

Microbial community changes along the Ecology Glacier ablation zone (King George Island, Antarctica)

Jakub Grzesiak¹ · Marek K. Zdanowski¹ · Dorota Górniak² · Aleksander Świątecki² · Tamara Aleksandrak-Piekarczyk³ · Katarzyna Szatraj³ · Joanna Sasin-Kurowska⁴ · Marta Nieckarz⁵

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Abstract In recent years, glacial surfaces have received much attention as microbial habitats of diverse photoautotrophic and heterotrophic cells. Supraglacial ecosystems are annually covered and uncovered by snow. The aim of this study is to investigate the microbial community response to changing environmental conditions in a transect following the receding snow line on the surface of Ecology Glacier (King George Island, Antarctica). Parameters of surface ice and cryoconite holes included chemical composition of ice and sediment, *Bacteria* diversity by denaturing gradient gel electrophoresis, microbial functional diversity (Biolog Ecoplates), and microbial counts (epifluorescence microscopy, colony-forming units). Data demonstrated profound differences between surface ice and cryoconite holes. Changing environmental factors along the transect influenced composition and abundance of the microbiocenosis in both habitat types. Several parameters correlated positively with distance from the glacier edge, including the cell morphotype Shannon index, chlorophyll

a, nitrogen, and seston concentrations. Suspended solid content positively correlated with microbial abundance and diversity. Nitrogen and phosphorus were limiting factors of microbial growth as amounts of organic nitrogen and phosphorus positively correlated with the cell numbers, fission rates, and photoautotroph contribution. Our findings indicate that microbial community shows a response in terms of abundance and diversity to exposure of the glacial surface as snow-cover melts. To our knowledge, this is the first study to recognize a microbial development pattern on a glacier surface in connection with the receding snow line. This may help better understand variability within supraglacial habitats, correct sampling procedures, and inform biocenotic development models.

Keywords Glacial ice · Cryoconite holes · Microbiocenosis · Microbial abundance · Snow line

Introduction

A significant part of the biosphere is permanently below 5 °C (Anesio and Laybourn-Parry 2012). This includes glaciers and ice sheets, which occupy 11 % of Earth's surface, and which comprise ~70 % of surface global freshwater (Shiklomanov 1993; Paterson 1994).

Glacial systems result from the perennial accumulation of snow. Their subsequent deformation leads to their flow to lower elevations, where they ultimately melt, sublimate or calve in the ablation zone. The glacial surface is annually covered by fresh snow whose melting during summer months exposes the ice below. As summer progresses, the ablation zone expands as the 'snow line' advances up the glacier (Hooke 2005; Anesio and Laybourn-Parry 2012; Edwards et al. 2014). The gradually exposed ice is

✉ Jakub Grzesiak
jgrzesiak@arctowski.pl

¹ Department of Antarctic Biology, Institute of Biochemistry and Biophysics, Polish Academy of Sciences, Pawińskiego 5a, 02-106 Warsaw, Poland

² Department of Microbiology, University of Warmia and Mazury, Oczapowskiego 1A, 10-719 Olsztyn, Poland

³ Department of Microbial Biochemistry, Institute of Biochemistry and Biophysics, Polish Academy of Sciences, Pawińskiego 5a, 02-106 Warsaw, Poland

⁴ Institute of Genetics and Biotechnology, Warsaw University, Pawińskiego 5a, 02-106 Warsaw, Poland

⁵ Department of Applied Microbiology, Faculty of Biology, Institute of Microbiology, University of Warsaw, Iłji Miecznikowa 1, 02-096 Warsaw, Poland

subjected to wind, solar radiation, and aeolian inputs of mineral and organic materials, resulting in its melting; the water produced circulates over and in the ice structure (Hodson et al. 2008). This water often carries a suspension that originates from wind-carried debris from local and from more distant sources (Zarsky et al. 2013). The dark dust deposit is known as cryoconite, and in sufficient quantities reduces ice surface albedo and accelerates melting. As the ice melts, a water-filled hole forms, into which the dark material sinks (Cameron et al. 2012).

