

## Structural Characterization of Microcystins by LC/MS/MS under Ion Trap Conditions

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*This article is dedicated in memory of Professor Kenneth L. Rinehart at the University of Illinois*

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**Abstract** LC/MS/MS under ion trap conditions was used to analyze microcystins produced by cyanobacteria. Tandem mass spectrometry using MS<sup>2</sup> was quite effective since ions arising from cleavage at a peptide bond provide useful sequence information. The fragmentation was confirmed by a shifting technique using structurally-related microcystins and the resulting fragmentation pattern was different from those determined by triple stage MS/MS and four sector MS/MS. Analysis of a mixture of microcystins in a bloom sample was successfully performed and two new microcystins were identified by LC/MS/MS under ion trap conditions. Thus, LC/MS/MS under ion trap conditions is effective for the structural characterization of microcystins.

**Keywords** LC/MS/MS under ion trap conditions, cyclic peptide, microcystin, structural analysis, sequence information, fragmentation pattern

### Introduction

Cyclic peptides have been often found and isolated from microorganisms, cyanobacteria, fungi and marine sponges as bioactive secondary metabolites, and some of them have been developed as medicines. For instance,

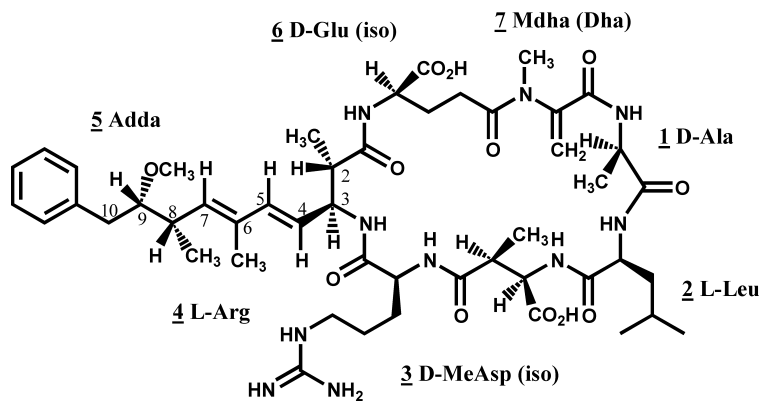
cyclic peptide antibiotics such as bacitracin [1], colistin [2], vancomycin [3] and micafungin [4], are widely used for medical treatments. Moreover, cyclosporin [5] is an immunosuppressive agent, which has been used for organ transplant rejection or bone marrow [6].

For the structure determination of cyclic peptides, the following information is required: structures, absolute configurations and sequence of constituent amino acids. Although partial acid hydrolysis had been used for the structure determination of these cyclic peptides, it has been currently performed by instrumental methods such as 2D-NMR (two dimensional nuclear magnetic resonance) and MS/MS (tandem mass spectrometry) techniques. However, these techniques have the following inherent drawbacks: overlapped signals derived from carbonyl carbons and  $\alpha$ -hydrogens of the constituent amino acids and non-reproducible fragmentation by tandem mass spectrometry with high- (four sector MS/MS) or low- (triple stage MS/MS) energetic CID (collision activation dissociation). Therefore, a more reliable and effective methodology has been eagerly required for cyclic peptides [7].

Microcystins (Fig. 1), a family of cyclic heptapeptides isolated from cyanobacteria, show potent inhibition of protein phosphatases 1 and 2A [8]. Their structures were determined in 1984 [9], and 67 related compounds have been isolated since then [10]. Although triple stage MS/MS

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| Microcystins (abbreviation)            | M.W. | 1     | 2            | 3            | 4            | 5    | 6     | 7            |
|--|------|-------|--------------|--------------|--------------|------|-------|--------------|
| Microcystin-LR (MCLR)                  | 994  | D-Ala | L-Leu        | D-MeAsp      | L-Arg        | Adda | L-Glu | Mdha         |
| Microcystin-YR (MCYR)                  | 1044 | D-Ala | <u>L-Tyr</u> | D-MeAsp      | L-Arg        | Adda | L-Glu | Mdha         |
| Microcystin-RR (MCRR)                  | 1037 | D-Ala | <u>L-Arg</u> | D-MeAsp      | L-Arg        | Adda | L-Glu | Mdha         |
| Microcystin-LF (MCLF)                  | 985  | D-Ala | L-Leu        | D-MeAsp      | <u>L-Phe</u> | Adda | L-Glu | Mdha         |
| Microcystin-LW (MCLW)                  | 1025 | D-Ala | L-Leu        | D-MeAsp      | <u>L-Trp</u> | Adda | L-Glu | Mdha         |
| 3-Desmethylmicrocystin-LR (3-Des-MCLR) | 980  | D-Ala | L-Leu        | <u>D-Asp</u> | L-Arg        | Adda | L-Glu | Mdha         |
| 7-Desmethylmicrocystin-LR (7-Des-MCLR) | 980  | D-Ala | L-Leu        | D-MeAsp      | L-Arg        | Adda | L-Glu | <u>Dha</u>   |
| Dihydromicrocystin-LR (Dihydro-MCLR)   | 996  | D-Ala | L-Leu        | D-MeAsp      | L-Arg        | Adda | L-Glu | <u>MeAla</u> |

Adda: [(2*S*,3*S*,8*S*,9*S*)-3 amino-9-methoxy-10-phenyl-2,6,8-trimethyldeca-4(*E*),6(*E*)-dienoic acid, Mdha: *N*-methyldehydroalanine, Dha: Dehydroalanine, MeAsp:  $\beta$ -erythro-methyl-D-Asp, MeAla: *N*-methylalanine.

