Altered expression of two light-dependent genes in a microcystin-lacking mutant of Microcystis aeruginosa PCC 7806

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Microcystin is a potent inhibitor of eukaryotic protein phosphatases and has been implicated in causing hepatotoxicity to humans and animals worldwide. It is produced primarily by the bloom-forming cyanobacterium Microcystis aeruginosa, although the function of the peptide in this micro-organism is unknown. In this study, a microcystin-related protein, MrpA, was identified using a microcystin-lacking mutant of M. aeruginosa, PCC 7806. Comparative two-dimensional protein electrophoresis showed that MrpA was strongly expressed in wild-type PCC 7806, but was not detectable in the mcyB mutant. MrpA showed similarity to the RhiA protein from Rhizobium leguminosarum, which is encoded by the rhiABC operon and controlled by quorum-sensing mediators. Sequencing of mrpA flanking regions in M. aeruginosa PCC 7806 revealed the presence of a rhiB homologue, mrpB, directly downstream of mrpA. Northern blot analyses of mrpA expression in cells exposed to different light conditions revealed a rapid decline of transcription under high light conditions. Most striking was a strong increase in transcript levels from cultures irradiated with blue light. The mrpA transcription level was strongly reduced in two independent microcystin-lacking mutants under all light conditions investigated.

Keywords: RhiA, transcription, endotoxin, Rhizobium leguminosarum

INTRODUCTION

Cyanobacteria (blue-green algae) are a particularly rich source of bioactive secondary metabolites, including peptides, polyketides and alkaloids. Under favourable environmental conditions these algae grow to such an extent that blooms develop. The toxic metabolites produced by these blooms have been implicated in serious health problems, and even death in humans and animals (Carmichael, 1994; Jochimsen et al., 1998; Namikoshi & Rinehart, 1996). Among the secondary metabolites synthesized by bloom-forming cyanobacteria the potent hepatotoxin microcystin has been most extensively investigated. More than 60 isoforms of this heptapeptide are currently known, sharing the structure cyclo(-d-Ala-L-X-d-MeAsp-L-Z-Adda-d-Glu-Mdha), where X and Z represent variable L-amino acids. Microcystins are specific inhibitors of the eukaryotic protein phosphatases 1 and 2A (Runnegar et al., 1993), but not of a similar cyanobacterial protein phosphatase (Shi et al., 1999). They are synthesized by various cyanobacterial freshwater species belonging to the genera Microcystis, Anabaena, Oscillatoria and Nostoc (Chorus & Bartram, 1999). Microcystis aeruginosa, which is the most widely distributed microcystin-producing species, also synthesizes other peptides, including the depsipeptides cyanopeptolin and micropeptin, the tricyclic microviridins, and the linear microginins and aeruginosins. Most of these peptides are protease inhibitors (Namikoshi & Rinehart, 1996), and the majority of Microcystis strains naturally synthesize more than one of these compounds.

Abbreviations: Adda, (2S,3S,8S,9S)-3-amino-9-methoxy-2,6,8,-trimethyl-10-phenyl-4,6-decadienoic acid; MALDI-TOF, matrix-assisted laser desorption/ionization-time of flight spectroscopy; Mdha, N-methyl-dehydroalanine; TFA, trifluoroacetic acid.

The GenBank accession number for the sequence reported in this paper is AJ271976.
The role of microcystin within the producing organism has not been elucidated. However, as its function is likely to be related to the conditions that enhance the production of peptides, factors such as culture age, temperature, light, nutrient, salinity, pH and micro-nutrient concentrations have been investigated and shown to affect the microcystin content of *M. aeruginosa*, Anabaena flos-aquae and Oscillatoria agardhii (Chorus & Bartram, 1999). A correlation between microcystin production and growth rate, regardless of environmental influences, has also been reported (Orr & Jones, 1998). However, with the inability to compare results from various studies due to different culturing and analysis methods used, no conclusions can be made about the effect of growth rate or environmental factors regulating toxin production. More recently, molecular techniques have been employed to show an effect of light on microcystin biogenesis in two strains of *M. aeruginosa* (Nishizawa et al., 1999; Kaebernick et al., 2000).

