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Unusual biogenic calcite structures in two shallow lakes, James Ross Island, Antarctica

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Abstract

The floors of two shallow endorheic lakes, located on volcanic surfaces on James Ross Island, are covered with calcareous organosedimentary structures. Their biological and chemical composition, lake water characteristics, and seasonal variability of the ther-

- ⁵ mal regime are introduced. The lakes are frozen down to the bottom eight-nine months per year and their water chemistry is characterized by low conductivity and neutral to slightly alkaline pH. The photosynthetic microbial mat is composed of filamentous cyanobacteria and microalgae that are considered to be Antarctic endemic species. The mucilaginous black biofilm is covered by green spots formed by a green microalga
- and the macroscopic structures are packed together with fine material. Thin sections consist of rock substrate, soft biofilm, calcite spicules and mineral grains originating from different sources. The morphology of the spicules is typical of calcium carbonate monocrystals having a layered structure and worn surface, which reflect growth and degradation processes. The spicules chemical composition and structure correspond
- to pure calcite. Lakes age, altitude, morphometry, geomorphological and hydrological stability, including low sedimentation rates, together with thermal regime predispose the existence of this community. We hypothesize that the precipitation of calcite is connected with the photosynthetic activity of the green microalgae that were not recorded in any other lake in the region. This study has shown that the unique community producing biogenia calcite pricules is guite different to any yet described.
- ²⁰ ducing biogenic calcite spicules is quite different to any yet described.

1 Introduction

The floors of most Antarctic lakes are covered with photosynthetic microbial mats (Vincent and Laybourn-Parry, 2008). However, the degree of disturbance plays a key role in the development of microbial mats. When growing in low-disturbance habitats, inter-

²⁵ actions between benthic microbial communities and their environments can produce complex emergent structures on a range of spatial scales. Such structures are best



developed in extreme environments, including benthic communities of deep, perennially ice-covered Antarctic lakes, where physical and chemical conditions, and/or geographical isolation preclude larger organisms that could otherwise disrupt organised microbial structures (Wharton, 1994; Andersen et al., 2011). Many organisedimentary

- ⁵ structures that emerge in these conditions are laminated and accrete through episodic trapping of sediments or grains and precipitation of minerals within a growing biogenic matrix (e.g. Arp et al., 2001; Reid et al., 2003). In perennially ice-covered lakes, the seasonality of growth imposed by the summer-winter light-dark conditions can induce annual growth laminations (Hawes et al., 2001), reinforced by calcite precipitation during growth and accieve diagonasis (Wharton et al., 1092; Wharton, 1094; Sutherland)
- ing growth and sediment diagenesis (Wharton et al., 1982; Wharton, 1994; Sutherland and Hawes, 2009). Calcite precipitation is not, however, a prerequisite for laminated, stromatolite-like communities (Walter, 1976; Schieber, 1999; Yamamoto et al., 2009).

Precipitation of calcite by expulsion (segregation) is also a common process in the nature related with the freezing of common low ionic strength Ca^{2+} - HCO_3^- waters. Cal-

- cite precipitation related to water freezing was observed and described also from various polar-alpine settings, e.g. from lake bottoms of Dry Valleys in Antarctica (Nakai et al., 1975); as a results of aufeis (icing, naled) formation in Northern Canada (Clark and Lauriol, 1997); as crystalline precipitates that form subglacially on bedrock, reported from numerous locations (Ng and Hallet, 2002); as fine-grained calcite powders in sub-
- glacial deposits and in aufeis formations, Svalbard (Wadham et al., 2000); in basal ice and subglacial clastic deposits of continental glaciers of Switzerland (Fairchild et al., 1993); as calcite pendants occurring beneath coarse clasts in well-drained sediments, Svalbard (Courty et al., 1994); as calcite coatings in cavities in cold-climate Pleistocene deposits of Western Transbaikalia, Russia, and in modern surface deposits at Seymour
- Island, Antarctica (Vogt and Corte, 1996); cryogenic calcite powder from modern cave environment (Clark and Lauriol, 1992).

James Ross Island belongs to a transitory zone between the maritime and continental Antarctic regions (Øvstedal and Lewis Smith, 2001). Air temperature records indicate progressive warming trends from 1.5 to 3.0 °C over the Antarctic Peninsula



during the past 50 years (Turner et al., 2014). More than 80 % of the island surface is covered with ice (Rabassa et al., 1982). Only the northernmost part of the island, the Ulu Peninsula, is significantly deglaciated and represents one of the largest ice-free areas in the northern part of the Antarctic Peninsula. The origin of the lakes on James