**Fig. 1** Structures of microcystin LR (MCLR) and related compounds.

and four sector MS/MS were applied to characterize the microcystin structures [11~14], the resulting fragmentations were complicated and it was difficult to use them for the structural characterization of unknown microcystins. Recently, LC/MS/MS (liquid chromatography/tandem mass spectrometry) under ion trap conditions has been applied to the structure elucidation of various compounds with low to high molecular weights, and mainly to the amino acid sequence determination in the proteomics study and *in vivo* or *in vitro* metabolite screening during the process of drug development [15~17]. From LC/MS/MS under ion trap conditions, it is possible to acquire much information with a small amount of sample and then use the MS<sup>n</sup> technique. For the MS<sup>2</sup> and MS<sup>3</sup> analyses of microcystin using LC/MS/MS under ion trap conditions, the resulting fragmentation pattern was simpler than those with other MS/MS methods. Poon *et al.* reported good results for the analysis of microcystins using this system. However, the elucidation of the fragmentation for microcystin-LR (MCLR) was not enough to apply it to the structure characterization of unknown microcystins [18]. Moreover, although Deihnelt *et al.* analyzed various microcystins using ion trap Fourier transform ion cyclotron,

this equipment has not been widely used and not been a standard tool [10]. Therefore, we have made an attempt to apply LC/MS/MS under ion trap conditions to extend its applicability for the structure elucidation of unknown microcystins.

In the present study, we confirmed the fragmentation pattern of microcystins using LC/MS/MS under ion trap conditions, which was used for the structure characterization of non-isolated microcystins after simple extraction from a bloom sample [19].

## Materials and Methods

### Chemicals and Reagents

Microcystins, -LR, -YR, -RR, -LW, -LF, 3-desmethyl-LR and 7-desmethyl-LR were isolated and purified by silica-gel chromatography and gel filtration chromatography from the lyophilized cells of cyanobacteria collected in various sources or fermented strains [20]. Dihydromicrocystin-LR was prepared by the reduction of microcystin-LR with NaBH<sub>4</sub> [21]. These microcystins were then dissolved in methanol (10  $\mu$ g/ml). Each injection volume was 5  $\mu$ l for

the analysis. A TSK-gel Super ODS column, 2.0×100 mm (Tosoh Co. Ltd., Tokyo, Japan) was used for the fractionation of the microcystins. Methanol and formic acid were purchased from Nacalai tesque Co. Ltd. (Kyoto, Japan). Pure water was obtained using the E-pure distillation and ion exchange system (Barnstead, Germany). A 5% acetic acid solution was used for the extraction of the microcystins from lyophilized cells collected from Lake Suwa, Japan. The Sep-pak cartridge (ODS, 500 mg) used for the extraction of microcystin from the lyophilized cells was purchased from Waters Co., Ltd. (MA, USA).

### Instrumentation

The Agilent Technologies (Waldbronn, Germany) 1100 series was used as the HPLC system that consisted of binary pump, column compartment module, degasser, and UV-VIS detector. The LC/MS/MS system was a Thermo Electron LCQ Deca XP plus (San Jose, CA, USA) equipped with an electrospray ionization (ESI) interface.

### LC/MS Conditions

The chromatographic system consisted of an ODS column with the mobile phase of A: water and B: methanol containing 0.1% formic acid. The LC gradient elution conditions were initially 40% B to 90% B over 30 minutes. The flow rate was set at 0.2 ml/minute. The MS<sup>2</sup> analysis was used in the product ion scan in centroid mode. In the MS<sup>2</sup> mode, detection was in the positive ionization mode. The source voltage was 5.0 kV. The temperature of the heated capillary was 300°C. Full-scan mass spectra were acquired over  $m/z$  400~1500 within 1.0 second. Helium was used as the collision gas in the range of 1~2 mTorr. The protonated molecule was activated at 35% different collision energy level.

### Extraction of Microcystins

Extraction was carried out using the established method as follows [19]: a 100 mg sample of lyophilized cells of cyanobacteria collected from Lake Suwa was extracted with 10 ml of a 5% acetic acid solution for 30 minutes, and the resulting extract was centrifuged at 3000 *g* for 10 minutes. This extraction was repeated 3 times. After the filtration, the supernatant was applied to an ODS cartridge, and the cartridge was washed with 5 ml of water, followed by 5 ml of 20% methanol. The desired compounds were eluted with 5 ml of 90% methanol and then 5 ml of methanol. Both fractions containing the microcystins were collected and evaporated to dryness, and the residue was dissolved in methanol.