So far detailed investigations into the function of microcystin and other similar cyanobacterial secondary metabolites have been hampered by the lack of clearly defined mutants incapable of producing one of these substances. *Microcystis* spp. may produce a number of secondary metabolites but biosynthesis genes have only been identified for microcystin. The toxin microcystin is synthesized non-ribosomally via a giant multifunctional enzyme complex (microcystin synthetase), which includes peptide synthetases, polyketide synthases and modifying activities (Dittmann et al., 1997; Nishizawa et al., 1999, 2000; Tillett et al., 2000). Three of the genes (*mcyA*, *mcyB* and *mcyD*) involved in microcystin synthetase production in *M. aeruginosa* PCC 7806 have been insertionally inactivated and the respective mutants are characterized by the complete absence of all variants of microcystin. The mutants lack the whole microcystin synthetase complex, but still produce other non-ribosomal peptides (Dittmann et al., 1997; Tillett et al., 2000). These mutants are ideal subjects for studying the function(s) of microcystins.

Putative roles for microcystin have been suggested (Chorus & Bartram, 1999; Rohrback et al., 1999): as a feeding deterrent against zooplankton, a suppressor of the growth of competing species, or as an iron-scavenging molecule. Most of these functions, and those investigated for secondary metabolites in other bacteria and fungi, are usually linked with export of the substance from the cells under certain environmental conditions or growth states (Perry et al., 1999; Pirkin et al., 1996). In contrast to most microbial secondary metabolites, microcysts are produced from early exponential phase to stationary phase and are regarded as endotoxins (Chorus & Bartram, 1999). According to most recent data, however, an ABC-transporter protein is encoded as part of the gene cluster for microcystin biosynthesis (Tillett et al., 2000). This protein may play a role in the active export of microcysts from the cells. So far, microcystin detected outside the cells in the culture medium, under high light, has largely been attributed to cell lysis (Rapala et al., 1997). Yet this may also suggest an export, and point to an extracellular role for microcystin under these conditions (Kaebernick et al., 2000; E. Dittmann & M. Kaebernick, unpublished data).

Here we describe a molecular approach to the elucidation of microcystin function. A protein of approximately 30 kDa was found to be strongly expressed in wild-type cells, but was not detectable in protein preparations from *mcyB* mutant cells. It is similar to the RhiA protein from *Rhizobium leguminosarum*. Expression of RhiA is cell-density-dependent and involves quorum-sensing mediators. Transcriptional analysis of the ‘Microcystis’ homologue indicates regulation by light.

METHODS

**Cyanobacterial strains and culturing.** The microcystin-producing strain *M. aeruginosa* PCC 7806 was kindly provided by J. Weckesser (Freiburg University, Freiburg i.Br., Germany) and Dr. Rippka (Institut Pasteur, Paris, France). Mutants of this strain unable to produce microcystins were obtained by natural transformation and insertion of a chloramphenicol-resistant cartridge into the peptide synthetase genes *mcyB* (Dittmann et al., 1997) and *mcyA* (Tillett et al., 2000). Wild-type and mutant cell lines were grown at 23 °C in batch cultures with BG-11 or Z8 medium (Rippka, 1998), respectively. The mutants used in this study were selected and kept on BG-11 medium containing 5 µg chloramphenicol ml⁻¹ (Dittmann et al., 1997). For comparative studies of the two variants chloramphenicol was removed from the mutant medium in order to exclude any influence the antibiotic may have had on physiological processes. Growth of the wild-type and the mutant strains was followed by monitoring the OD₆₆₆ nm with a UVikon 930 spectrometer (Contron Instruments) and the cell number by microscopic counting using a haemocytometer (Thoma). Cells were grown under white light (low light), 5 µmol photons m⁻² s⁻¹ (very low light) or 68 µmol photons m⁻² s⁻¹ (high light). Light intensities were measured using a LI-COR LI-250 light meter (Walz). Different light qualities were obtained by the use of red and blue plastic cut-off filters (Lee Filters). The blue light filter had a bandwidth of 400–510 nm with a peak at 472 nm and the red filter opened from 620 nm with a bandwidth at 666 nm. For both cases a light intensity of 16 µmol photons m⁻² s⁻¹ was applied in a dark room using a Fibre Illuminator FL-440 (Walz) and unilateral irradiation.

