Microcystin Production in the Tripartite Cyanolichen *Peltigera leucophlebia*

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Submitted 30 October 2008. Accepted 25 January 2009.

We show that the cyanobacterial symbionts of a tripartite cyanolichen can produce hepatotoxic microcystins *in situ*. Microcystins were detected with high-performance liquid chromatography mass spectrometry both from cephalodia of the tripartite cyanolichen *Peltigera leucophlebia* and from a symbiotic *Nostoc* strain isolated from the same lichen specimen. Genetic identities of symbiotic *Nostoc* strains were studied by amplifying and sequencing the 16S rRNA gene. Also, the presence of the microcystin synthetase gene *mcyE* was confirmed by sequencing. Three highly toxic microcystins were detected from the lichen specimen. Several different *Nostoc* 16S rRNA haplotypes were present in the lichen sample but only one was found in the toxin-producing cultures. In culture, the toxin-producing *Nostoc* strain produced a total of 19 different microcystin variants. In phylogenetic analysis, this cyanobacterium and related strains from the lichen thallus grouped together with a previously known microcystin-producing *Nostoc* strain and other strains previously isolated from the symbiotic thalloid bryophyte *Blasia pusilla*. Our finding is the first direct evidence of *in situ* production of microcystins in lichens or plant–cyanobacterial symbioses. Microcystins may explain why cyanolichens and symbiotic bryophytes are not among the preferred food sources of most animal grazers.

Microcystins are a family of cyclic peptide hepatotoxins produced by a number of cyanobacterial genera (Sivonen and Jones 1999). They are potent inhibitors of eukaryotic protein phosphatases 1 and 2A (MacKintosh et al. 1990). Microcystins are implicated in the toxicosis of wild and domestic animals throughout the world (Ressom et al. 1994). These toxins are transported into the liver cells via the bile acid route and lead to overphosphorylation and disruption of the cytoskeleton in hepatocytes (Carmichael 1994). Subsequent hepatocyte necrosis leads to pooling of blood in the liver. In acute poisoning, death is the result of intrahepatic hemorrhage and hypovolemic shock. Microcystins are also suspected to act as tumor promoters (Nishiwaki-Matsushima et al. 1992) and the use of water contaminated with the toxin in renal dialysis is held responsible for the deaths of 60 patients in Brazil (Jochimsen et al. 1998).

Microcystins have a specific chemical structure and contain a number of nonproteinogenic amino acids (Sivonen and Börner 2008; Sivonen and Jones 1999). There are at least 85 known microcystin variants having the common structure of cyclo(D-Ala¹-X²-D-MeAsp³-Z⁴-Adda⁵-D-Glu⁶-Mdha⁷), where X and Z are variable L-amino acids, D-MeAsp is D-erythro-β-methylaspartic acid, Adda is (2S,3S,8S,9S)-3-amino-9-methoxy-2,6,8-trimethyl-10-phenyldeca-4,6-dienoic acid, and Mdha is N-methyldehydroalanine (Sivonen and Börner 2008). All of the amino acids have been reported to vary but, in addition to the variable amino acids in positions 2 and 4, the most common substitutions involve demethylation of the amino acids in positions 3 and 7. Also, the peculiar Adda structure may be demethylated into DMAAdda or ADMAdda, which has an acetoxyl instead of the methoxyl group (Namikoshi et al. 1992; Sivonen and Jones 1999). Microcystins are synthesized on large, mixed nonribosomal and polyketide synthetases in a programmed biosynthetic event (Christiansen et al. 2003; Nishizawa et al. 1999, 2000; Rouhiainen et al. 2004; Tillett et al. 2000). The *mcyE* gene was chosen for the genetic analysis because it is involved in the synthesis of Adda and the formation of the bond between Adda and D-glutamate, which vary less than other amino acids in the molecule and are essential for toxicity. The *mcyE* gene has also been shown to be unaffected by horizontal gene transfer (Rantala et al. 2004; Tillett et al. 2000).

Microcystins are produced by many different cyanobacteria, and the genera *Anabaena*, *Hapalosiphon*, *Microcystis*, *Nostoc*, and *Planktothrix* all contain microcystin-producing strains. A toxic strain usually produces several microcystin variants simultaneously but only one or two of these variants are dominant. Microcystin-producing strains are relatively rare among cyanobacteria and typically, even within a single species, some but not all strains actually produce microcystins. Environmental conditions, such as temperature and light, affect microcystin production. Microcystins are chemically very stable but some bacteria in aquatic environments have an ability to break them down (Sivonen and Jones 1999; van Apeldoorn et al. 2007).