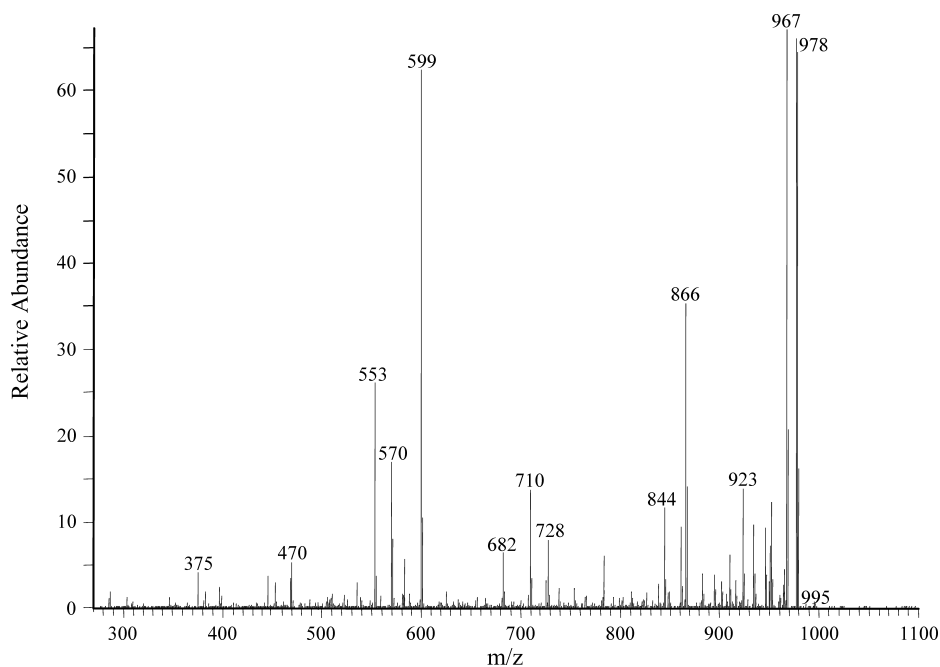
## Results

### Fragmentation Pattern of Arg Containing Microcystin

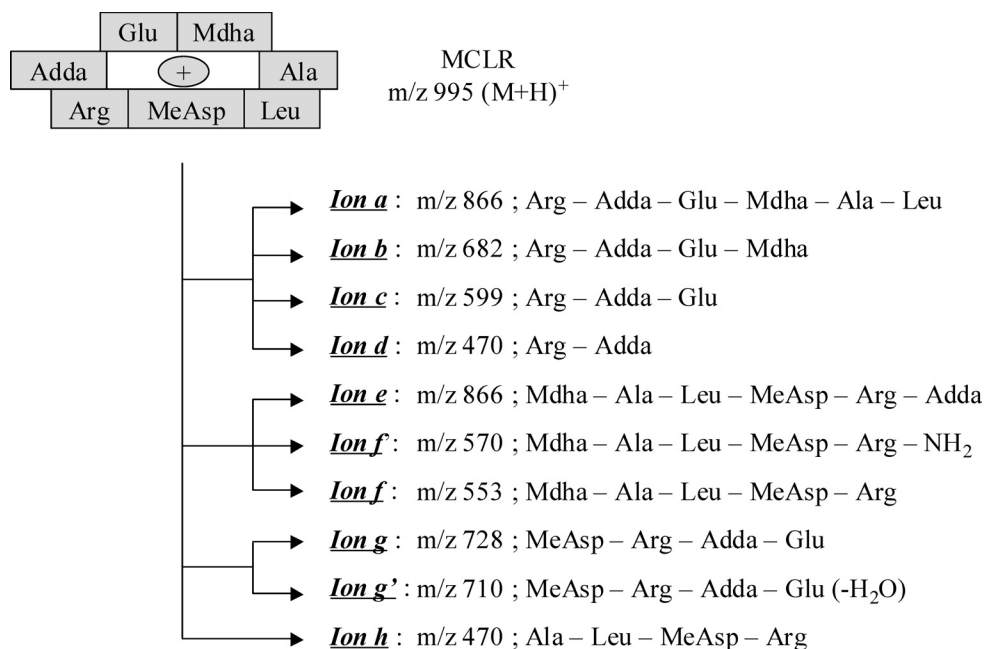
The general structure of microcystin is cyclo (-D-Ala<sup>1</sup>-R<sub>2</sub><sup>2</sup>-D-MeAsp<sup>3</sup>-R<sub>4</sub><sup>4</sup>-Adda<sup>5</sup>-D-Glu<sup>6</sup>-Mdha<sup>7</sup>-), where R<sub>2</sub> and R<sub>4</sub> are the two variable L-amino acids, and their abbreviations are used as the suffix of the amino acid name (Fig. 1). Adda, Mdha and MeAsp denote (2*S*,3*S*,8*S*,9*S*)-3-amino-9-methoxy-10-phenyl-2,6,8-trimethyldeca-4(*E*),6(*E*)-dienoic acid [22], *N*-methyldehydroalanine and β-methylaspartic acid, respectively. MCLR (Fig. 1, M.W. 994) is the most popular member found among the microcystins, where its two L-amino acid positions are Leu as R<sub>2</sub> and Arg as R<sub>4</sub>. The MS<sup>2</sup> spectrum of the protonated molecule (M+H)<sup>+</sup> at  $m/z$  995 of MCLR obtained from the LC/MS/MS on ion trap analysis is shown in Fig. 2.