**Two-dimensional gel electrophoresis.** This was performed as described by Otto et al. (1996) as modified by Scheler et al. (1998). Using the Bio-Rad 2-D system. Twenty milligrams of dried cells was suspended in sonication buffer [9 M urea, 70 mM DTT and 2 % ampholyte (Serva 2-4)] and sonicated for 5 min. After centrifugation (12000 g, 10 min) the first dimension (IF) was performed in a glass tube (0.15 cm x 9.3 cm) using 20 µg of the protein extract. For the second dimension the IF gel was incubated with loading buffer for 15 min and subsequently loaded onto a 10 % SDS-PAGE gel. Gels were stained with Coomassie R 250.

**Isolation of a two-dimensional spot and sequence analysis of peptides.** Protein was isolated from a spot as described by Otto et al. (1996) and Scheler et al. (1998). The protein was digested with trypsin (1 µg in 600 µl digestion buffer; 100 mM Tris/HC1 pH 8.5 in 50 % acetonitrile) overnight at 37 °C. The reaction was stopped with 2 % TFA (1 h, 60 °C) and the gel was mixed with reversed-phase material (Scheler et al., 1998).
After binding (0.1% TFA) and washing (0.1% TFA) peptides were eluted with elution buffer (60% acetonitrile in 0.1% TFA). Peptides were separated using a FPLC chromatography system (Pharmacia), analysed by MALDI-TOF (Perspective Biosystems) and sequenced using the Procise Protein Sequencing system (Applied Biosystems).

**Degenerate PCR and sequencing, partial genomic library.** The two primer pairs Deg1F/Deg2R (5’-GTNGTNGATHGAYAAYTTY-3’/5’-RTTRAARTCNRATARTA-3’) and Deg2F/Deg1R (5’-TAYTYATGNYATTTAYAYAY-3’/5’-RARTTGTCDATNACN-3’) were designed from the peptide fragments LIAQAANVVIDNFSSFDQ and TYYVDFNNQTDR. PCR was performed using Qiataq (Qiagen) and 45 °C as the annealing temperature. A number of bands from both amplifications were eluted from the gel and inserted into the pGem-T vector (Promega) according to the manufacturer’s instructions. Sequence reactions were carried out using a Dye-deoxy Terminator Cycle Sequencing Kit (Applied Biosystems) and analysed on an automated DNA sequencer (model 373; Applied Biosystems). Southern blotting and the construction and screening of a partial (XbaI) library in pUC19 were performed as described by Sambrook et al. (1989).