It is estimated that 13% of all lichen species have cyanobacterial symbionts and the most common cyanobacterial genus in lichen symbioses is *Nostoc* (Rikkinen 2002). Symbiotic *Nostoc* strains can also establish well-defined symbiotic associations with the glomeromycete *Geosiphon* (Kluge et al. 2002), some thalloid bryophytes (Rikkinen and Virtanen 2008), cycads (Costa et al. 2004), and the angiosperm *Gunnera* (Bergman 2002).

In a previous study, a lichen-associated *Nostoc* strain was found to produce microcystins in culture (Oksanen et al. 2004). The toxin-producing cyanobacterium was isolated

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Nucleotide sequence data is available in the National Center for Biotechnology Information GenBank database.

*The e-Xtra logo stands for “electronic extra” and indicates that a supplementary table and supplementary material about microcystins in *Peltigera leucophlebia* are published online.

from a thallus fragment of the bipartite cyanolichen *Protopannaria pezizoides* from central Finland. However, the ecology of this cyanobacterium remained somewhat uncertain, because the identity of the lichen symbiont was not confirmed by sequencing the intact lichen. Here, we report the results of further studies on microcystin-production in lichen-associated cyanobacteria and focus on the cephalodial symbionts of the tripartite cyanolichen *Peltigera leucophlebia* (Nyl.) Gyeln. (Fig. 1).

RESULTS

Genetic diversity of *Nostoc* cyanobionts of *P. leucophlebia*.

We amplified the 16S rRNA gene from genomic DNA extracted from a pooled sample (approximately 200 in total) of *P. leucophlebia* cephalodia and constructed a clone library. All the cephalodia were collected from the upper surface of a single lichen thallus (size approximately 15 cm²). The 39 clones sequenced represented 29 different *Nostoc* 16S rRNA gene haplotypes (National Center for Biotechnology Information [NCBI] GenBank accession number FJ815293-321). In our analysis of the 29 16S rRNA sequences and other related sequences obtained from NCBI GenBank neighbor-joining, maximum likelihood and maximum parsimony produced phylogenetic trees with only minor differences. Most *Nostoc* hap-

lotypes from the cephalodia were placed into three closely related groups among a diversity of sequences previously obtained from symbiotic *Nostoc* strains of different *Peltigera* spp. and other terricolous cyanolichens, but also from the thalloid bryophytes *Blasia* and *Anthoceros*, various *Gunnera* spp., and the cycad *Macrozamia* (Fig. 2).

Single-strain cultures of symbiotic cyanobacteria were obtained from nine different cephalodia of *P. leucophlebia*. All the cultured cyanobacteria were filamentous, heterocystous, and showed no evidence of branching. They formed slowly spreading, gelatinous colonies typical of *Nostoc* spp. Sequencing the 16S rRNA gene revealed that all six analyzed cultures belonged to the same *Nostoc* haplotype (UK18), which had also been the most common haplotype among the sequences obtained directly from lichen cephalodia. The most similar sequence in NCBI GenBank was that of *Nostoc* sp. strain IO-102-I, a microcystin-producing cyanobacterium that had been found to be associated with the bipartite cyanolichen *Protopannaria pezizoides* in Sysmä, central Finland. Also, several other *Nostoc* haplotypes amplified directly from *Peltigera leucophlebia* cephalodia were nearly identical to these two strains (Fig. 2).

The *mcyE* microcystin synthetase gene fragment was amplified directly from the pooled sample of *P. leucophlebia* cephalodia (NCBI GenBank accession number FJ815292). The same

Fig. 1. Toxin-producing cyanolichen. **A**, *Peltigera leucophlebia* in natural habitat (scale bar 1 cm). **B**, Cephalodia (fungal structures containing symbiotic cyanobacteria) on the upper surface of the lichen thallus (scale bar: 1 mm). The white spots on the lichen surface are pine pollen. **C**, Isolated microcystin-producing *Nostoc* sp. strain UK18 in pure culture (scale bar: 20 µm).