- ⁵ Ross Island is related to the last glaciations of the Antarctic Peninsula ice sheet and retreat of the James Ross Island ice cap during the late Pleistocene and the Holocene (Nývlt et al., 2011; Nedbalová et al., 2013). Interactions between volcanic landforms and glacial geomorphology during previous glacial-interglacial cycles, the Holocene paraglacial and periglacial processes and relative sea level change have resulted in
- the complex present-day landscape of James Ross Island (Davies et al., 2013). All of these processes have influenced the development of the lakes which are found on the Ulu Peninsula at altitudes from <20 m above sea level (a.s.l.) near the coast to 400 m a.s.l. in the mountain areas (Nedbalová et al., 2013).</p>

During two Czech research expeditions (2008 and 2009) to James Ross Island, lake ecosystems of the Ulu Peninsula were studied in respect to their origin, morphometry, physical, chemical and biological characteristics (Nedbalová et al., 2013, together with detailed cyanobacterial and microalgal diversity descriptions (Komárek and Elster, 2008; Komárek et al., 2011; Kopalová et al., 2013; Škaloud et al., 2013; Komárek et al., 2015). As part of this study, we encountered 1 to 5 mm scale calcareous organosedi-

- ²⁰ mentary structures on the floor of two endorheic lakes, 1 and 2, which are quite different to any microbially mediated structures yet described from modern environments. These shallow lakes on higher-lying levelled surfaces originated after the deglaciation of volcanic mesas which became ice-free some 6.5–8 ka ago (Johnson et al., 2011) and are considered among one of the oldest in the region. However, a later appear-
- ²⁵ ance of these lakes is also possible, as we have no exact dates from their sediments (Nedbalová et al., 2013).

The aim of this paper is to describe in detail the chemical and biological composition of the organosedimentary structures (stromatolites sensu Allwood et al., 2006) together with the limnological characteristics of the two lakes. A hypothesis concerning



the formation of calcite spicules is also presented. The results of this study can serve as a baseline for understanding microbial behaviors in forming these organosedimentary structures, which will provide insight into the interpretation of fossil forms from early Earth.

5 2 Materials and methods

2.1 Study site

Endorheic lake 1 (63°54′11.7″ S, 57°46′49.9″ W, altitude 65 m a.s.l.) and endorheic lake 2 (63°53′54.6″ S, 57°46′33.8″ W, altitude 40 m a.s.l.) are shallow lakes located near Andreassen Point on the E coast of the deglaciated Ulu Peninsula, in the northern part of James Ross Island, NE Antarctic Peninsula (Fig. 1). They are shallow with maximum depth of 1.1 and 0.9 m, and mean depth of 0.5 and 0.3 m. Their catchment areas are 340 515 and 368 586 m², lake area 4220 and 2970 m² and water volume 2183 and 1037 m³, respectively (Nedbalová et al., 2013). Melt water from the surrounding snowfields feed the lakes for a few weeks during the austral summer. The water level in both lakes fluctuated dramatically. Water is mainly lost through evaporation from the ice free water surface. During this period, intense evaporation in both lakes is coupled with macroscopic changes in the littoral belt (Fig. 2).

Climate conditions of the Ulu Peninsula are characterized by mean annual air temperatures around -7°C and mean summer temperatures above 0°C for up to four ²⁰ months (Láska et al., 2011a). The mean global solar radiation is around 250 W m⁻² in summer (December-February), with large day-to-day variation affected by extended cyclonic activity in the circumpolar trough and orographic effects over the Antarctic Peninsula (Láska et al., 2011b). The bedrock is composed of two main geological units, namely Cretaceous back-arc basin sediments and mostly subglacial Neogene

to Quaternary volcanic rocks (Olivero et al., 1986). The terrestrial vegetation is limited to non-vascular plants and composed predominantly of lichen and bryophyte tundra. A



large number of lakes can be found in this area, formed by glacial erosion and deposition, followed by glacier retreat during the Holocene (Nedbalová et al., 2013).

2.2 Sampling procedures

Lake 1 was sampled on 22 February 2008. In 2009, lake 1 was sampled on 5 January,
lake 2 on 12 January. Air temperature at 2 m above ground was measured by an automatic weather station (AWS) located nearby (Fig. 1). Incident global solar radiation was monitored with a LI-200 pyranometer (LI-COR, USA) at Mendel Station, located 11 km northwest of the study site (Fig. 1). The LI-200 spectral response curve covers wavelengths from 400 to 1100 nm with absolute error typically of ±3% under natural daylight conditions. Global radiation was measured at 10s time interval and stored as 30 min average values, while air temperature was recorded at 1 hour intervals from 1 February 2009 to 30 November 2010. In lake 1, water temperature was monitored from 10 February 2009 to 30 November 2010 at 1 hour intervals using a platinum resistance thermometer with Minikin T data logger (EMS Brno, Czech Republic) installed on the lake bottom.

