate the Cyanobacteria and to support identification.

Environmental parameters were measured at every station. The parameters were temperature, pH, light intensity, transparency, DO, conductivity, N and P concentrations. DO values was measured using DO meter, whereas N and P concentrations were determined at laboratory following the methods described in Hutagalung *et al.* (1997).

Identification was done with Nikon SE light microscope with gradual magnification from (10x4),(10x10),(10x40),and (10x100). Identification references used were Edmonson (1959), Pentecost (1984), Geitler (1985), Santra (1993), and Whitton (2002). The density of microalgae m³ (D) was estimated from the sub-samples and application of the formula, $D = q \times 1/f \times 1/v$

(Wickstead, 1965), where q = cell counts, f = fraction examined, and v = volume of water filtered (m³). V was estimated by the formula, $v = \pi r^2 x$ t (Greenberg *et al.*, 1992), where r is plankton-net radius (10 cm), and t is distance tow (5m).

RESULTS

The results of environmental measurement in the three lakes were different. The data showed that water temperature in Lake Sunter 2 (28.5 to 32°C) was warmer than other two lakes. The pH in Lake Sunter 2 was alkaline (8), and light intensity was more than 500,000 lux. Beside that, the dissolved oxygen and conductivity were higher than the other two lakes. Other environmental data can be seen at Table 1.

The numbers of microalgae found in the three lakes are shown in Table 2. Ten genera were found in Lake Sunter 2, 40 genera in Lake Babakan, and 14 genera in Lido Lake.

The total numbers of genera identified easily were 45 comprising with Chromophyta (seven genera), Chlorophyta (22 genera), Cyanobacteria / Cyanophyta (14 genera), Dinophyta (one genus), and Euglenophyta (three genera).

The Cyanobacteria were found dominantly in Sunter 2 and Babakan (Fig. 1). Two genera were found blooming in Sunter 2 i.e. *Planktothrix* (53.48 %) and *Arthrospira* (42.54 %), whereas *Choococcus dispersus* was blooming in Babakan (68.52%). Lido was not dominated by Cyanobacteria, but *Chlamydomonas* (Chlorophyta) was blooming in the lake (70.15 %).

Table 1. Environmental data of Lakes Sunter 2, Babakan, and Lido

Sampling site	Stasiun	Temp (°C)	Light Int. (lux)	Transpa rancy (cm)	рН	DO	Conduc tivity (µMHos)	N cont. (ppm)	Pcont. (ppm)
Sunter 2	inlet	28.5	>500000	14	8	7.4	1000	11.81	0.23
Lake	outlet	30.5	>500000	15	8	13.9	1150		
	centre	32	>500000	16	8	10.3	1200		
Babakan Lake	inlet	27	163100	27	7	4.0	310	12.75	0.08
	outlet	29	193700	37	7	4.7	250		
	centre	29	>500000	29	7	8.4	250		
Lido Lake	inlet	27	>50000	170	6	6.3	168	21.4	0.17
Ν	outlet	27	>50000	207	6	7.3	170		
	centre	27	>50000	27	6	4.1	171		

Table 2. Composition and density of plankton in the lakes of Sunter 2, Babakan and Lido