Fig. 2. Phylogenetic relationships among symbiotic *Nostoc* strains (maximum-likelihood tree of 16S rRNA sequences). Symbiotic strains are named after their host species. Known microcystin producers are on black and the sequences from our clone library of *Peltigera leucophlebia* cephalodia are on gray backgrounds. Bootstrap values >50% are given at the node for maximum likelihood, maximum parsimony, and neighbor-joining, respectively.

mcyE gene haplotype was detected in all cloned samples ($n = 16$). The length of these gene fragments was 769 bp. They were nearly identical to the sequence previously obtained from *Nostoc* sp. strain IO-102-I (single difference at base number 183).

Microcystin production by the *Nostoc* cyanobionts.

We detected three highly toxic microcystins from a pooled sample of cephalodia collected from the surface of a *P. leucophlebia* thallus (Table 1; Fig. 1B). The major microcystin [ADMAdda⁵]MC-LR had a mass of 1,022 Da and accounted for approximately 60% of the total intensity of the microcystin peaks. Two other microcystins, [DMAdda⁵]MC-LR and [D-Asp³, ADMAdda⁵]MC-LR, each accounted for 20% of the total intensity and had a mass of 980 and 1,008 Da, respectively.

Strains were isolated and cultured from nine separate cephalodia of *P. leucophlebia* and all of them produced the same 19 microcystin variants with practically the same abundance (Table 1). The total amount of microcystin produced by the strains was approximately 0.4 µg/mg of dry weight. In addition to the microcystins detected from the *P. leucophlebia* cephalodia, several other previously known variants and new microcystin structures were found. The four most intensive microcystin peaks in the chromatograms obtained from the cultured *Nostoc* strains were [ADMAdda⁵]MC-LR (m/z 1,023, [M+H]⁺), [ADMAdda⁵]MC-(H₄)YR (m/z 1,077, [M+H]⁺), [D-Asp³, ADMAdda⁵]MC-LR (m/z 1,009, [M+H]⁺), and [ADMAdda⁵]MC-HilR (m/z 1,037, [M+H]⁺), and they accounted for approximately 70, 15, <10, and <10% of the total intensity, respectively. Rare nonproteinogenic amino acids such as tetrahydrotyrosine (H₄)Y and homoisoleucine Hil are proposed in second position on the grounds of mass spectrometric (MS) analysis.

The assignment of the main ions of the microcystins detected from *P. leucophlebia* cephalodia and in the four major variants of the cultured *Nostoc* strains is presented in Table 2. In the case of [ADMAdda⁵]MC-LR, the important ADMAdda diagnostic ion m/z 627 [Arg-ADMAdda-Glu + H⁺] and ion m/z 553 [Mdha-Ala-Leu-MeAsp-Arg + H⁺] are present (Table 2). Also, the DMAdda structure diagnostic ion m/z 585 [Arg-DMAdda-Glu + H⁺] and the ion m/z 553 can be seen in the spectrum of [DMAdda⁵]MC-LR. In both spectra, [D-Asp³,

ADMAdda⁵]MC-LR ions m/z 627 and m/z 539 [Mdha-Ala-Leu-Asp-Arg + H⁺] were intensive.

DISCUSSION

Three microcystin variants were detected from the cephalodia of the tripartite cyanolichen *P. leucophlebia*. This finding proves that some lichen-symbiotic *Nostoc* strains do produce microcystins in situ.

We found a considerable diversity of different 16S rRNA gene haplotypes in our pooled sample of *P. leucophlebia* cephalodia. Generally, lichen-forming fungi are believed to be quite selective in their choice of cyanobacterial symbionts and, typically, only a limited number of closely related *Nostoc* strains are found from the thalli of an individual lichen species (Myllys et al. 2007; Paulsrud and Lindblad 1998; Paulsrud et al. 2000). Previous studies have shown that this is also true of some tripartite *Peltigera* spp., such as *P. aphthosa* (Paulsrud et al. 2001), but more variation has been detected in other species, such as *P. venosa* (Paulsrud et al. 2000) and *P. leucophlebia* (Myllys et al. 2007). The diversity of *Nostoc* haplotypes found in the present study may have been partly related to the sampling method used. In order to obtain sufficient biomass for chemical analysis, hundreds of minute cephalodia had to be collected from the lichen surface and pooled into one sample. All the detected haplotypes belonged to a rather diverse group of symbiotic *Nostoc* strains typical of the *Peltigera* guild (Rikkinen et al. 2002). In our trees, the only major group gaining strong bootstrap support was that of GenBank sequences of *Nostoc* symbionts of lichens that belong to the *Nephroma* guild. This pattern is in good accordance with previous results (Myllys et al. 2007; O'Brien et al. 2005; Rikkinen et al. 2002).