Conductivity, pH, temperature and dissolved oxygen were measured in situ with a portable meter (YSI 600) at the time the lakes were ice free. Water samples were collected from the surface layer, immediately filtered through a 200 μ m polyamide sieve to remove zooplankton and coarse particles. Chlorophyll *a* was extracted from particles

- retained on Whatman GF/F glass microfiber filters according to Pechar (1987). After centrifugation, chlorophyll *a* was measured by a Turner TD-700 fluorometer equipped with a non-acidification optical kit. The remaining water was kept frozen until analyzed at the Institute of Hydrobiology (Czech Republic). The chemical analytical methods are given in Nedbalová et al. (2013). The stones covered by photoautotrophic mats –
- ²⁵ biofilm collected in the field were transported to the Czech Republic in a frozen and/or dry state, documented with stereomicroscope (Bresser, HG 424018) and used for (a) phytobenthos community description and isolation of dominant species, (b) fix for thin



section analyses, (c) scanning electron and optical microscopy, and (d) determination of the structure and chemical composition of calcium carbonate spicules.

2.3 Thin section analyses

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Thin section analyses were made to observe both rock substrate and inorganic particles within biofilms. Dry microbial mat were saturated with epoxy resins in vacuum, subsequently cut perpendicularly and saturated again with epoxy resin. The sample was cemented to a glass slide after grinding and polishing, and a thin section was prepared by final sectioning, grinding and polishing to a desired thickness of 50–55 µm. Thin sections of rocks were studied in transmitted (PPL) and polarized (XPL) light (Olympus BX-51M) and documented in transmitted light of a Nikon SMZ-645 optical microscope using NIS-Elements software.

2.4 Biofilm scanning electron and optical microscopy

The morphology of photoautotrophic mats and calcareous spicules was studied using standard methods of scanning electron microscopy (SEM) using back-scattered electrons (BSE; Jeol JSM-6380, Faculty of Science, Charles University) and optical microscopy (Nikon SMZ-645 using NIS-Elements software). Calcareous spicules were collected directly from the surface of biofilms. Samples studied in SEM were completely dried for 5 months at room temperature, then mounted on stubs with carbon paste and coated with gold prior to photomicrographing.

20 2.5 Structure and chemical composition of calcium carbonate spicules – EDS and EBSD analyses

The chemical composition of the analyzed spicules was measured by using the Link ISIS 300 system with 10 mm² Si-Li EDS detector on a CamScan 3200 scanning electron microscope (Czech Geological Survey, Prague). Analyses were performed using an accelerating voltage of 15 kV, 2 nA beam current, 1 µm beam size and ZAF cor-



rection procedures. Natural carbonate standards (calcite, magnesite, rhodochrosite, siderite and smithsonite) were used for standardization. Subsequent structural identification was confirmed by electron backscattered diffraction (EBSD). Identification data and crystallographic orientation measurements were performed on the same scan-

- ⁵ ning electron microscope using an Oxford Instruments Nordlys S EBSD detector. The thick sections used for EBSD applications were prepared by the process of chemo-mechanical polishing using colloidal silica suspension. The acquired EBSD patterns were indexed within Channe I5 EBSD software (Schmidt and Olensen, 1989) applying calcite and aragonite crystallographic models (Effenberger et al., 1981; Caspi et al., 1981;
- ¹⁰ 2005). Orientation contrast images were collected from a 4-diodes forescatter electron detector (FSD) integrated into the Nordlys S camera. EBSD pattern acquisition was carried out at 20 kV acceleration voltage, 3 nA beam current, 33 mm working distance and 70° sample tilt.

3 Results

3.1 General description of the lakes and water chemistry

Pictures and detailed bathymetric parameters of both lakes together with marked lines of water level and the maximum extent of the photosynthetic microbial mat littoral belt in lake 1 are presented in Fig. 2.

The physico-chemical characteristics of the lake water for both lakes are given in Ta-

- ²⁰ ble 1. The sampling of lake 1 (pH 7.4–7.9, saturation of oxygen 98.9 %) was performed during cloudy days. Oxygen supersaturation (128%) together with a relatively high pH (8.6) was observed in lake 2 during a sunny day. Conductivity was below $100 \,\mu\text{S}\,\text{cm}^{-1}$ in both lakes. The concentrations of dissolved inorganic nitrogen forms were low, whereas the concentration of dissolved reactive phosphorus (SRP) was $19.3 \,\mu\text{g}\,\text{L}^{-1}$ in lake 2.
- ²⁵ Relatively high concentrations of dissolved organic carbon, particulate nutrients and chlorophyll *a* were also characteristic for lake 2 (Table 1). The autotrophic biomass in



lake water was mostly formed by detached benthic species; no substantial phytoplankton developed in the lakes. The comparison of the two sampling dates available for lake 1 suggested high fluctuations of dissolved nutrient concentrations.