The prominent fragment ions at  $m/z$  866 (*Ion a, e*), 728 (*Ion g*), 710 (*Ion g'*), 682 (*Ion b*), 599 (*Ion c*), 570 (*Ion f'*), 553 (*Ion f*) and 470 (*Ion d, h*) were observed together with some ions in the higher mass region of the MS<sup>2</sup> spectrum. These ions are generated by the cleavage of peptide bonds, because the mass numbers of the ions correspond to the difference between the precursor ion ( $m/z$  995) and the liberated peptide moieties. Generally, these ions must be formed by the two-bond fissions of the precursor ion, in which the initial fission of the cyclic structure to a linearized one and subsequent fission(s) at the peptide bonds are included. For the MCLR, the following peptide bonds, Arg-MeAsp, MeAsp-Leu, Ala-Mdha and Mdha-Glu, were initially cleaved as shown in Fig. 3. The resulting fragment ions were classified into four groups: (A) *Ions a~d* by the cleavage at Arg-MeAsp, (B) *Ion e, f'* by the cleavage at Mdha-Glu, (C) *Ions g* and *g'* by the cleavage at MeAsp-Leu and (D) *Ion h* by the cleavage Ala-Mdha. As shown in Fig. 3, it was found that the observed ions always contain the Arg moiety.

In order to rationalize the fragmentation of MCLR using the LC/MS/MS under ion trap conditions mentioned above, we compared the fragmentation of MCLR with those of three related microcystins that are structurally, but slightly different: dihydromicrocystin-LR [21] (dihydro-MCLR, M.W. 996) contains MeAla (*N*-methylalanine) instead of Mdha, microcystin-YR (MCYR, M.W. 1044) has Tyr replaced by Leu in MCLR, and 3-desmethymicrocystin-LR (3-des-MCLR, M.W. 980) is a desmethylated analogue of MCLR at the 3 position (Fig. 1). The predominant fragment ions observed in the MS<sup>2</sup> spectra of the related microcystins are summarized in Table 1. This is a shifting technique [23], and these ions are classified into two groups, the shifted or non-shifted groups as follows: in the



**Fig. 2** MS<sup>2</sup> spectrum of MCLR.



**Fig. 3** Fragmentation scheme of MCLR using LC/MS/MS under ion trap conditions.

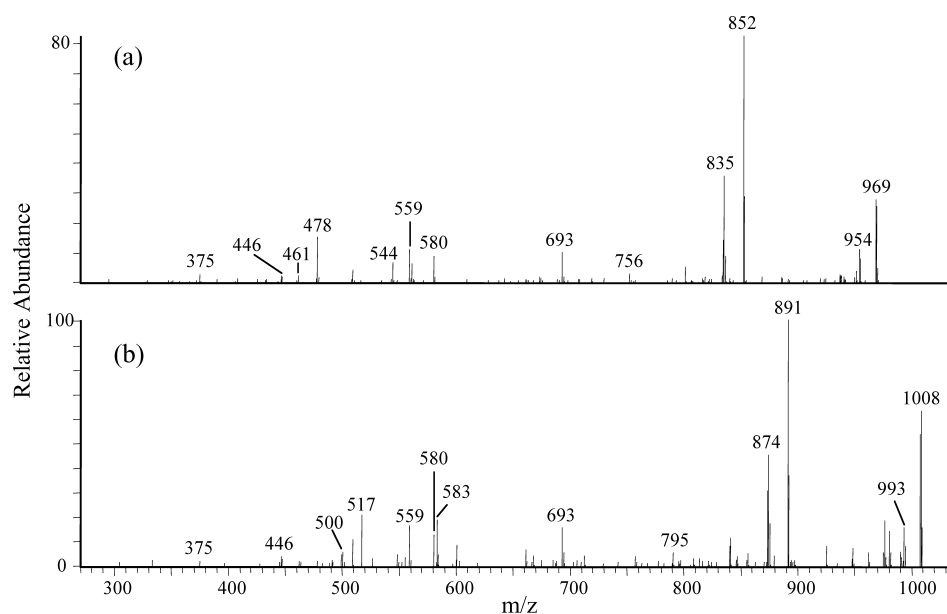
case of dihydro-MCLR, the MeAla containing fragment ion is shifted by 2 mass units; in the case of MCYR, the Tyr containing fragment ion is shifted by 50 mass units; and in the case of 3-des-MCLR, the Asp containing fragment ion is shifted by -14 mass units. The principal fragment ions of MCLR are compared with those of three related

microcystins (Table 1). As discussed above, they are classified into two groups, the shifted or non-shifted groups. This shifting behavior definitely supported the fact that the fragmentation of MCLR and related microcystin using LC/MS/MS under ion trap conditions is reproducible, and these fragment ions are available for sequencing the

**Table 1** Fragment ions of MCLR, dihydro-MCLR, MCYR and 3-des-MCLR observed in the MS<sup>2</sup> spectra using LC/MS/MS under ion trap conditions.