Although no cyanobacteria were seen on cephalodial surfaces in microscopic examination, it is quite possible that some *Nostoc* haplotypes in the pooled sample were surface contaminants. However, a few incidental cells from outside the cephalodia could not have produced microcystins in detectable amounts. Also, all nine strains cultured from the *P. leucophlebia* cephalodia produced microcystins and belonged to the same haplotype that was most common among the sequences obtained directly

Table 1. Microcystin variants detected from the tripartite cyanolichen *Peltigera leucophlebia* and its cultured symbiont (*Nostoc* sp. strain UK18)^a

Microcystin variant	m/z [M+H] ⁺	t_R (min)	Cep. variant (%) ^b	UK18 variant (%) ^c
[DMAdda ⁵]mc-(H ₄)YR	1,035	28.5	...	<0.5
Microcystin	967	33.3	...	<0.5
[D-Asp ³ , DMAdda ⁵]mc-LR	967	33.8	...	<0.5
[DMAdda⁵]mc-LR	981	34.2	20	<2
[DMAdda ⁵]mc-FR	1,015	35.3	...	<0.5
[D-Asp ³ , ADMAdda ⁵]mc-(H ₄)YR	1,063	37.0	...	<2
[DMAdda ⁵]mc-HilR	995	37.6	...	<0.5
[ADMAdda ⁵]mc-(H ₄)YRHar	1,091	38.0	...	<0.5
[ADMAdda ⁵]mc-(H ₄)YR	1,077	38.1	...	15
[ADMAdda ⁵]mc-YR	1,073	38.9	...	<0.5
Microcystin	1,009	40.4	...	<0.5
[ADMAdda ⁵]mc-LR	1,023	40.9	...	<2
[D-Asp³, ADMAdda⁵]mc-LR	1,009	41.5	20	<10
[ADMAdda⁵]mc-LR*	1,023	41.7	60	70
[ADMAdda ⁵]mc-FR	1,057	42.3	...	<2
[ADMAdda ⁵ , Dha ⁷]mc-LR	1,009	42.7	...	<2
[ADMAdda ⁵]mc-HilR	1,037	44.0	...	<10
[ADMAdda ⁵]mc-HilR	1,037	45.2	...	<2
Microcystin	1,051	48.3	...	<0.5

^a Protonated molecular ion [M+H]⁺, retention time t_R , and the relative proportions of the variants detected by liquid chromatography mass spectrometry (MS) are given. The three microcystins detected from both the lichen sample and *Nostoc* sp. UK18 are in boldface. Structural differences were not detected in MS analysis in two pairs of microcystin isomers (m/z 1,023 and 1,037 Da) and * indicates the known late-eluting isomer. Three microcystin structures remained unresolved.

^b Relative proportion of the microcystin variant in *Peltigera leucophlebia* cephalodia.

^c Relative proportion of the microcystin variant in the *Nostoc* sp. strain UK18.

from lichen cephalodia. Hence, there is strong reason to believe that the toxin producers were the true cyanobiont of *P. leucophlebia*.

The microcystin-producing *Nostoc* sp. IO-102-I was originally isolated from a Finnish specimen of the bipartite cyanolichen *Protopannaria pezizoides*. However, because the identity of the cyanobacterial symbiont or microcystin production was not confirmed by analyzing the intact lichen, the exact ecology of *Nostoc* sp. IO-102-I remained unclear (Oksanen et al. 2004). Because 16S rRNA sequences now show *Nostoc* sp. UK18 to be the closest known relative of *Nostoc* sp. IO-102-I, it is almost certain that the later strain is also a lichen symbiont. Both strains were isolated from lichens in Finland but from locations more than 500 km apart. The lichen genus *Peltigera* is quite widespread on most continents (Martinez et al. 2003) and the cyanobacterial genus *Nostoc* is cosmopolitan (Wright et al. 2001). All this suggests that microcystin-producing lichen cyanobionts will not turn out to be only a local phenomenon.