**RNA isolation and Northern blot hybridization.** For RNA isolations 30–50 ml cultures (mid-exponential phase; OD$_{600}$ 0.7–1.0) were immediately chilled under the particular light conditions being investigated. Cultures were centrifuged (2770 g, 10 min, 4 °C) and stored at −20 °C. The pellet was homogenized in liquid nitrogen using a pestle and mortar. RNAs were isolated using the Trizol kit (Gibco-BRL) according to the manufacturer’s instructions. RNA gel electrophoresis, Northern blotting and hybridization were performed as described by Sambrook et al. (1989). Probes to the psbB gene fragment of Synechocystis PCC 6803 were kindly provided by Dr Annetget Wilde (Humboldt University, Berlin, Germany). A gene fragment of the cpc operon of PCC 7806 was amplified using the primer pair PCf and PceR (Neilan et al., 1995) and the 825 bp mrpA PCR fragment cloned into the pGEM-T vector (see above) was used as a gene-specific reverse primer. The primer pair mdpBF (5’-CATACGGTTGGATGTTGTGC-3’) and mdpBR (5’-CATTCCCTCTGGTCCAATTGC-3’) was used for use as a probe was amplified using the primer pair 16S (5’-TGTAAAACGACGGCCAGTGAAGCTGTAACAAAGC-3’) and 16S (5’-TGCATGACCTGTGCGATGGA-3’). The reverse transcription was performed using an RT kit (Gibco-BRL) according to the manufacturer’s instructions. After the reaction RNA was further purified using DNase I (Gibco-BRL). The mrpBRT primer (5’-CAAGCTCTGAGCTGCTGATTGC-3’) was used as a gene-specific reverse primer. The subsequent PCR was performed with DNA, DNA-free RNA, as a positive and a negative control, respectively, and cDNA with the primer mrpBRT and the primer MB35/63F (5’-GAAGGTGACTCCAGTGTTGATG-3’).

**RESULTS**

**Identification of the differentially expressed MrpA protein**

In order to compare the proteomes of *M. aeruginosa* PCC 7806 and the microcystin-lacking mcyB mutant (Dittmann et al., 1997) we performed two-dimensional protein electrophoresis. Visual analysis revealed several qualitative and quantitative differences between the wild-type and mutant. A protein of approximately 30 kDa was strongly expressed in the wild-type extract, but was not detectable in the mutant extract (Fig 1a, b). At least three more proteins showed different abundances. Two (22 kDa, 25 kDa) were more strongly and one (40 kDa) was more weakly expressed in the mcyB mutant. The most prominent protein, of approximately 30 kDa, was extracted from the gel and trypsinized for further analysis. Tryptic peptides were purified and fractionated using MALDI-TOF and FPLC. Four of the resulting peptides were subsequently sequenced (Fig. 1c). A blast (http://www.ncbi.nlm.nih.gov/blast/blast.cgi) search of these peptides against the complete proteome of *Synechocystis* sp. PCC 6803 (Kaneko et al., 1995) revealed no significant homologies and a search against the entire *Synechocystis* database indicated that this was not a conserved protein among cyanobacteria.

The sequence information from peptides 1 and 2 (Fig. 1c) was used to design degenerate PCR primers, as these sequences had the lowest degree of redundancy in their amino acid composition. As the order of the peptides in the protein was unknown, two complementary primer pairs were synthesized. Both pairs yielded a number of PCR products (data not shown). Three abundant bands from each of the two PCR approaches were eluted from the agarose gel and inserted into the pGEM-T vector. Sequencing of a number of clones revealed an 825 bp PCR product encoding a protein starting with the sequence TYYVDFNQNTDR and terminating with the sequence LIAQAANVVIDNFSSFDQ. Further sequence analysis of the whole clone led to the identification of a continuous ORF encoding a protein with identity to all partial peptides we had sequenced before. The 825 bp sequence encodes a protein of approximately 30 kDa that represents the major part of the gene encoding the microcystin-related protein, MrpA. A blast search revealed 42% similarity and 23% identity of the N-terminus to RhiA from *R. leguminosarum*. Further homologues of the mrpA gene were identified in the complete genomes of *Yersinia pestis* and *Yersinia pseudotuberculosis* (http://pedant.gsf.de/).

To complete the sequence encoding MrpA, and to identify neighbouring genes, we established a partial genomic library of strain PCC 7806. Using Southern blot hybridization with the 825 bp PCR fragment as a probe we identified a single 3-kb *XbaI* fragment. Based on this information, *XbaI* fragments with a size range of 3–2–3 kb were cloned after insertion into a pre-digested pUC19/XbaI vector. Subsequently, a positive *Escherichia coli* clone containing a 3-kb fragment of *M. aeruginosa* DNA was identified by colony hybridization. The sequence was determined for both strands and has been deposited in the EMBL/GenBank database under the accession number AJ271976.