No.	Division	Genus/Species	Density (%)				
	DIVISION	denus/opecies	Lake Sunter 2	Lake Babakan	Lake Lido		
1	Chromophyta	Cyclotella	0.00	0.79	0.00		
2	Chromophyta	Cymbella	0.00	0.00	0.10		
3	Chromophyta	Melosira	0.00	0.14	19.69		
4	Chromophyta	Navicula	0.00	0.25	0.00		
5	Chromophyta	Nitzchia	0.00	0.29	0.00		
6	Chromophyta	Pinnularia	0.00	0.03	0.10		
7	Chromophyta	Pseudonitszchia sp.	0.22	0.00	0.00		
8	Chlorophyta	Actinastrum	0.00	0.29	0.00		
9	Chlorophyta	Ankyra	0.00	0.00	0.10		
10	Chlorophyta	Chlamydomonas	0.00	0.63	70.15		
11	Chlorophyta	Chlorella	0.00	0.04	0.00		
12	Chlorophyta	Closterium	0.00	0.01	0.05		
13	Chlorophyta	Coelastrum	0.58	0.13	1.35		
14	Chlorophyta	Coelosphaerium	0.00	0.49	0.00		
15	Chlorophyta	Crucigenia	0.00	0.15	0.00		
16	Chlorophyta	Crucigeniella	0.00	0.19	0.00		
17	Chlorophyta	Dictyosphaerium	0.00	0.99	0.34		
18	Chlorophyta	Golenkinia	0.00	0.01	0.00		
19	Chlorophyta	Kirchneriella	0.00	0.04	0.00		
20	Chlorophyta	Micractinium	0.73	0.00	0.00		
21	Chlorophyta	Monaraphidium	0.00	0.16	0.00		
22	Chlorophyta	Pandorina	0.11	0.06	4.06		
23	Chlorophyta	Pediastrum	0.00	0.16	0.00		
24	Chlorophyta	Scenedesmus	0.02	1.83	0.00		
25	Chlorophyta	Selenastrum	0.00	0.03	0.00		
26	Chlorophyta	Staurastrum	0.00	0.07	0.00		
27	Chlorophyta	Tetrastrum	0.00	0.14	0.00		
28	Cyanobacteria	Aphanothece	0.00	0.08	0.00		
29	Cyanobacteria	Arthrospira	42.54	7.75	0.00		
30	Cyanobacteria	Chroococcus	0.00	1.23	0.00		
31	Cyanobacteria	Chroococcus dispersus	0.00	68.52	0.00		
32	Cyanobacteria	Gloeocapsa sp.	0.00	0.00	0.48		
33	Cyanobacteria	Merismopedia	0.00	2.75	0.00		
34	Cyanobacteria	Microcystis aeruginosa	0.00	2.89	0.68		
35	Cyanobacteria	Microcystis sp.	0.00	1.97	0.00		
36	Cyanobacteria	Oscillatoria sp.1	0.00	0.38	0.29		
37	Cyanobacteria	Oscillatoria sp.3	0.00	2.88	0.00		
38	Cyanobacteria	Planktothrix agardhii	53.48	0.25	0.00		
39	Cyanobacteria	Romeria	0.00	0.83	1.40		
40	Cyanobacteria	Spirulina	0.00	0.30	0.00		
41	Cyanobacteria	Synechococcus	0.00	0.18	0.00		
42	Dinophyta	Peridinium	0.00	0.07	0.00		
43	Euglenophyta	Euglena	2.29	0.90	0.00		
44	Euglenophyta	Phacus	0.03	0.07	0.00		
45	Euglenophyta	Trachelomonas	0.01	2.03	1.21		
otal (Genera		10	40	14		

DISCUSSION

Among the 45 genera found in the three lakes, 14 genera were belonging to the division Cyanobacteria. In Lake Babakan, 13 genera of Cyanobacteria were common, whereas in Lake Lido were four genera, and in Lake Sunter 2 only two genera were found.

Even though only two genera of Cyanobacteria were easily found in Sunter 2, they were dominant.

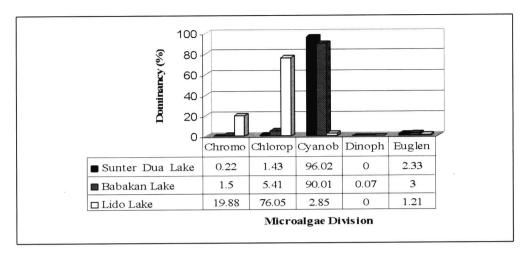


Figure 1. Microalgal dominancy in Lakes Sunter 2, Babakan, and Lido

The genus or species Planktothrix / Planktothrix agardhii posses the highest densities (53.48%), whereas Arthrospira become the second majority (42.54 %). The abundant Cyanobacteria (Planktothrix and Arthrospira) in the Sunter 2 probably were caused by the water condition. Cyanobacteria have been known to have range of wide habitat condition if compared to others aquatic microalgae. The water condition of Sunter 2 support Cyanobacteria to grow abundantly, especially because of pH 8, water temperature range from 28.5 to 32°C, and light intensity more than 500,000 lux. The water pH can inhibit the distribution of Cyanobacteria (Sze, 1998). Generally, Cyanobacteria are uncommon in acidic pH waters, but tend to grow better in neutral and alkaline pH, and enough light intensity (Nicholas, 1980). More over, polluted lakes in tropical country illuminated by high sun light intensity, and high temperature can supply optimal condition for the growth of Cyanobacteria (Kumar and Singh, 1979). These optimal conditions will be conducive if there is an increasing P and N concentration in the water (Odum, 1993). Finally, seasons are also able to influence the occurrence of microalgae blooms (Willen and Willen, 1978).

