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[Article begins on next page]
A novel heterocyst glycolipid detected in a pelagic N$_2$-fixing cyanobacterium of the genus *Calothrix*

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ABSTRACT

Previous studies have shown that heterocyst glycolipids (HGs) are unique markers for N$_2$-fixing heterocystous cyanobacteria. In this study, the HGs of a marine pelagic Calothrix sp. CCY1611 isolated from the tropical western North Atlantic were analyzed by ultra-high pressure liquid chromatography–high resolution mass spectrometry and it was shown that this organism contains an unusual C$_{28}$ triol HG with a methylated C$_6$ sugar (methyl-HG$_{28}$ triol) head group. Gas chromatography–mass spectrometry analysis of the sugar released from the novel HG by acid methanolysis revealed that the sugar is likely 6-O-methyl-β-D-glucopyranose. We propose that this methyl-HG$_{28}$ triol is a potential biomarker for pelagic members of the genus Calothrix.

1. Introduction

In all heterocystous cyanobacteria studied to date, the heterocyst cell wall contains heterocyst glycolipids (HGs) (Nichols and Wood, 1968; Abreu-Grobois et al., 1977; Gambacorta et al., 1995; Bauersachs et al., 2009a; 2014). These HGs almost universally comprise a hexose head group (hereafter C$_6$) glycosidically bound to long-chain diols, triols, or hydroxyketones (Bryce et al., 1972; Gambacorta et al., 1998; Bauersachs et al., 2009b; 2011), except for some marine endosymbiotic cyanobacteria which contain pentose head groups (Schouten et al., 2013; Bale et al., 2015; 2018). Previous studies reported that HGs show structural diversity depending on the family level within the cyanobacteria divisions (Bauersachs et al., 2009a; 2014; 2017). The heterocystous cyanobacteria of the genus Calothrix are characterized by the presence of the 1-(O-hexose)-3,25,27-octacosanetriol (C$_6$ HG$_{28}$ triol) and 1-(O-hexose)-27-keto-3,25-octacosanediol (C$_6$ HG$_{28}$ keto-diol) (Gambacorta et al., 1998; Bauersachs et al., 2009a; Wörmer et al., 2012). However, these studies on Calothrix focused predominantly on benthic strains, while this genus is also known from the pelagic where it occurs as a symbiont of marine diatoms, specifically of
Chaetoceros (Foster et al., 2010; 2011). These symbioses are known as diatom-diazotroph associations. Previously, while analyzing the HG content of Calothrix sp. UTEX 2589, Schouten et al. (2013) found that alongside the C_{6} HG_{28} triol and C_{6} HG_{28} keto-diol there was also an unknown glycolipid eluting several minutes earlier than the known glycolipids. Based on mass spectral fragmentation patterns and molecular weight, this novel glycolipid was tentatively described as a HG_{28} triol containing a C_{6} sugar moiety which contained either an additional keto group, e.g. glucuronic acid instead of glucose, or an additional methyl group, e.g. by methylation of one of the hydroxyl groups (Schouten et al., 2013). This novel glycolipid was not present in the majority of benthic Calothrix species examined to date, but was present in a Calothrix isolated from an intertidal microbial mat (CCY0202; Schouten et al., 2013). Here, we identified this novel HG in a pelagic Calothrix sp. CCY1611 that was isolated from the surface water of the tropical western North Atlantic using ultra-high pressure liquid chromatography–high resolution mass spectrometry (UHPLC–HRMS) and acid methanolysis.