3.2 Thermal regime

- Fig. 3a shows the annual variation of daily mean water temperature in lake 1 and of daily mean air temperature measured 2 m above ground in the Solorina Valley (locations of air temperature, global radiation and water temperature sensors are marked in Figs. 1 and 2). Lake 1 is frozen to the bottom from the end of March to the end of October or beginning of November. Minimum daily mean winter temperatures on the bottom of the lake were about -12 and -10°C for 2009 and 2010, respectively. Minimum daily mean air temperatures in the same period were between -32 and -25°C. Positive summer daily mean water temperatures on the bottom of the lake were between 5 to 9°C from the beginning of November to the end of March. Air temperatures
- tween 5 to 9 °C from the beginning of November to the end of March. Air temperatures were frequently lower than water temperatures. Mean monthly water temperatures in
 the lake ranged from -10.4 (August 2009) to 5.8 °C (February 2010), while monthly mean air temperatures were between -18.7 to 0.7 °C. The differences were greater at the beginning of the winter season (June–July), due to a rapid drop of air temperature. Diurnal water and air temperature amplitudes in lake 1 and the Solorina Valley are shown in Fig. 3b. The highest night-day air temperature fluctuations (up to 28 °C) were
- 20 recorded during the winter months, while the lowest, in contrast, occurred in summer. Conversely, the highest night-day amplitudes of water temperature on the bottom of the lake were recorded from November to February, with typical values between 2 and 4°C.

The course of global solar radiation (Fig. 3c) is smooth, with the maximum daily ²⁵ mean of 385 W m⁻² during clear sky conditions around the summer solstice. Global radiation reaches the bottom of both lakes during the ice free period.

The relative frequency of hourly values of water temperature at the bottom of lake 1 and air temperature measured 2 m above ground in the Solorina Valley in the period



from 10 February 2009 to 30 November 2010 is shown in Fig. 4. Water temperature fluctuation is narrow, ranging from –15 to 8 °C. This plot also documents that the bottom of the lake is dry frozen, with temperatures from -15° C to most frequently –2 to –6°C (Fig. 4a). The growing season, with liquid water at temperatures from 2 to 4°C, covers only two-three months. With respect to air temperature (Fig. 4b), the seasonal thermal

regime in the Solorina Valley is much wider (typically from –38 to 8 °C) in comparison with the thermal regime at the bottom of lake 1, where was from –16 up to 8 °C.

The occurrence of days with water temperature higher than -4 °C in lake 1 and monthly mean global solar radiation in the period from February 2009 to November

- ¹⁰ 2010 is shown in Fig. 5. Owing to the variability of the weather conditions, lake water at the bottom is completely frozen only during the coldest one-two months per year (July–August); water temperature is higher than -4 °C in the rest of the year. The water temperature is above 0 °C (liquid phase) from November to April (139 days in average). A number of days with temperature between 0 and -4 °C remains the same as far liquid water accurate with ample abapted in the start and and dates towards to
- for liquid water occurrence with small changes in the start and end dates towards to the transition period (February–June and September–November, respectively). In such thermal conditions, the benthic littoral community can be metabolically active.

3.3 Littoral phytobenthos – biofilm community description

The littoral benthic community in lakes 1 and 2 are dominated by the heterocytous
 cyanobacterium *Calothrix elsteri* Komárek 2011, which forms a flat black biofilm on the upper surface of bottom stones (Fig. 6), followed by *Hassallia andreassenni* Komárek 2011 and *Hassallia antarctica* Komárek 2011. *Hassallia andreassenni* is associated with calcium precipitation, as described later. *Hassallia antarctica* was found in stone crevices, being only loosely attached to the substrate. Littoral benthic mats – biofilms
 on stones are co-dominated on the surface of the blackish cyanobacterial biofilm by the green filamentous and richly-branched alga *Hazenia broadyi* Škaloud et Komárek 2013

(Ulotrichales, Chlorophyceae). *Hazenia broadyi* grew in macroscopic colonies producing green spots (Fig. 6b, d). Later in the summer season, the green spots connected



micro fortified mucilaginous lines (Fig. 6c, d). Figure 6a shows the community in early spring whereas Fig. 6b, d originated from later summer when the littoral benthic community was already well developed with a dense coverage of *Hazenia broadyi* green spots. More detailed pictures (Fig. 6e, f) documented the structure of the black leather

⁵ like biofilm with mucilaginous marble on its surface covered by green spots. When the biofilm gets dry, the net of precipitated micro fortified mucilage mixed with soft mineral particles and crystals of calcium carbonate is visible (Fig. 6g, h).

Scanning electron micrographs document the structure of the biofilm (Fig. 7). Figure 7a shows a lateral view (cross section) of a biofilm with cyanobacterial filaments

¹⁰ (*Calothrix elsteri* and *Hassallia andreassenni*). A biofilm upper view (Fig. 7b, d) shows the structure of the cyanobacterial-microalgae community producing the mucilaginous micro fortified net of filaments with spots on its surface.