Both genera mentioned above are known to produce toxin, which is harmful to the lake. The toxins produced includes anatoxins (neurotoxins), microcystin (hepatoxins), and saxitoxin (neurotoxins) (Carmichael, 1995). Recent studies have shown that *Planktothrix* produces potent toxins in addition to the well-characterised microcystin (Blom *et al.*, 2003; Tonk *et al.*, 2005;

Lance et al., 2006). Beside that, Arthrospira is also potentially toxic Cyanobacteria. Like Planktothrix, most members of this genus also can produce hepatotoxins and neurotoxins (Carmichael, 1995; Stewart et al., 2006).

A member of Cyanobacteria (90.01%) also dominated in Babakan, which *Chroococcus dispersus* contributed about 68.52%. Some potential toxic Cyanobacteria found in Babakan were *Arthrospira* (7.75%), *Microcystis* aeruginosa (2.89%), *Microcystis* sp. (1.97%), and *Planktothrix agardhii* (0.25%). Cyanobacteria was not only the dominant genera, but also the highest genera number compared to those in Sunter 2 (10 genera) and Lido (14 genera). This condition maybe occur because the domination of *Chroococcus* does not influence the growth of other genera in the lake.

The water condition of Babakan was a little bit different from Sunter 2. Water temperature, pH, and light intensity were from 27 to 29°C, 7, and 163,100 to more than 500,000 lux in range, respectively. This condition is optimal for Cyanobacteria growth, and *Chroococcus dispersus* was blooming at that time. The Lake Babakan is relatively not polluted because it is a tourism area and treated with a good management.

Lido has different condition from the other two lakes, especially for water temperature and pH. Water temperature was cooler than in Sunter 2 and Babakan. The pH was also more acidic than the other two lakes. These conditions seem to be favourable for *Chlamydomonas* (division Chlorophyta) which contributed up to 70.15%.

Eventhough enough light intensity (more than 500,000 lux), Cyanobacteria were not common to be found in Lido. Only four genera were found, i.e. *Gloeocapsa* sp. (0.48%), *Microcystis aeruginosa* (0.68%), *Oscillatoria* sp 1 (0.29%), and *Romeria* (1.40%). Isolation of all these Cyanobacteria have been done for culturing in order to study the toxicity of those Cyanobacteria.

CONCLUSION

The composition of aquatic microalgae comprise of ten genera in Sunter 2, 40 genera in Babakan, and 14 genera in Lido. These genera belong to five division of microalgae i.e. Chromophyta (class Bacillariophyceae), Chlorophyta, Cyanobacteria, Dinophyta, and Euglenophyta. The dominant genus/species was Planktothrix agardhii (Cyanobacteria) in Sunter 2, Chroococcus dispersus (Cyanobacteria) in Babakan, and Chlamydomonas (Chlorophyta) in Lido. Some potential toxic Cyanobacteria were found in these three lakes i.e. Planktothrix agardhii and Arthrospira in Sunter 2, Planktothrix agardhii, Arthrospira, Microcystis aeruginosa and Microcystis sp1 in Babakan, and Microcystis aeruginosa in Lido

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REFERENCES

- Blom, J.F., B. Bojan, D. Bischoff, G. Nicholso, G. Jung, R. Sfissmuth, and F. Juttner. 2003. Oscillapeptin a new grazer toxin of the freshwater cyanobacterium Planktonic rubescens. *J. Nat. Prod.*, 66: 431–434.
- Budiman, F.R. 1995. Produkstivitas primer dan komunitas fitoplankton situ baru dan situ rawa besar, Depok, Jawa barat. Skripsi Jurusan Biologi FMIPA UI 1995: viii + 66p.
- Carmichael, W.W. 1995. Cyanobacterial toxins. *In:* Hallegraeff, G.M., D.M. Anderson. & A.D. Cembella (eds). 1995. *Manual on harmful marine microalgae*. Unesco, Paris: 163–175p.
- Edmondson, W.T. 1959. *Freshwater biology*. 2nd ed. John Wiley and Sons Inc., New York: 1203p.