Reindeer are ruminants that selectively forage according to digestibility of their food plants and fungi (Danell et al. 1994; Hofmann 1989). Lichen species are important winter feed for reindeer and caribou and, for example, in Northern Finland, *Cladonia* spp. with green algal symbionts are heavily grazed (Danell et al. 1994; den Herder et al. 2003). The nitrogen content of lichens with only green algae as a photobiont is so low that reindeer fed on a lichen diet composed mainly of *Cladonia* tend to lose weight (Storeheier et al. 2002). Nitrogen content of cyanolichens is substantially higher (Rai 2002), which would indicate a better nutritional value compared with green algal lichens. However, reindeer distinctly avoid eating cyanolichens even during starvation and the only previously proposed explanation for this phenomenon is the apparently poor digestibility of some cyanolichen species (Hofmann 1989; Storeheier

et al. 2002). The microcystin concentration of *Nostoc* UK18 was 0.4 µg/mg (dry weight), which is slightly more than the reported microcystin production of the strain IO-102-I (Oksanen et al. 2004). We estimate that, in the *Peltigera leucophlebia* thalli studied, the cephalodial *Nostoc* cyanobionts accounted for approximately 1% of the cyanolichen dry weight. For mature reindeer feeding on reindeer lichen, the intake is approximately 60 g/kg of dry lichen per body weight in a day (Holleman et al. 1979). Hence, a reindeer grazing solely on *P. leucophlebia* could theoretically accumulate microcystins at approximately 240 µg/kg in a day, while the acute toxicity value LD₅₀ of [ADMAAdda⁵]microcystin-LR is approximately 60 µg/kg of body weight in mice (Sivonen and Jones 1999). The microcystin content could be significantly higher in the case of bipartite cyanolichens like *Protopannaria pezizoides* (Rikkinen 2002). In heteromorous bipartite lichens, the proportion of photobiont cells is typically 5 to 15% (Honegger 1991). Whether the distribution of toxic cyanobacteria among lichen symbionts and the levels of microcystins in lichens could pose a risk to reindeer under some conditions should be studied further.

Nostoc is the most common cyanobacterial genus in lichen symbioses (Rikkinen 2002). In addition to lichen-forming fungi, symbiotic *Nostoc* strains also establish well-defined associations with several types of plants, including some thalloid liverworts and all hornworts, cycads, and species of *Gunnera* (Bergman 2002; Costa et al. 2004; Rikkinen and Virtanen 2008). The *Nostoc* symbionts of thalloid liverworts are closely related to those found in many terricolous lichens. In fact, sequence-identical *Nostoc* haplotypes have been identified from the liverwort *Blasia pusilla* (Blasiales) and several *Peltigera* spp. (Rikkinen and Virtanen 2008). Also, in the present study, both toxin-producing lichen symbionts grouped together with several *Nostoc* strains from *B. pusilla*—pointing toward the

Table 2. Assignment of the main ions in microcystins detected from *Peltigera leucophlebia* cephalodia and in the four major microcystin variants of the cultured cyanobionts in liquid chromatography mass spectrometry analysis