3.4 Inorganic compounds of biofilms

Thin sections, showing both biofilms and rock substrate (Fig. 8), provided information

- on various inorganic compounds associated with the soft tissue of the cyanobacterial – microalgal community. These inorganic compounds are represented by (1) allochthonous mineral grains that are overgrown and incorporated by biofilms and (2) calcareous spicules of different sizes ranging from 0.5 mm to 1 cm that are precipitated within the cyanobacterial-microalgal community.
- The rock substrate of biofilms is formed by subangular to subrounded pebbles to boulders of basaltic rock, which is dark-grey in colour, compact and usually with a microcrystalline porhyric texture. The rock is not homogenous, but contains numerous ball-like empty voids, which are often partly filled with feldspathoids (Fig. 8a). Crystals of plagioclase (feldspar group) and augite (pyroxene group) are easily recognizable in thin sections (Figs. 8a–c).

Biofilms are often partly covered with various mineral grains and rock fragments, but all specimens studied also contain these particles incorporated directly within soft cyanobacterial – microalgal filaments (Fig. 8a–c).



Mineral grains embedded within biofilms close to the basaltic rock surface are mainly angular to subangular crystal fragments of plagioclase and augite (Figs. 8b, c), i.e. the main mineral components of the basaltic rock substrate described above. In the upper part of biofilms, however, partly or fully incorporated grains of quartz occur, being typically rounded or partly rounded (Figs. 8a, b). One of the thin sections shows a

calcareous spicule in situ and mineral grains within the biofilm (Figs. 8c, d).

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The structure and morphology of calcareous spicules was studied on SEM (Fig. 9). The spicules (see also Methods) show an intensively worn surface (Fig. 9a), partial or intense recrystallization (Figs. 9a, b) and dissolution (Fig. 9b). Crystal facets on the surface and cleavage (crystallographic structural planes) in the interior of the spicules (Figs. 9a, b) are typical characteristics of calcium carbonate monocrystals.

A non-recrystallized superficial layer of microcrystalline calcite (e.g., Fig. 9b) shows the structure of parallel needle-like calcite microcrystals (Figs. 9d–f). Partial corrosion and dissolution of spicules show distinct layering of these needle-like microcrystals (Fig. 9d). The layered structure of even partly recrystallized spicules is confirmed in the ring-like structures with a cyanobacterial filament in the centre (Fig. 10).

The chemical composition of the studied calcareous spicules determined by FDS corresponds to pure $CaCO_3$. Following chemical composition, calcite and aragonite structural models were applied for the EBSD study focused on structural identification

- of the crystals forming the spicule. Structural identification of the studied specimen especially prepared for the EBSD study confirmed the absolute agreement between the recorded EBSD patterns and modelled patterns for calcite. The presence of aragonite was not confirmed. FSD images acquired for chemical and orientation contrasts (Fig. 10) show a layered structure especially visible in orientation contrast. This fea-
- ²⁵ ture reflects continual growing processes on layers with very similar crystallographic orientation. Absolute angular differences between individual layers are below 0.8°.



4 Discussion

4.1 Environmental properties

The Ulu Peninsula is a region of high limnological diversity that is related to differences in lake age, bedrock and altitude. The endorheic lakes under study are characterized by

⁵ a low content of major ions due to their volcanic bedrock and lower marine influence. In comparison with other lakes of this area, the two lakes show no specific lake water chemistry characteristics with moderate SRP and nitrate concentrations frequently below the detection limit (Nedbalová et al., 2013). High pH together with oxygen supersaturation recorded in lake 2 could be associated with high photosynthetic activity of the mats at the time of sampling.

Because water in either liquid or solid form has a large heat storage capacity, it acts as an important buffer to temperature change. Local climatic conditions of shallow freshwater lakes is the principal external factor controlling their ecological functionality. Lake 1 is frozen to the bottom with minimum daily mean temperatures between -12

and -10°C approximately eight-nine months per year. For most of the year, however, the temperature of the littoral and lake bottom is only from -2 to -4°C. In such conditions, a thin layer of water probably covers the surface of the littoral benthic community (the community can be metabolically active at temperatures of about -4°C) (Davey et al., 1992). The growing season, with liquid water at temperatures between 2 to 4°C, covers only two-three months.