2. Methods

2.1. Isolation and culturing

Calothrix sp. CCY1611 was isolated from surface water from the tropical North Atlantic Ocean collected during a research cruise onboard the R/V Pelagia in 2014 (Station 20, 64PE393, cf. Bale et al., 2018). A surface water sample (1.5 L) was filtered over a 47 mm GFF (Whatman, Maidstone, UK). The filter was placed in a disk filled with agarose (0.6%) solidified seawater from the same location and subsequently stored at –80 °C until transport to the laboratory at NIOZ. Isolation of diazotrophic cyanobacteria was performed by transferring the GFF filter to a Petri dish with a solidified artificial seawater T^{0} medium (modified from Chen et al., 1996) with agarose (7 g L^{-1}) as the solidifying agent. The medium was supplemented with glucose (2 g L^{-1})
and the incubation was carried out in an incubator (model MLR-350, SANYO, Osaka, Japan) at 27 °C, with a 12-12 h light-dark cycle and a light intensity (photon density) of 20–30 µmol m\(^{-2}\) s\(^{-1}\). Once colonies appeared on the filter, they were transferred to new agarose medium without glucose, and a pure culture was obtained after repeated transfers of single trichomes using standard microbiological techniques. The isolate was identified as a *Calothrix* sp. based on its morphology using a light microscopy and sequencing of the 16S rRNA gene (GenBank accession number MH364376). In order to characterize its HGs, the strain was grown for 40 days in T\(^{0}\) liquid medium at 27 °C and harvested at stationary phase and stored at –20 °C until analysis.

### 2.2. Lipid extraction and analysis

The extraction of lipids from freeze dried biomass was carried out using a modified Bligh-Dyer extraction as described previously (Bale et al., 2013). UHPLC–HRMS was carried out as described by Bale et al. (2017) using an Agilent 1290 Infinity UHPLC was used, equipped with thermostatic auto-injector and column oven, coupled to a Q Exactive Orbitrap MS with Ion Max source with heated electrospray ionization (HESI) probe (Thermo Fisher Scientific, Waltham, MA, USA).

To confirm the structure of the sugar in the novel HG, it was isolated using semi-preparative HPLC and the normal phase system as described by Bale et al. (2017). The column effluent was collected in 1 min fractions and the fractions containing the novel HG were pooled. Acid methanolysis was performed on the isolated compound and hydroxyl groups were converted into trimethylsilyl (TMS) ester derivatives using N,O-\textit{bis}(trimethylsilyl)trifluoroacetamide (BSTFA) and pyridine (1:1; 20 min at 60 °C) before analysis by GC–MS using a Thermo Trace DSQ as described in Schouten et al. (2013). The sugar was identified by comparison of its mass spectrum with library mass spectra (NIST Mass Spectral Library, Version 2.0, 2012) and three
3. Results and discussion

Analysis of *Calothrix* strain CCY1611 by UHPLC–HRMS (Fig. 1a) indicated the presence of a C₆ HG₂₈ triol ([M+H]⁺ m/z 621.493) and a C₆ HG₂₈ keto-diol ([M+H]⁺ m/z 619.478) and a novel HG ([M+H]⁺ m/z 635.508), previously reported by Schouten et al. (2013) in two other *Calothrix* species. Initial structural identification was based on the HRMS² spectrum generated from the protonated molecule (Fig. 1b). The spectrum contained the same five ions as described in the MS² spectrum of the C₆ HG₂₈ triol (Bauersachs et al., 2009b), at m/z 459.441, 441.430, 423.420, 405.409 and 387.398, suggesting that the alkyl chain is also a 3,25,27-octacosanetriol. The product ion at m/z 459.441 corresponded to a neutral loss of a head group of mass 176.067 Da (C₇H₁₂O₅). The accurate mass of the unknown HG [M+H]⁺ ion (m/z 635.508) allowed us to distinguish between the two hypothesized structures for the head group by Schouten et al. (2013) since the accurate mass of the HG with an additional keto group on the C₆ sugar, e.g. glucuronic acid, (C₃₄O₁₀H₆₆) is 635.473, whereas accurate mass of the HG with an additional methyl group on the C₆ sugar (C₃₅O₉H₇₀) is 635.509. This demonstrates that the unknown HG compound is a methylated C₆ HG with a C₂₈ triol core (methyl-HG₂₈ triol). This was confirmed by GC–MS analysis of the sugar released by acid methanolysis of the isolated (by preparative HPLC) novel HG. The mass spectrum (Fig. 2) provided evidence that there was no methylation at the C-2, C-3 or C-4 position and that, due to the methanolysis of the alcohol chain, there was a methylation at the C-1 position (Petersson and Samuelson, 1968). The additional methylation was therefore determined to be at the C-6 position. Furthermore, the spectrum was similar to the reported mass spectrum to 6-O-methyl-β-D-glucopyranose (NIST Mass Spectral
Library, Version 2.0, 2012), while its retention time and mass spectrum was different from our analysis of sugars with single methylations at the 1-O and 3-O position. Therefore, the novel HG was identified as 1-(O-6-O-methyl-β-D-glucopyranose)-3,25,27-octacosanetriol, a potential novel biomarker for cyanobacteria in the genus *Calothrix*.