- Greenberg, A.E., L.S. Clesceri and A.D. Eaton. 1992. Standard methods for the examination of water and wastewater. 18th ed. American Public Health Association, Washington: xxxi + 10–137p.
- Geitler, L. 1985. *Cyanophyceae*. Koeltz Scientific Books, Koenigstein: vi + 1196p.
- Hoek, C. Van den, D.G. Mann and H.M. Jahns. 1995. *Algae: An introduction to phycology*. Cambridge University Press, Melbourne: xi + 623p.
- Hutagalung, H.P., D. Setiapermana and S. H. Riyono (eds.). 1997. *Metode analisis air laut, sedimen dan biota. Buku* 2. Pusat Penelitian dan Pengembangan Oseanologi, Lembaga Ilmu Pengetahuan Indonesia, Jakarta: x + 182p.
- Junwinanto. 1998. Studi perbandingan struktur komunitas fitoplankton di situ rawa besar dan situ rawa kalong kabupaten Bogor, Jawabarat. Skripsi Jurusan Biologi FMIPA UI 1998: ix + 78p.
- Kumar, H.D. and F.N. Singh. 1979. *A textbook on algae*. The Macmillan Press, LTD, Tokyo: vii + 216p.
- Lance, E., L. Brient, M. Bormans, and C. Gerard. 2006. Interactions between cyanobacteria and gastropods I. Ingestion of toxic *Plantothrix agardhii* by Lymnaea stagnalis and the kinetics of microcystin bioaccumulation and detoxification. *Aquat. Toxicol.*, 79: 140–148.
- Nicholas, D.J.D. 1980. Mineral nutrients requirements of utilization by algal flora of freshwater lake. Technical Paper No. 50. Research project no 74/72, Australian Government Publishing Service, Canberra: viii + 51p.
- Odum, E.P. 1993. *Dasar-dasar ekologi*. Ed ke-3. Terjemahan. Dari Fundamentals of Ecology, oleh Samingan, Tjahjono. Gadjah Mada Uiversity Press, Yogyakarta: xv +697p.
- Pentecost, A. 1984. *Introduction to fresh water algae*. Richmond Publishing Co. Ltd., England: vii + 247p.
- Prihantini, N.B., W. Wardhana, A. Widyawan, and R. Rianto. 2006. Cyanobacteria dari beberapa situ dan sungai di kawasan Jakarta dan Depok, Indonesia. Prosiding Seminar Nasional Limnologi 2006, Puslit Limnologi, LIPI, Jakarta: 210-221.
- Rianto, R., A. Widyawan, N.B. Prihantini, W. Wardhana, and D. Hendrayanti. 2006. Aquatic microalgae in Agathis and Kenanga Ponds of University of Indonesia, Depok: Focus on Cyanobacteria division. Proc. of International Conference on Mathematics and Natural Sciences (ICMNS) Bandung, 2006: 318–321.
- Santra, S.C. 1993. *Biology of Rice-fields blue-green algae*. Daya Publishing House, New Delhi: vii + 229p.

- Sze, P. 1998. *A biology of the algae*. 3rd ed.WCB. McGraw-Hill, Boston: viii + 278p.
- Stewart, Ian, P.M Webb, P.J Schluter and G.R Shaw. 2006. Recreational and occupational field exposure to freshwater cyanobacteria a review of anecdotal and case reports, epidemiological studies and the challenges for epidemiologic assessment. BioMed Central Ltd., 5: 6.
- Tonk L., P.M. Visser, G. Christiansen, E. Dittmann, E.O.F.M. Snelder, C. Wiedner, L.R. Mur, and J. Huisman. 2005. The microcystin composition of the cyanobacterium Planktothrix agardhii changes toward a more toxic variant with increasing light intensity. *Appl. Environ. Microbial.*, 71: 5177–5181.
- Whitton, B.A. 2002. Phylum Cyanophyta (Cyanobacteria). *In:* Jhon, D.M. B.A. Whitton & A.J. Brook (eds). The freshwater alga flora of The British Isles: an identification guide to freshwater and terrestrial algae. Cambridge University Press, Cambridge: p.25–122.
- Wickstead, J.H. 1965. An introduction to the study of tropical plankton. Hutchinson Tropical Monographs, London: 160p.
- Willen, E. and T. Willen. 1978. About freshwater phytoplankton. *In*: Phytoplankton manual. UNESCO, Paris: 297–301.