Fragment of [M + H] ^{±b}	Microcystin ^a						
	<i>P. leucophlebia</i> cephalodia			<i>Nostoc</i> sp. strain UK18			
	1	2	3	4	2	3	5 ^c
M + H	981	1,009	1,023	1,077	1,009	1,023	1,037
M + H – H ₂ O	963	991	1,005	1,059	991	1,005	1,019
M + H – COOH	936	964	978	1,032	964	978	992
Ala-X-(Me)Asp-Arg-(A)DMAdda-Glu-Mdha – CO + H	953	981	995	1,049	981	995	1,009
Ala-X-(Me)Asp-Arg-(A)DMAdda-Glu-Mdha – NH ₃ + H	964	992	1,006	1,060	992	1,006^c	1,020
Ala-X-(Me)Asp-Arg + H	470	456	470	524	456	470	484
X-(Me)Asp-Arg – NH ₃ + H	382	...	382	436	368	382	396
(Me)Asp-Arg-(A)DMAdda-Glu + H	714	742	756	756	742	756	756
(Me)Asp-Arg-(A)DMAdda-Glu – NH ₃ + H	697	725	739	739	725	739	739
Arg-(A)DMAdda-Glu-Mdha-Ala-X + H	852	894	894	948	894	894	908
Arg-(A)DMAdda-Glu-Mdha-Ala-X + NH ₂ + 2H	869	911	911	965	911	911	925
Arg-(A)DMAdda-Glu-Mdha-Ala-X – CO + H	824	866	866	920	866	866	880
Arg-(A)DMAdda-Glu-Mdha + H	668	710	710	710	710	710	710
Arg-(A)DMAdda-Glu + H	585	627	627	627	627	627^c	627
Arg-(A)DMAdda-Glu – CO + H	557	599	599	599	599	599	599
Arg-(A)DMAdda-Glu – NH ₃ + H	568	610	610	610	610	610	610
Arg-(A)DMAdda + H	456	498	498	498	498	498	498
Arg-(A)DMAdda – NH ₃ + H	439	481	481	481	481	481	481
Mdha-Ala-X-(Me)Asp-Arg-(A)DMAdda + H	852	880	894	948	880	894	908
Mdha-Ala-X-(Me)Asp-Arg-(A)DMAdda + NH ₂ + 2H	869	...	911	965	...	911	925
Mdha-Ala-X-(Me)Asp-Arg-(A)DMAdda – CO + H	824	...	866	920	852	866	880
Mdha-Ala-X-(Me)Asp-Arg + H	553	539	553	607	539	553	567
Mdha-Ala-X-(Me)Asp-Arg + NH ₂ + 2H	570	556	570	624	556	570	584
Mdha-Ala-X-(Me)Asp-Arg – CO + H	525	511	525	579	511	525	539

^a The most intensive peaks are in boldface and the base peaks of the spectra with a relative intensity of 100% are underlined. Column numbers indicate the following: 1 = [DMAdda⁵]mc-LR, 2 = [D-Asp³, ADMAAdda⁵]mc-LR, 3 = [ADMAAdda⁵]mc-LR, 4 = [ADMAAdda⁵]mc-(H₄)YR, and 5 = [ADMAAdda⁵]mc-HiLR.

^b X is the variable amino acid in position two: L, (H₄)Y, or HiL.

^c [ADMAAdda⁵]MC-LR had two base peaks with intensity of 100%.

possibility that microcystin-producing *Nostoc* strains could also occur in plant symbioses.

Nostoc-containing lichens are eaten and used in traditional medicine in China (Wang et al. 2001). Now that it is known that lichen-symbiotic *Nostoc* spp. do produce microcystins, it would be essential to screen for microcystins at least among the species that are consumed by humans. It has also been noticed that, even though certain *Adiantum* containing microcystins do strongly inhibit protein phosphatases, they show low cross-reactivity with antibodies used in the enzyme-linked immunosorbent assay, which is often used to test the toxicity of microcystins (Laub et al. 2002).

To conclude, we have now provided the first direct evidence of in situ production of microcystins in lichens. The toxin-producing lichen symbiont was very closely related to several bryophyte-associated *Nostoc* strains; therefore, one could expect to also find some toxin-producing strains among the cyanobacterial symbionts of *B. pusilla* and other thalloid bryophytes. Both plant and fungal hosts might benefit from the production of cyanobacterial microcystins because they are detrimental to animal grazers. However, the true significance of microcystin production for the producers themselves remains one interesting question to be explored in further studies.

MATERIALS AND METHODS

Symbiont culture.

Specimens of *Peltigera leucophlebia* were collected during week 40 in 2006 from the commune of Autti within the area of reindeer husbandry in northern Finland. Cyanobacterial symbionts were isolated from nine cephalodia and cultured on Z8 medium plates lacking nitrogen in continuous light at room temperature (Kotai 1972). In order to obtain clonal cultures, the cyanobacteria were replated repeatedly from single filaments and then grown in liquid Z8 medium to obtain sufficient biomass for liquid chromatography (LC)-MS analysis.

Microcystin extraction.

The upper surface of the lichen thallus was rinsed thoroughly with distilled water prior to collection of the cephalodia, after which cephalodia were scraped off. Cyanobacteria from the liquid cultures were harvested by centrifugation for 7 min at $7,400 \times g$ and 10 min at $10,600 \times g$. The cephalodia and the cultured cyanobacterial trichomes were freeze dried, homogenized with a mortar, and stored at -18°C . A sample of freeze-dried cephalodia (approximately 0.04 g dry weight) or cultured cyanobacteria was shaken in a Fast Prep (FP120; Bio 101, Savant, Thermo Electron Corporation, Milford, MA, U.S.A.) cell disrupter three times for 20 s with 1.0 ml of methanol and glass beads (0.5 mm) and then centrifuged at $10,600 \times g$ for 5 min. The supernatant was passed through a 0.2- μm -pore-size filter (GHP Acrodisc; PALL Life Sciences, New York). In the case of cultured strains, the samples were also sonicated twice in between of shakings and the supernatant was not filtered.