| Ion series | Ion                                | Microcystins |              |            |            |
|------------|------------------------------------|--------------|--------------|------------|------------|
|            |                                    | MCLR         | Dihydro-MCLR | MCYR       | 3-Des-MCLR |
|            |                                    | <i>m/z</i>   |              |            |            |
|            | M+H                                | 995          | 997 (+2)     | 1045 (+50) | 981 (-14)  |
|            | Fragment ion                       |              |              |            |            |
| <i>a</i>   | Arg-Adda-Glu-R7-Ala-R2             | 866          | 868 (+2)     | 916 (+50)  | 866        |
| <i>b</i>   | Arg-Adda-Glu-R7                    | 682          | —            | 682        | —          |
| <i>c</i>   | Arg-Adda-Glu                       | 599          | 599          | 599        | 599        |
| <i>d</i>   | Arg-Adda                           | 470          | 470          | —          | —          |
| <i>e</i>   | R7-Ala-R2-R3-Arg-Adda              | 866          | 868 (+2)     | 916 (+50)  | 852 (-14)  |
| <i>f</i>   | R7-Ala-R2-R3-Arg-NH <sub>2</sub>   | 570          | 572 (+2)     | 620 (+50)  | 556 (-14)  |
| <i>f</i>   | R7-Ala-R2-R3-Arg                   | 553          | 555 (+2)     | 603 (+50)  | 539 (-14)  |
| <i>g</i>   | R3-Arg-Adda-Glu                    | 728          | 728          | 728        | 714 (-14)  |
| <i>g'</i>  | R3-Arg-Adda-Glu(-H <sub>2</sub> O) | 710          | 710          | 710        | 696 (-14)  |
| <i>h</i>   | Ala-R2-R3-Arg                      | 470          | 470          | 520 (+50)  | 456 (-14)  |
|            | Constituent amino acid             |              |              |            |            |
|            | R2                                 | Leu          | Leu          | Tyr        | Leu        |
|            | R3                                 | MeAsp        | MeAsp        | MeAsp      | Asp        |
|            | R7                                 | Mdha         | MeAla        | Mdha       | Mdha       |

—: Not observed.

**Fig. 4** MS<sup>2</sup> Spectra of MCLF (a) and MCLW (b).

constituent amino acids in the microcystins.

### Fragmentation Pattern of Microcystin without Arg

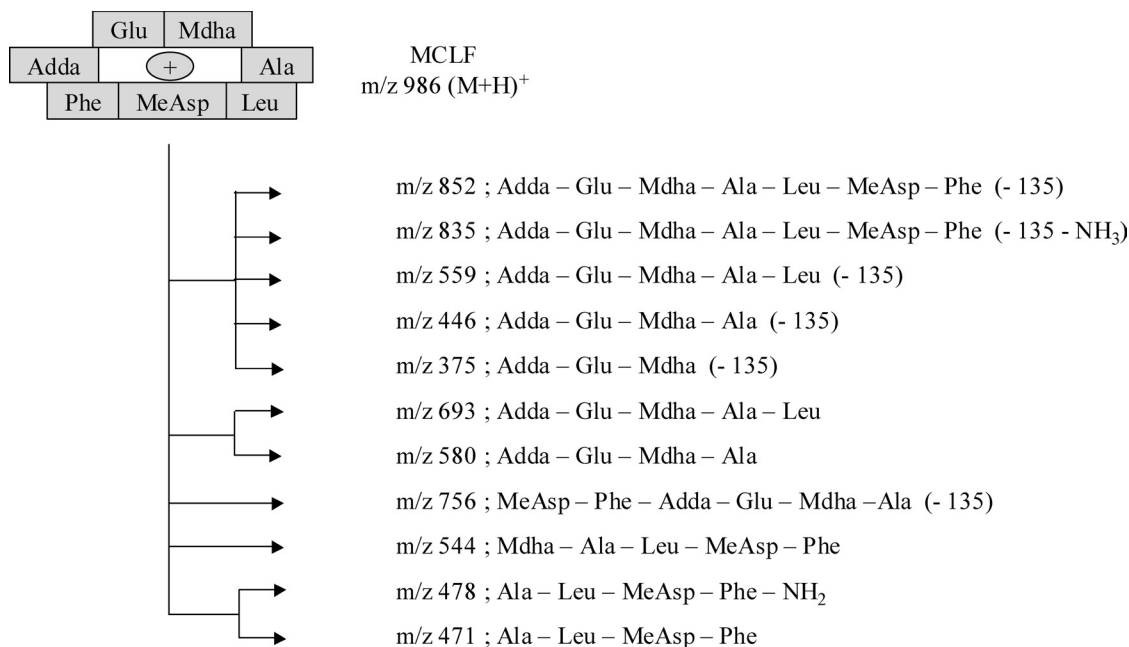
As mentioned above, the mass spectrometric behavior of MCLR and related microcystins was discussed and it was found that any detected fragment ion contained the Arg moiety. It is well known that microcystins without Arg as the constituent amino acid are also produced by cyanobacteria and they often are designated as a “hydrophobic microcystin”. Microcystin-LF (MCLF, M.W. 985) and -LW (MCLW, M.W. 1024) contain Phe and Trp instead of Arg in the MCLR, respectively (Fig. 1). The MS<sup>2</sup> spectra of the protonated molecules for the two microcystins

are shown in Fig. 4 and the observed ions are summarized together with those of the MCLR in Table 2. The ion series *f* and *h* were only detected in MCLF and MCLW, but almost no other ions were identified as being in the ion series observed in the MCLR and related microcystins. Whereas the fragment ions at *m/z* 693, 580, 559, 446 and 375 were commonly observed in the MCLF and MCLW, other ions at *m/z* 891 (852), 874 (835) and 795 (756) individually appeared. The fragment ions at *m/z* 891 and 852 were found as the base peaks in the mass spectra of the MCLW and MCLF, respectively. These ions were 134 mass units less than the precursor ions ((M+H)<sup>+</sup>) and considered to be generated by the cleavage between the 8 and 9 positions in