Analysis of the 3-kb DNA fragment revealed the presence of three ORFs, designated mrpA, mrpB and ORF3. The molecular mass of MrpA was determined to be 35.9 kDa (pI 5.11). The second ORF, *mrpB*, was
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70

kDa 4·5 7·0 pI 4·5 7·0 pI

(a) (b)

(c)

1. TYYVDFNNQTDK
2. LIAQAANVVDNFSFDQXA
3. FNIIQGKTKK
4. DVLSSSQAQFK

Fig. 1. Two-dimensional protein maps of (a) wild-type M. aeruginosa PCC 7806 and (b) the mcyB mutant (Dittmann et al., 1997). Protein extracts were isolated from late-exponential phase cultures. Positions of qualitative and quantitative differences are indicated. The white arrow denotes a protein spot of approximately 30 kDa which was predominantly expressed in the wild-type PCC 7806, but not detectable in the mcyB mutant. This protein has been designated MrpA (microcystin-related protein A). Black arrows indicate further differentially expressed proteins. (c) Partial peptide sequences derived from tryptic digestion of the MrpA protein.

located 117 bp downstream of mrpA. Its deduced amino acid sequence exhibits 71 % similarity and 56 % identity to RhiB. RhiB is encoded together with RhiA and a third protein, RhiC, in the rhiABC operon of R. leguminosarum. The molecular mass of MrpB was calculated to be 15·8 kDa with a pI of 4·47. A protein with a pI in this range was not detectable by two-dimensional gel electrophoresis under the conditions used in this study. The third ORF was located 275 bp downstream of mrpB on the same DNA strand. Database analysis of this ORF did not show significant similarity to known proteins. Analysis of MrpA and MrpB, using PROSITE and the PSORT software, detected no significant matches with motifs in the database for MrpA; however, a possible ATP/GTP binding motif (P-loop) was observed in MrpB. MrpA is predicted to be a cytosolic protein, as is its homologue in Rhizobium. Cytosolic localization was also predicted for MrpB, with a lower value compared to MrpA.

Transcript analyses

To investigate whether different amounts of MrpA in wild-type and mutant cells correlate with altered transcript levels we performed Northern blot hybridizations with a fragment of mrpA as the probe. High amounts of a transcript of approximately 1·5 kb were detected in RNA preparations from the wild-type, compared to the reduced level found in samples prepared from mutant cells (Fig. 2). As already suggested by its length, this transcript also hybridized with a mrpB probe, therefore indicating a bicistronic messenger (data not shown). The same RNA samples were used to compare the abundances of cpcBA and psbB transcripts, encoding proteins of the light-harvesting antennae and of the photosystem II reaction centre, respectively. For both transcripts, no differences between wild-type and mutant cells were observed. In all cases 16S rRNA served as an internal standard (Fig. 2). The co-transcription of mrpA and mrpB was confirmed by RT-PCR (Fig. 3). The intergenic region between mrpA and mrpB was successfully amplified from cDNA using a primer pair binding to mrpA and mrpB, respectively.

Fig. 2. Northern blot analysis of M. aeruginosa PCC 7806 (WT) and mcyB mutant (M) RNA isolated from late-exponential phase cultures under medium light condition (20 µg RNA per lane). DNA fragments of (a) cpcBA, (b) psbB, (c) mrpA and (d) M. aeruginosa 16S rRNA served as probes.