In regards to heat balance, the studied shallow lakes are pond (wetlands) environments which freeze solid during the winter. This inevitability is a strong habitat-defining characteristic, which places considerable stress on resident organisms (Hawes et al., 1992; Elster, 2002). In summer, they must withstand drying in large parts of the lit-

toral zone due to a considerable drop in water level. In freezing and desiccation resistance studies of freshwater phytobenthos in shallow Antarctic lakes, several ecological measurements have recorded seasonal, diurnal, and year round temperature fluctuations and changes in water state transitions (e.g., Davey, 1989; Hawes et al., 1992, 1989; Hawes et al., 1



Hawes et al., 1999). In localities with steady moisture and nutrient supplies, the abundance and species diversity of algae is relatively high. However, as the severity and instability of living conditions increases (mainly due to changes in mechanical disturbances, desiccation-rehydration and subsequent changes in salinity), algal abundance

- and species diversity decreases (Elster and Benson, 2004). The speed at which water state can change between liquid, ice, and complete dryness, is one of the most important ecological and physiological factors of these lakes. Studies based on field or laboratory experiments have shown that some cyanobacteria and algae are able to tolerate prolonged periods of desiccation. It is also obvious that there are strain/species specific
- differences in the overwintering strategies, and also between strains/species inhabiting different habitats (Davey, 1989; Hawes et al., 1992; Jacob et al., 1992; Šabacká and Elster, 2006; Elster et al., 2008). The ice and snow which cover the endorheic lakes for about eight-nine months per year serve as a natural incubator which moderate potential mechanical disturbances and stabilise the thermal regimes of the lakes.

15 4.2 Biodiversity

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Patterns of endemism and alien establishment in Antarctica are very different across taxa and habitat types (terrestrial, freshwater or marine; Barnes et al., 2006). Environmental conditions, as well as dispersal abilities, are important in limiting alien establishment (Barnes et al., 2006). Antarctic microbial (cyanobacteria, algae) diversity is still poorly known, although recent molecular and ecophysiological evidence support a high level of endemism and speciation/taxon distinctness (Taton et al., 2003; Rybalka et al., 2009; de Wever et al., 2009; Komárek et al., 2011; Strunecký et al., 2012; Škaloud et al., 2013).

The floors of the studied lakes are covered with photosynthetic microbial mats composed of previously described species of heterocytous cyanobacteria, mostly *Calothrix elsteri* Komárek 2011 followed by *Hassallia andreassenni* Komárek 2011 and *Hassallia antarctica* Komárek 2011 (Komárek et al., 2011). They are co-dominated by a newly described species of green filamentous and richly branched algae *Hazenia*



broadyi Škaloud et Komárek 2013 (Ulotrichales, Chlorophyceae; Škaloud et al., 2013). All the previously mentioned recently described species have special taxonomic positions together with special ecology and are considered at present as Antarctic endemic species.

- ⁵ The black leather like biofilm with mucilaginous marble on its surface is covered by green spots. These macroscopic structures form mats a few mm thick consisting of the above mentioned species packed in mucilage glued together with fine material. The regular leather biofilm structure with distinct cyanobacterial-microalgal composition and incorporated mineral grains is a modern analogue of some of the oldest well-described
- ¹⁰ Archean stromatolites (sensu Allwood et al., 2006). During the limnological survey of the whole Ulu Peninsula (Nedbalová et al., 2013), this specific biofilm structure was observed only in these two endorheic lakes, although lakes with very similar morphometric and chemical characteristics are found in the area. The mat structure is thus apparently tightly linked to the species composition (Andersen et al., 2011).
- ¹⁵ The low abundance of benthic diatoms in the lakes is unusual, but not unprecedented as there are other areas in Antarctica where diatoms are scarce or absent (Broady, 1996; Wagner et al., 2004). The reason underlying the absence of diatoms is not immediately obvious, because diatoms are quite a common and frequently dominant component of microbial communities in most freshwater habitats of the Ulu Penin-
- ²⁰ sula, James Ross Island (Kopalová et al., 2013). Local geographical separation of lakes 1 and 2 together with founder effect may have precluded successful colonization by the subset of diatoms that are common in the surrounding freshwater habitats. Although it has long been held that diatoms are dispersed widely, some recent reports document very small scale microbial distributions and endemism (Kopalová et al., 2012; Kopalová et al., 2013).

4.3 Inorganic compounds of biofilms

Based on the character of the rock substrate and lake sediments it is suggested, that one of the main prerequisites for existence of this cyanobacterial-microalgal community



producing unusual biogenic calcite structures is; (1) flat and stable substrate in both lakes and (2) low sedimentation rate.

The substrate for biofilms is composed of boulders and pebbles of the stony littoral zone, petrographically corresponding to compact and massive basaltoids (Smellie et

al., 2008; Svojtka et al., 2009). Rounded or sub-rounded quartz grains that are incorporated ("trapped") within biofilms cannot originate from basaltic volcanic rocks forming the bottom of both lakes and substrate of the studied biofilms. This is evidenced by the petrographic character of the basaltoids, which do not contain any quartz. The presence of abraded quartz grains in lake 1 and 2 can be easily explained by wind transport (e.g., Shao, 2008).