**Table 2** Fragment ions of MCLF and MCLW observed in the MS<sup>2</sup> spectra using LC/MS/MS under ion trap conditions

| Ion series | Ion   | Microcystins |          |            |
|------------|---|--------------|----------|------------|
|            |   | MCLR         | MCLF     | MCLW       |
| <i>m/z</i> |   |              |          |            |
|            | M+H   | 995          | 986 (−9) | 1025 (+30) |
|            | Fragment ion  |              |          |            |
| <i>a</i>   | R4-Adda-Glu-Mdha-Ala-Leu                              | 866          | —        | —          |
| <i>b</i>   | R4-Adda-Glu-Mdha                                      | 682          | —        | —          |
| <i>c</i>   | R4-Adda-Glu   | 599          | —        | —          |
| <i>d</i>   | R4-Adda   | 470          | —        | —          |
| <i>e</i>   | Mdha-Ala-Leu-MeAsp-R4-Adda                            | 866          | —        | —          |
| <i>f</i>   | Mdha-Ala-Leu-MeAsp-R4-NH <sub>2</sub>                 | 570          | —        | —          |
| <i>f</i>   | Mdha-Ala-Leu-MeAsp-R4                                 | 553          | 544 (−9) | 583 (+30)  |
| <i>g</i>   | MeAsp-R4-Adda-Glu                                     | 728          | —        | —          |
| <i>g'</i>  | MeAsp-R4-Adda-Glu(H <sub>2</sub> O)                   | 710          | —        | —          |
| <i>h'</i>  | Ala-Leu-MeAsp-R4-NH <sub>2</sub>                      | —            | 478      | 517        |
| <i>h</i>   | Ala-Leu-MeAsp-R4                                      | 470          | 461 (−9) | 500 (+30)  |
|            | Adda-Glu-Mhda-Ala-Leu                                 | —            | 693      | 693        |
|            | Adda-Glu-Mdha-Ala                                     | —            | 580      | 580        |
|            | Adda-Glu-Mdha-Ala-Leu-MeAsp-R4(-135)                  | —            | 852      | 891        |
|            | Adda-Glu-Mdha-Ala-Leu-MeAsp-R4(-135-NH <sub>3</sub> ) | 844          | 835      | 874        |
|            | Adda-Glu-Mdha-Ala-Leu(-135)                           | —            | 559      | 559        |
|            | Adda-Glu-Mdha-Ala(-135)                               | 446          | 446      | 446        |
|            | Adda-Glu-Mdha(-135)                                   | 375          | 375      | 375        |
|            | MeAsp-R4-Adda-Glu-Mdha-Ala(-135)                      | —            | 756      | 795        |
|            | Constituent amino acid                                |              |          |            |
|            | R4  | Arg          | Phe      | Trp        |

—: Not observed.



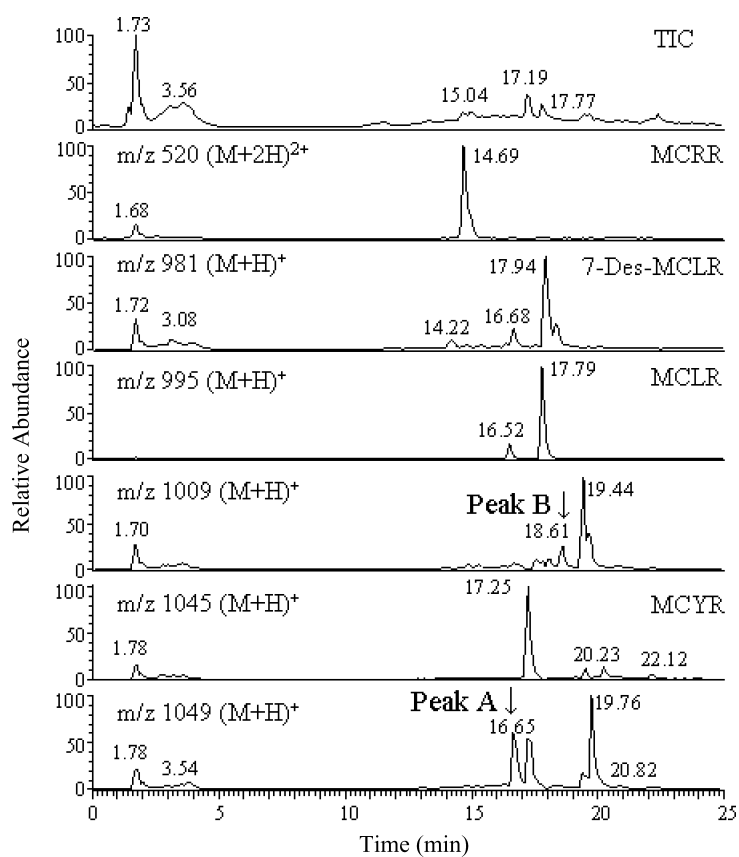
**Fig. 5** Fragmentation scheme of MCLF using LC/MS/MS under ion trap conditions.

the side chain of Adda. This cleavage has been previously reported [11]. Thus the fragment ions detected in the MS<sup>2</sup> spectra of MCLF and MCLW are grouped into three categories: the first one contains the initial carbon–carbon cleavage at the Adda moiety followed by fission of the cyclic structure into a linearized one and subsequent fission of the peptide bonds, the second consists of the fission of the cyclic structure at a peptide bond between Adda and Leu (Trp) into linearized one, but the carbon–carbon cleavage does not occur, and the third one is found in the case of the MCLR. The fragmentation scheme for MCLF is depicted in Fig. 5 and provides informative sequence information even if the carbon–carbon cleavage occurs at the Adda moiety.