The rhiABC transcription in R. leguminosarum was shown to be under the control of quorum-sensing mediators, therefore showing a strong increase in the stationary phase (Rodelas et al., 1999). In contrast, the mrpAB transcript did not show significant differences at different time points of the growth curve (data not
shown). Genes of the microcystin biosynthesis gene cluster are differentially expressed under various light intensities and qualities (Kaebernick et al., 2000). With the possibility of an influence of microcystin on MrpAB production, we also studied the response of mrpAB transcripts to different light quantities and qualities. Wild-type PCC 7806 was cultured to mid-exponential phase (OD_{660} 0.66) under low light conditions (16 μmol photons m^{-2} s^{-1}). Subsequently, aliquots were exposed for 30 min or 2 h, to different quantities of white light (16 μmol photons m^{-2} s^{-1}, 68 μmol photons m^{-2} s^{-1}, 400 μmol photons m^{-2} s^{-1}), red light or blue light (each at 16 μmol photons m^{-2} s^{-1}) and complete darkness (Fig. 4a). RNA was isolated from each of the samples and analysed by Northern blot hybridization. Transcript amounts seen under low light (16 μmol photons m^{-2} s^{-1}) were almost abolished after 2 h of darkness, or exposure to high light and very high light (68 μmol photons m^{-2} s^{-1} and 400 μmol photons m^{-2} s^{-1}, respectively) (Fig. 4a). Most striking was the strong increase in the mrpAB transcript level in cells exposed to blue light for 2 h (Fig. 3a). Results were standardized by hybridization with a DNA probe encoding the 16S rRNA gene of the total RNA samples (Fig. 4a). The microcystin-free mcyB mutant, grown to mid-exponential phase and subsequently exposed to the same light conditions as above, revealed strongly reduced transcript amounts, even though a fourfold amount of RNA was used in Northern blot hybridizations. Interestingly, no significant differences in the transcript amounts were observed for the different light conditions (Fig. 4b). The same results were obtained for the mcyA mutant (Fig. 4c). An extracellular microcystin proportion can be detected, especially under high light (E. Dittmann & M. Kaebernick, unpublished data). In order to find out if this extracellular microcystin proportion could potentially influence the mrpAB transcription (as seen for the rhiABC transcription in Rhizobium, which is strongly dependent on AHL mediators), 300 ng microcystin ml^{-1} was added under low light and high light conditions to the mcyB mutant culture. However, extracellular microcystin could not restore the wild-type transcription level (data not shown). All experiments with the wild-type and mutant strains were repeated twice, using independent material; reproducible results were obtained.

**DISCUSSION**

There has been considerable interest in understanding the function of microcystin for bloom-forming species of cyanobacteria. The microcystin-lacking mcyB mutant opened a way for investigations into the role(s) such metabolites may play by enabling comparison of wild-type and mutant cells at the molecular level. This study describes the light-regulated mrpA and mrpB genes,
which were previously thought to be unique to Rhizobium, and shows that the expression of these genes, but not of other light-regulated genes, is drastically reduced in the mcyB mutant at both the protein (not shown for MrpB) and the mRNA level. We conclude that the down-regulation of mrpA and mcyB expression is directly or indirectly caused by the lack of microcystin in two independent mutants.

MrpA and MrpB show significant similarities to the RhiA and RhiB proteins of Rhizobium leguminosarum, respectively. While effects on nodulation, especially in the absence of nodFEL, have been observed in R. leguminosarum, the biochemical role of these proteins remains unclear (Gray et al., 1996). The genes rhiA and rhiB, together with rhiC, form an operon which is located on the sym plasmid and is flanked by nifH and the nod gene cluster (Cubo et al., 1992). Several plasmids of different sizes exist in M. aeruginosa (Schwabe et al., 1988; E. Dittmann & G. Christiansen, unpublished data). Although the localization of mrpAB has not been investigated it is known that the mcy genes encoding the microcystin synthetase reside on the chromosome (Tillett et al., 2000).