Freeze-dried microcystin-producing *Nostoc* sp. strains IO-102-I and 152 were also analyzed in addition to the samples of *P. leucophlebia* cephalodia and cephaloidal cyanobacteria in culture. Samples of all cultured cyanobacterial strains were prepared in the same way as the cephalodia sample.

Microcystin analysis.

Microcystins were identified by LC-MS/MS according to their microcystin characteristic protonated molecular ions $[\text{M}+\text{H}]^+$, fragment ions of the $[\text{M}+\text{H}]^+$, byproduct ion fragmentation (MS^3), and by comparing the results to reference strains *Nostoc* sp. 152 and IO-102-I (Oksanen et al. 2004; Sivonen et al. 1992). The extracts were analyzed (injection volume 10 μl)

with an Agilent 1100 Series LC/MSD Trap System high-performance liquid chromatograph (Agilent Technologies, Palo Alto, CA, U.S.A.), which has an XCT Plus model ion trap as a mass detector. The ionization method used was electrospray ionization (ESI) in positive mode. The column used in high-performance LC was Phenomenex Luna C18 (150 by 2.0 mm, 5 μm) (Phenomenex, Torrance, CA, U.S.A.). The mobile phase was composed of 0.1% formic acid in water (A) and 0.1% formic acid in 2-propanol (B). The gradient run was from 5% B to 100% B over 50 min at a flow rate of 0.15 ml min^{-1} at 40°C .

In ion source nebulizer gas (N_2) pressure was 207 kPa, desolvation gas flow rate 8 liters/min, and the desolvation temperature was 350°C . The capillary voltage was set to 3,270 V, the capillary exit offset was 152.4 V, the skimmer potential was 40.0 V, and the trap drive value was 106.1. Spectra were recorded 50 to 1,200 m/z and a scanning rate of 26,000 m/z s^{-1} .

The structure of certain microcystin product ions was confirmed by fragmentation. The MS^2 fragmentation amplitude was 0.5 V and MS^3 fragmentation amplitude was 1.0 V. The total microcystin concentration of the strain was approximated with a microcystin-LR standard (a gift from Z. Grzonka, Faculty of Chemistry, University of Gdansk, Poland).

DNA extraction, amplification, and sequencing.

DNA was extracted from the same biological material that had been used in the chemical analysis. Trace methanol was evaporated in a vacuum centrifuge and DNA was extracted by using the DNEasy plant mini kit (Qiagen GmbH, Hilden, Germany).

Amplification of cyanobacterial 16S rRNA genes from the lichen sample was performed with three primer pairs: 27F and 23S30R, 359F and 23S30R, and 359F and 781Ra/b (Bustil et al. 2002; Nübel et al. 1997). Amplification was performed in an 80- μl volume containing 22 ng of genomic DNA, 200 μM deoxynucleoside triphosphate (Finnzymes Espoo, Finland), 0.75 μM each primer, 0.1 U of DynaZyme EXT DNA polymerase (Finnzymes), and 2 μl of dimethyl sulfoxide. The heating cycle was as follows: the initial denaturation of 3 min at 94°C was followed by 35 cycles of 1 min at 94°C , 1 min at 54°C , and 2 min at 72°C , with a final extension of 10 min at 72°C . The 16S rRNA gene amplification from cultured cyanobacteria was performed in a 40- μl volume with the same reagent concentrations and conditions, but only with primers 27F and 23S30R and enzyme DynaZyme II DNA polymerase (Finnzymes).