The specific cyanobacterial-microalgal community described above can prosper in the two shallow endorheic lakes, because of low sedimentation rates resulting from minor water input. Low sedimentary input is the main necessary ecological parameter which facilitates the existence of this special microbial community. The community

- is, however, well adapted to seasonally elevated sedimentation rates coming from frequent and intense winds. During wind storms, the wind is carrying a relatively large amount of small mineral grains and rock microfragments (intense eolic erosion; e.g., Shao 2008 and references therein). These grains and particles are usually derived from erosion of the rocks either in the very close vicinity of the locality (weathering of
- ²⁰ basaltic rocks), but mainly come from remote locations where especially Upper Cretaceous marine sedimentary sequences are outcropping (Smellie et al., 2008; Svojtka et al., 2009). Even elevated amounts of mineral grains transported into the lake by wind do not stop the growth of cyanobacterial-microalgae biofilms, due to their ability of incorporating and "trapping" mineral grains within the living tissue.

This study has shown that inorganic substances precipitated by microbial lithogenetic processes are exclusively represented by calcite spicules. Precipitation of carbonate outside of microorganisms during photosynthesis as a mechanism of carbonate construction was described for many filamentous cyanobacterial species (Schneider and Le Campion-Alsumard, 1999). However, the biogenic calcite structures in both lakes



are quite different to any microbially mediated structures yet described from modern environments (Kremer et al., 2008; Couradeau et al., 2011). Although there are many lakes with thick mats and similar chemical characteristics on the Ulu Peninsula, the calcite spicules were found exclusively in the two endorheic lakes. We believe that their

- formation is linked to the specific photoautotrophic mats present in the lakes. From Figs. 6g, h it is clearly visible that the calcareous organosedimentary structures keep contours of viable photosynthetic microbial mat after desiccation or calcite spicules precipitation. More specifically, the co-dominance of a green microalga is unique since mats in Antarctic lakes are most frequently formed by filamentous cyanobacteria (Vin-
- ¹⁰ cent and Laybourn-Parry 2008). Therefore, we hypothesise that the more rapid photosynthesis rate of *Hazenia* in comparison with cyanobacteria may induce conditions necessary for carbonate precipitation in the lakes (Schneider and Le Campion-Alsumard, 1999; Vincent, 2000). However, abiotic precipitation of calcite is also possible. The segregation of Ca²⁺ and HCO₃⁻ between ice and the residual solution depend on the
- ¹⁵ freezing rate and hydrogen oxygen isotope fractionation (O'Neil, 1968; Žák et al., 2004). From our measurements we cannot clearly decide if the winter abiotic calcite precipitation accompany microbial lithogenetic processes.

It is striking that some calcite spicules exhibit recrystallization, forming spicules with the structure of calcite monocrystals. Determining the structure and material of precipitated inorganic substances brought another relevant question: "Do calcite spicules have fossilisation potential"? Microcrystalline calcite forming the recrystallized spicule is a typical material of calcite shells of fossil invertebrates (e.g., Vodrážka, 2009). Although calcite fossils may be partly or completely dissolved during diagenetical processes in the fossil record (e.g., Schneider et al., 2011; Švábenická et al., 2012), their preservation potential is relatively high. Therefore, we expect to find fossil and/or sub-

25 preservation potential is relatively high. Therefore, we expect to find fossil and/or sub fossil calcite spicules from the Quaternary lake sediments of the studied area.

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Table 1. Physico-chemical characteristics and chlorophyll *a* concentrations in lake water. Samples were collected from surface of lakes. ND – not determined, ANC – acid neutralization capacity, PN – particulate nitrogen, DP – dissolved phosphorus, PP – particulate phosphorus, SRP – dissolved reactive phosphorus, DOC – dissolved organic carbon, PC – particulate carbon, * – laboratory values.

Lake		Green 1		Green 2
Date		22.2.2008	5.1.2009	12.1.2009
Temperature	°C	3.5	ND	12.3
O ₂	mg L ⁻¹	13.1	ND	13.7
O ₂ saturation	%	98.7	ND	128.0
рН		7.9	7.4*	8.6
Conductivity (25°C)	µScm⁻¹	54	48*	97
ANC	mmol L^{-1}	236	246	455
Na ⁺	mgL^{-1}	4.7	5.9	12.5
K ⁺	mg L ⁻¹	0.24	0.29	0.60
Ca ²⁺	mg L ⁻¹	2.12	1.26	2.32
Mg ²⁺	mgL^{-1}	1.24	0.77	1.65
SO_4^{2-}	mgL^{-1}	1.74	1.33	2.60
CI	mgL^{-1}	5.3	5.1	10.6
NO ₃ -N	μ g L ⁻¹	< 5	11	< 5
NO ₂ -N	μ g L ⁻¹	0.6	0.2	0.1
NH ₄ -N	μ g L ⁻¹	6	< 5	< 5
PN	μ g L ⁻¹	20	50	73
DP	μ g L ⁻¹	7.8	20.2	30.4
PP	μ g L ⁻¹	4.6	5.9	11.7
SRP	μ g L ⁻¹	4.0	11.6	19.3
DOC	mgL^{-1}	1.25	1.13	2.17
PC	mgL^{-1}	0.13	0.41	1.33
Si	mgL^{-1}	1.45	0.87	2.85
chl a	μ g L ⁻¹	0.9	ND	6.0

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Figure 1. Location of lake 1 and 2 and air temperature measurements (ASW) in the Solorina Valley.