The RhiA protein is one of the most abundant cytoplasmic proteins in R. leguminosarum (Cubo et al., 1992). Similarly, we detected MrpA as one of the most predominant spots in the two-dimensional pattern of protein preparations from M. aeruginosa PCC 7806 wild-type cells. A rhiC homologue is missing in the direct vicinity of mrpA and mcyB. In contrast to RhiA and RhiB, a periplasmic localization is assumed for RhiC (Cubo et al., 1992). A rhiC homologue may well be positioned distinct from mrpAB within the Microcystis genome. Alternatively the function of RhiC is unnecessary in Microcystis.

Expression of RhiA, B and C is controlled by the RhiR protein, which is a LuxR-type regulator. The LuxR/LuxI type of quorum-sensing regulation has been shown to control the expression of an increasing number of genes, including peptide synthetase genes, exopolysaccharide genes and virulence factors, especially in host-associated bacteria such as Pseudomonas aeruginosa, Erwinia carotovora and Agrobacterium tumefaciens (Rhizobium tumefaciens) (Whiteley et al., 1999; Bassler, 1999). In Gram-negative bacteria quorum-sensing systems consist of an N-acylhomoserine lactone (AHL) signal molecule (autoinducer), which is synthesized by a LuxI homologue, and requires an autoinducer-dependent transcriptional activator protein, as represented by LuxR (Whiteley et al., 1999). At least two different types of AHLs were shown to regulate the rhiABC operon in a cell-density-dependent manner (Rodelas et al., 1999). This type of quorum sensing has not been described for cyanobacteria so far. However, in the natural environment cell-to-cell contact plays an important role for Microcystis and a type of quorum sensing may be required during the formation of a colony habit and of blooms. No evidence exists yet for a role of microcystin as an extracellular signalling molecule. However, the existence of a putative ABC transporter linked to microcystin, as inferred by the identification of a gene homologous to those of known ABC-transporters and located directly upstream of the mcy gene cluster (Tillett et al., 2000), supports the proposal of an active efflux mechanism for microcystin. As significant extracellular amounts of microcystin are detectable, especially under high light (Rapala et al., 1997; E. Dittmann & M. Kaebnerick, unpublished data), this ABC-transporter might be active under these conditions. Such conclusions can also be drawn from preliminary transcription data of the corresponding gene (E. Dittmann, unpublished data). The involvement of efflux pumps in quorum-sensing processes has been postulated not only for AHLs, although diffusible (Whiteley et al., 1999), but also for quinolone as an alternative quorum-sensing mediator (Pesci et al., 1999).

Especially interesting with regard to a function of microcystin is the influence of light on the mrpAB transcript. Presently, there are no data which indicate a role for light in the expression of rhiABC in R. leguminosarum. In PCC 7806 wild-type, the effect of light on mrpAB is seen by strong transcript increases when cells are exposed to blue light. The lack of microcystin in the mcyB mutant leads to a reduction in both the MrpA protein and the level of mrpAB transcripts observed under all light conditions tested. The mutation, however, had no influence on other light-regulated genes (cpcBA, psbB). The different responses of mrpAB transcript accumulation by mutant (no response) and wild-type cells (distinct increase) exposed to blue light indicate the microcystin-dependent nature of mrpAB transcription. Therefore, it appears that the presence of microcystin is a pre-condition for the blue light effect on mrpAB expression. In contrast to the well-investigated mechanisms sensing cell density in other bacteria, bloom-forming cyanobacteria may sense light intensity and/or quality and use this mechanism to control production and (possibly) export of microcystin. Microcystin in turn may act under high-light conditions as an intercellular signal. However, mrpAB is not directly regulated by extracellular microcystin, as seen for rhiABC regulated by AHLs. Since the addition of microcystin to the mutant cells did not restore the wild-type transcription level, there could either be another key factor which is regulated by microcystin, or the intracellular microcystin proportion may be somehow involved in the regulation of mrpAB; alternatively, microcystin reception could be affected due to the lack of microcystin in the mutants. We conclude that additional light-sensing mechanisms may exist in bloom-forming cyanobacteria, which share features with quorum-sensing mechanisms of other bacteria.

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