The microcystin synthetase *mcyE* gene fragment was amplified and sequenced directly from *P. leucophlebia* cephalodia. The amplification was performed with 2 and 11 ng of extracted DNA with the same reagent concentrations as the 16S rRNA from the lichen sample but with primers *mcyE*-F2 and *mcyE*-R4 (Rantala et al. 2004). The heating cycle started with the initial denaturation of 3 min at 94°C and was followed by 35 cycles of 30 s at 94°C , 30 s at 52°C , and 1 min at 72°C , with a final extension of 10 min at 72°C . The *mcyE* gene fragment was sequenced with a Big Dye cycle sequencing kit (Applied Biosystems, Foster City, CA, U.S.A.). Then, 1 μl of amplified DNA, 5 $\times$ buffer, 1 μl of Big Dye, and 10 μM concentrations of the primers *mcyE*-F2, *mcyE*-R4, pEintF (5'-aatataactatcaagaacc-3'), and pEintR (5'-cctaatactccggtctaa-3') were used in a volume of 10 μl . The heat cycle consisted 25 cycles of 10 s at 96°C , 5 s at 50°C , and 4 min at 60°C . The products were purified by salt and ethanol precipitation and analyzed by an ABI Prism 310 DNA sequencer (Applied Biosystems).

The 16S rRNA gene amplification products and some products from the *mcyE* gene amplification were purified with gel electrophoresis and QIAQuick gel extraction kit (Qiagen).

Purified DNA was cloned with the TOPO TA cloning kit (Invitrogen, Groningen, The Netherlands). The clones amplified with primer pairs 27F-23S30R and 359F-23S30R were now also amplified with primers 359F and the cyanobacteria-specific 781Ra/b to determine whether the cloned DNA was cyanobacterial 16S rDNA. Clones with cyanobacterial DNA were then amplified with M13F and M13R (Invitrogen user manual) with 4 µl of bacterial colony resuspended in Luria-Bertani medium, with the same reagent concentrations and conditions as the genes from the cultured cyanobacteria.

Amplification products were purified with the Montage polymerase chain reaction clean-up kit (Millipore, Billerica, MA, U.S.A.) and Bio 101 Systems Geneclean Turbokit (MP Biomedicals Europe, Illkirch, France). The *mcyE* gene fragment was sequenced with primers M13F and M13R and 16S rRNA gene with primers 27F, pEF, pDF, and pER (Edwards et al. 1989). Sequencing was performed by MacroGen Inc. in South Korea. The chromatograms of *mcyE* and 16S rRNA gene sequences were checked and edited with the Chromas 2.2 program (Technelysium Pty, Tewantin, Australia) and the contig assembly and the alignment of the sequences were performed with Bioedit sequence alignment editor.

Phylogenetic DNA sequence analyses.

The new 16S rRNA gene sequences and related cyanobacterial sequences from GenBank at the NCBI were analyzed with phylogenetic methods. Additional sequences used to construct phylogenetic trees were searched with BLAST through the website of the NCBI and chosen on the grounds of overall similarity and in order to attain representative sampling from ecologically different *Nostoc* strains.

The length of the sequences used in the phylogenetic analyses was 1,361 bp. Sequences were aligned with ClustalW in the program BioEdit. The alignment was also edited manually. Phylogenetic analyses were conducted by using PAUP* (version 4) with 1,380-character-long aligned sequences of the 16S rRNA genes (Swofford 1998). Phylogenetic trees were inferred by using neighbor-joining, maximum parsimony, and maximum likelihood. To test the stability of monophyletic groups, 1,000 bootstrap replicates were analyzed in neighbor-joining and maximum parsimony and 100 replicates in maximum likelihood. In distance analyses, the neighbor-joining method used maximum-likelihood distances. Maximum-likelihood and maximum-parsimony analyses were performed with 10 heuristic searches, random addition sequence starting trees, and tree bisection and reconnection branch arrangements, with the number of rearrangements limited to 10,000. However, in maximum likelihood, bootstrap analyses were performed without the 10 random sequence addition replicates. The GTR model of DNA substitution with a γ distribution of rates and constant sites removed in proportion to base frequencies was used in maximum-likelihood analyses. Phylogenetic trees were rooted using *Azolla filiculoides* cyanobiont (AY742450).

ACKNOWLEDGMENTS

The work was supported by the grants from the Academy of Finland to J. Rikkinen (122288), K. Sivonen (53305—Research Center of Excellence and 214457), and D. P. Fewer (1212943). We thank M. Wahlsten for valuable help in extracting microcystins and L. Saari for handling cultures.

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AUTHOR-RECOMMENDED INTERNET RESOURCE

Invitrogen TOPO TA Cloning User Manual, Version U:
tools.invitrogen.com/content/sfs/manuals/topota_man.pdf