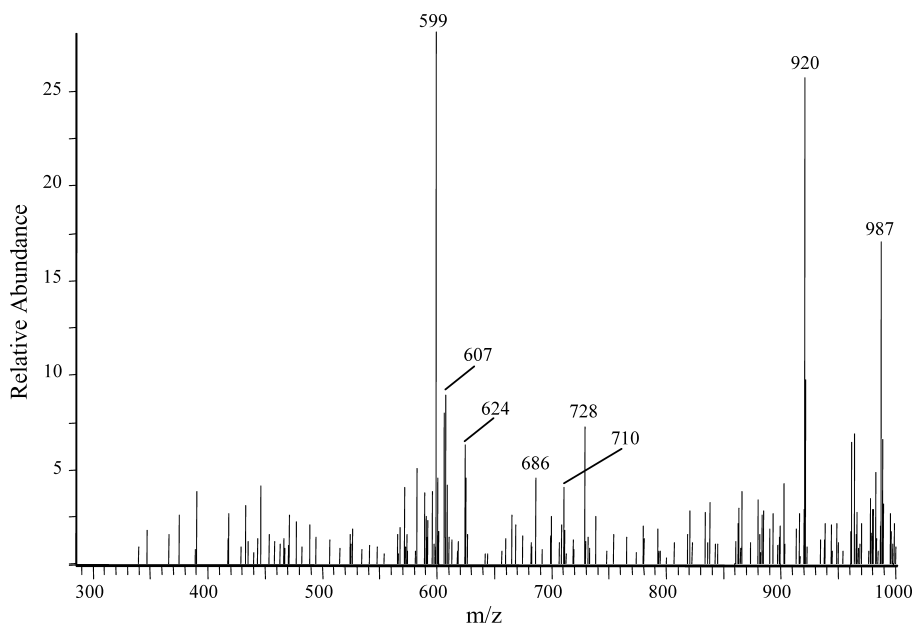
### Structural Characterization of Microcystins in Cyanobacterial Blooms by LC/MS

The systematic study mentioned above strongly suggested that the fragment ions of microcystins generated in the MS<sup>2</sup> analysis of the LC/MS/MS under ion trap conditions are informative and available for the sequencing the constituent amino acids. Therefore, the established fragmentation pattern was applied to identify the microcystins in the crude extracts from a bloom sample. The total ion chromatography (TIC) and mass chromatograms of the microcystins detected in the LC/MS/MS under ion trap conditions of the crude extracts from the lyophilized cells of cyanobacteria collected from Lake Suwa in August 2005 are shown in Fig. 6. Four ion peaks were readily identified

as MCLR and 7-des-MCLR, MCRR, MCYR based on their molecular weights and retention times. However, it was difficult to identify two peaks eluted at 16.65 minutes (peak A) and 18.61 minutes (peak B) using only the information concerning their molecular weights and retention times. Peak A showed the protonated molecule at *m/z* 1049, which is 4 mass units higher than MCYR. The MS<sup>2</sup> spectrum of peak A is shown in Fig. 7, in which the prominent fragment ions were observed at *m/z* 920, 728, 710, 686, 624, 607 and 599 in the MS<sup>2</sup> analysis of the protonated molecule at *m/z* 1049 as the precursor ion. The resulting fragmentation pattern showed that peak A contains Arg. The fragment ions at *m/z* 728, 710 and 599 were commonly observed in the spectrum of the MCYR, and identified as *Ion g*, *g'* and *c*. Four fragment ions at *m/z* 920, 686, 624 and 607 appeared 4 mass units higher than *Ions a* or *e*, *b*, *f'* and *f* of the MCYR, respectively. Thus, a linear structure, Tyr-MeAsp-Arg-Adda-Glu, was determined as the partial structure of peak A. A fragment ion at *m/z* 920 was detected 129 mass units less than the protonated molecule at *m/z* 1049, and this difference corresponded to loss of Glu or MeAsp. This indicated that the structural variation between peak A and MCYR is due to the Mdha or Ala moiety. The ion at *m/z* 686 was 4 mass units higher than *m/z* 682 (Arg-Adda-Glu-Mdha) observed in the MS<sup>2</sup> analysis of the MCYR, suggesting that this fragment ion was Arg-Adda-Glu-Ser. Therefore, peak A was considered to be [Ser<sup>7</sup>]-MCYR, in which Mdha in the MCYR was replaced by Ser and the predicted fragmentation pattern for



**Fig. 6** Total ion chromatogram and mass chromatograms of microcystins of bloom extracts collected from Lake Suwa.



**Fig. 7** MS<sup>2</sup> spectrum of Peak A.



peak A coincide with the established one. Thus, peak A was identified to [Ser<sup>7</sup>] MCYR apart from the absolute configuration of Ser.