To date, there have been no environmental reports of the methyl-HG$_{28}$ triol, likely due to limitations of previous analytical methods such as selective reaction monitoring (SRM) (Bauersachs et al., 2009b; Bale et al., 2015), which did not include the transition of the methyl-HG$_{28}$ triol. Further research examining more species of pelagic and benthic *Calothrix* should reveal if this novel HG is associated with pelagic strains of *Calothrix* or whether it has a wider distribution within the genus.

**Acknowledgments**

We would like to thank John Volkman and Bart van Dongen for their helpful comments. We thank J. Ossebaar, A. Mets and S. Vreugdenhil for analytical and technical support. We thank Professor T. Villareal for helpful discussions on this subject. This work was supported by the Netherlands Organisation for Scientific Research (NWO) through grant 822.01.017 to S.S. This project has also received funding from the European Research Council (ERC) under the European Union’s Horizon 2020 research and innovation programme (grant agreement No 694569 to J.S.S.D). S.S. and J.S.S.D were financially supported by the Netherlands Earth Systems Science Centre (NESSC), a Gravitation grant (024.002.001) from the Dutch Ministry of Education, Culture and Science.

*Associate Editor–Bart van Dongen*
References


Figure 1. (a) UHPLC-HRMS partial base peak chromatogram (Gaussian smoothed) showing the distribution of heterocyst glycolipids (filled peaks) in the Bligh and Dyer extract of *Calothrix* sp. CCY1611. Insert: proposed structure of the novel methyl-C\textsubscript{6} HG\textsubscript{28} triol, 1-(O-6-O-methyl-β-D-glucopyranose)-3,25,27-octacosanetriol. (b) MS\textsuperscript{2} spectrum of the novel methyl-C\textsubscript{6} HG\textsubscript{28} triol with [M+H]\textsuperscript{+} 635.508. For interpretation of the acyl chain fragments see Bauersachs et al. (2009b).
Figure 2. Mass spectra of trimethylsilylated (TMS) sugar moiety of the novel heterocyst glycolipid with structure shown as insert.
Dear Professor Volkman

Thank you for considering our manuscript for publication in Organic Geochemistry as a Note. We thank you for your helpful editorial edits and we have additionally made all the edits requested by the reviewer.

Yours sincerely,
Dr. Nicole Bale, on behalf of co-authors

Editorial changes:
use correct degree symbol for °C, and space before °C.
We have made this change

Put subsection headings in italics, not bold.
We have made this change

Insert a blank line between sections.
We have made this change

In m/z only the m and z are in italics.
We have made this change

Reviewer #1: This is generally well written note with an interesting new identification of what could be a potential new biomarker. I suggest accepting it after some minor corrections.

We thank the reviewer for the comments. We have addressed the specific points below.

However, I feel that it may be too early to make this a potential biomarker for the genus Calothrix, as stated in the abstract (lines 18-19), particularly since it was not present in the majority of the benthic Calothrix species (line 42-23). Could it be that this is a potential biomarker for pelagic Calothrix species? In addition, this conclusion is not completely in line with the final line of the MS (lines 110-110). Here the authors mention that 'Further research examining more species of Calothrix as well as other heterocystous cyanobacteria should reveal if this novel HG is restricted to this genus', which basically means that they themselves would not call it a biomarker for this genus yet.

We agree that it is not correct to associate the novel biomarker with the entire genus Calothrix at this stage. A wide range of benthic strains did indeed not contains the novel HG and we theorized ourselves that the biomarker is associated with pelagic Calothrix. Unfortunately, we are not able to confirm whether the two additional Calothrix strains that were found to contain the glycolipid were pelagic. We have rewritten the lines mentioned above as well as at other points in the manuscript to focus in on pelagic Calothrix rather than making broad statements about the entire genus.

Other minor comments:
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