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Figure 2. Bathymetric parameters of lake 1 (a) and 2 (b) together with marked lines of water level and maximum extent of the photosynthetic microbial mat littoral belt in lake 1.





Figure 3. (a) Annual variation of daily mean water temperature in lake 1 (L1water) and annual variation of daily mean air temperature 2 m above ground in the Solorina Valley (SV air), **(b)** Diurnal temperature amplitudes in lake 1 (L1 water) and diurnal air temperature amplitudes in the Solorina Valley (SV air), respectively. **(c)** Daily mean global radiation at Mendel Station. All parameters measured from February 2009 to November 2010.



Figure 4. Relative frequency of hourly values of water temperature in lake 1 (a) and air temperature measured 2 m above ground in the Solorina Valley (b) from February 2009 to November 2010.





Figure 5. Occurrence of days with lake 1 water temperature (L1 water) higher than -4 °C complemented with monthly mean global solar radiation at Mendel Station both from February 2009 to November 2010.





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Figure 6. Photoautotrophic mats in lakes 1 and 2. (a) Rapid development of the mat in January 2009 (lake 1). The two photos show the mat at a one week interval, note the growth of gelatinous clusters of densely agglomerated filaments of the green alga *Hazenia broadyi*, (c, d) fully developed mats with mosaic-like structures on the surfaces of stones in the littoral zone of lake 2, (e, f) detail, (g) drying of the mat in the littoral zone leaves a characteristic structure on the surface of stones, (h) calcium carbonate spicule in situ (arrowed).





Figure 7. SEM macrographs showing the structure of the dried mat in the lakes. (a) Transversal section of the mat with visible cyanobacterial filaments, (b) surface structure of the mat, (c) general view of the surface structure of the mat with the net formed by mucilage (compare with Fig. 7d), (d) detail of the same mucilaginous structure.





Figure 8. Perpendicular thin sections of rock substrate overgrown by biofilms. Note that biofilms are partly detached from the surface of the rock due to complete drying of the sample. (a) Conspicuous U-shaped empty void (arrowed "2") near the surface of basaltic rock partly infilled with crystals of feldspathoids (tectosilicate minerals, arrowed "1"); empty void is bridged by biofilm (arrowed "3") with partly incorporated mineral clasts, represented by semi-rounded quartz grains (arrowed "4"), (b) rather thick biofilm with numerous incorporated mineral grains. Note that close to the rock substrate the angular grains of plagioclase (feldspar group) and augite (pyroxene group) dominate, being derived from basaltoids, whereas close to the surface rounded grains of quartz occur (arrowed), (c) in situ calcium carbonate spicule penetrating biofilm and surrounded by incorporated grains of feldspars (two arrows on the left) and pyroxene (arrow on the right), (d) close up of the same calcium carbonate spicule with a cyanobacterial filament in its centre.







Figure 9. SEM macrographs showing the morphology of partly recrystallized calcium carbonate spicules. Spicules were washed away from the living tissue and collected directly from the surface of biofilms, although residence time on the bottom cannot be determined. (a) Calcareous spicule showing intensively worn surface and complete recrystallization of the spicule interior. The spicule shows crystal facets on the surface and cleavage (crystallographic structural planes) in the interior (arrowed) – i.e. typical characteristics of calcium carbonate monocrystal, (b) detail of previous image; two parallel systems of deep furrows on the surface are crystallographic structural planes of calcite monocrystal; remnants of a superficial layer of microcrystalline calcite are, however, preserved in places on the surface of the crystal (arrowed), (c-f) poorly recrystallized spicule, formed mainly by microcrystalline calcite, (c) lateral view of the spicule, (d) detail of the surface showing corrosion of needle-like calcite microcrystals with distinct layering, (e) parallel needle-like calcite microcrystals on the surface of the central part of the spicule, (f) tops of parallel needle-like calcite microcrystals on the surface of the terminal part of the spicule; the view is perpendicular with respect to the previous macrograph.





Figure 10. FSD image of a transversely sectioned, partly recrystallized calcite spicule acquired in (a) chemical contrast, (b) orientation contrast.

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