Peak B showed the protonated molecule at  $m/z$  1009 in the LC/MS/MS under ion trap conditions. The fragment ions were observed at  $m/z$  880, 710, 682, 599, 584, 567 and 484 in the MS<sup>2</sup> analysis of the protonated molecule at  $m/z$  1009 as the precursor ion (data not shown). The fragment ions at  $m/z$  710 and 599 were identified as *Ions g'* and *c*, respectively. Because the fragment ion at  $m/z$  682 corresponded to *Ion b*, a partial structure of peak B was supposed to be MeAsp-Arg-Adda-Glu-Mdha. Other fragment ions at  $m/z$  880, 584, 567 and 484 were 14 mass units higher than *Ions a* or *e*, *f'*, *f* and *h*. These data suggested that peak B contains the methylated Ala or Leu as the constituent amino acid, so that the structure was considered to be [methylated Ala<sup>1</sup>] or [methylated Leu<sup>2</sup>] MCLR. Such a microcystin has not yet been reported.

## Discussion

Many cyclic peptides are used in modern life and possibly in our future. As discussed in the introduction, no suitable methodology has been developed for the structural characterization of cyclic peptides, because cyclic peptides from natural sources have complicated structures including unusual amino acids. Particularly, no reproducible and reliable method has been established for the sequencing of the constituent amino acids in the case of a small amount of sample. Although tandem mass spectrometry, such as four sector MS/MS and triple stage MS/MS, has been used for this purpose, they have not provided satisfactory results [11~14]. In this study, a systematic investigation was planned to confirm the applicability of LC/MS/MS under ion trap conditions for the sequencing using microcystin as a typical cyclic peptide, and the final goal was to successfully apply the obtained information of known microcystins to characterize unknown ones.

First of all, the mass spectrometric behavior of the microcystin was carefully examined under MS<sup>2</sup> analysis conditions. The MS<sup>2</sup> spectrum of the protonated molecule (M+H)<sup>+</sup> provided many fragment ions which are mainly generated by the cleavage of peptide bonds. These ions were formed by the two-bond fissions of the precursor ion, in which an initial fission of the cyclic structure to a linearized one and subsequent fission(s) at the peptide bonds are included. They belong to the *bn* ions according to the notation by Biemann [24], which are desirable for the sequencing. In the case of the microcystin-containing Arg, the observed ions always contain the Arg moiety (Fig. 3).

The fragmentation of other microcystins without Arg showed a different pattern and the cleavage between the 8 and 9 positions in the side chain of Adda predominates prior to the fission of the peptide bonds (Fig. 5). The obtained fragmentation was confirmed by a shifting technique using structurally related microcystins. Diehnelt *et al.* reported the identification of the fragment ions of microcystins using an ion tap Fourier transform ion cyclotron [10]. They also analyzed MCLF and the fragment ions at  $m/z$  852, 835, 693, 580, 559 and 544 were similar to the fragment ions obtained from our experiment, which are available for the sequence information. In other microcystins, the fragment ions were identified by their  $m/z$  value and molecular composition, and most of them coincided with our proposed fragment ions. Moreover, they reported that the fragmentation of microcystin containing Arg or without Arg showed similarly the different pattern. These results indicate that fragment ions obtained from LC/MS/MS under ion trap conditions are reproducible and this system is applicable to the structural characterization of microcystins. Poon *et al.* also reported the analysis of microcystins using LC/MS/MS under ion trap conditions, and the fragment ions obtained from the MS<sup>2</sup> analysis was described in detail [18]. However, their proposed fragmentation pattern was not systematically discussed, and it was difficult to apply them for the structural characterization of unknown microcystins, because there was no discussion regarding the generation of fragment ions including the two-bond fission and the difference of fragmentation pattern of microcystins containing Arg or without Arg as discussed above.

The established fragmentation pattern was then used to identify the microcystins in the crude extracts from a bloom sample, and 4 known and 2 unknown microcystins were found. One of the unknown microcystins was identified to [Ser<sup>7</sup>] MCYR, but another corresponded to either the [methylated Ala<sup>1</sup>] or [methylated Leu<sup>2</sup>] MCLR. The structure was not fully elucidated, because there were no informative ions generated by the cleavage of the peptide bond between Ala and Leu. We considered that this method is difficult to apply to the structural characterization of unknown microcystins structurally modified at [Ala<sup>1</sup>] or [R<sub>4</sub>] position (Fig. 1). MS<sup>2</sup> spectrum using centroid mode was not always sufficient because obtained data was not scanned in profile mode. However we have analyzed isolated microcystins in centroid mode and made sure that the initial fission occurred the peptide bond definitely and fragment ions were generated as *b<sub>n</sub>* series ions. Further, we could find the fragmentation pattern of microcystins containing Arg or without Arg and clarified the difference. Therefore the obtained fragment ions using centroid mode

was applicable to the structural characterization of unknown microcystins and make possible to distinguish the fragment ion peaks from noise peaks. These experiments shown in this study strongly indicated that the obtained fragmentation pattern of the microcystins by LC/MS/MS under ion trap conditions is reproducible, and these fragment ions are available for sequencing the constituent amino acids in the microcystins. In a subsequent step, we have to confirm that the established fragmentation by LC/MS/MS under ion trap conditions is available or not for other types of cyclic peptides.

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