Scientific interest in the biology of glaciers was limited up to the 1980s because they were considered to be ‘ice deserts’ with little biological activity (Edwards et al. 2014). However, Kohshima (1984) identified a simple glacier-associated food web. Hodson et al. (2008) and Anesio et al. (2009) then radically changed old beliefs by focusing on the importance of high microbial activity on glaciers in the global carbon cycle. Microbes, which have adapted to living under very low temperatures, dominate such cryosphere environments, and may comprise 4×10^{25} to 7×10^{29} cells in glacial ice globally, and up to 1×10^{26} bacterial cells in the uppermost 1–2 m of ablating glacier ice areas (Irvine-Fynn and Edwards 2012). Microbiological investigations of glacial habitats have revealed that their microbial communities are active in biogeochemical transformations because they can perform metabolic functions involved in the C, N, and P cycles (Hodson et al. 2007; Stibal et al. 2008a; Telling et al. 2011). This has important implications at local and global scales. Due to favorable irradiance conditions and access to water, photoautotrophs are more abundant in surface layers than deeper in the ice (Quesada and Vincent 2012). Cell numbers in this glacial ‘euphotic zone’ are of the same order of magnitude as in photic habitats in warmer climates (Edwards et al. 2014).

We report here on microbial communities on the surface of Ecology Glacier, a rapidly receding glacier on King George Island, South Shetland Islands, Antarctica. We hypothesize that gradual exposure of ice under melting snow is one of the major factors causing spatial variations of chemical composition, bacterial phylogenetic and microbial functional diversity, and abundance in surface ice and cryoconite holes. We analyzed the data using simple regression and principal component analysis (PCA) in order to identify environmental controls of microbial processes. By identifying biological and physical variables for microbial community development in supraglacial habitats, we aim to supplement the still-scarce knowledge of these disappearing habitats, and better inform microbiocenotic development models. We also quantitatively and qualitatively investigated the culturable component of the glacial microbial community.

Materials and methods

Sites and sampling

Ecology Glacier is situated at the western shore of Admiralty Bay, on King George Island, South Shetland Archipelago, Antarctica. It is subjected to annual surface snow melt like other glaciers in the vicinity (Braun and Gossmann 2002). The altitude and distance from glacier terminus of the transient snow line has been observed from early November. Samples were taken in December 2011/January 2012 when the snow line reached over 200 m a.s.l. and nearly 2 km from the glacier terminus. Ice and cryoconite material were collected aseptically at six sites on the surface of Ecology Glacier into chemically clean and sterile containers along a 1841-m transect running from the glacier terminus to the snow line at the top of the ablation zone (Table 1; Fig. 1). Ice samples were termed EI (Ecology Ice), with EI1 being the first point in the ablation zone, and EI5 the snow line point. Additionally, site EDI was selected below the first sampling point on a fragment of the glacier, which in previous years was an integral part of the glacier and at the sampling period was recognized as dead-ice (Schomacker 2008). Cryoconite samples were labeled EC2 to EC5, the number indicating the same area as the surface ice samples (no cryoconite holes were found in the dead-ice and EI1 areas).

During the 2011/2012 austral summer, ice from the glacier’s surface was crushed with an 70 % EtOH sterilized and deionized water-washed Tonar ice auger (158 cm long, 130 mm in diameter), collected using sterile plastic spatulas and placed into sterile plastic bags. The crushed ice was gathered from five points per sampling site, in an area of 100 m². Pooled cores totaled 3 kg per site. Cryoconite holes were drained of water and sediment with a 160-ml sterile plastic syringe, and the material was transported in 500-ml sterile bottles to a field laboratory and processed within 2 h. Five cryoconite holes per site were drained and pooled. A duplicate set of samples was stored at -20°C for further research in Poland.

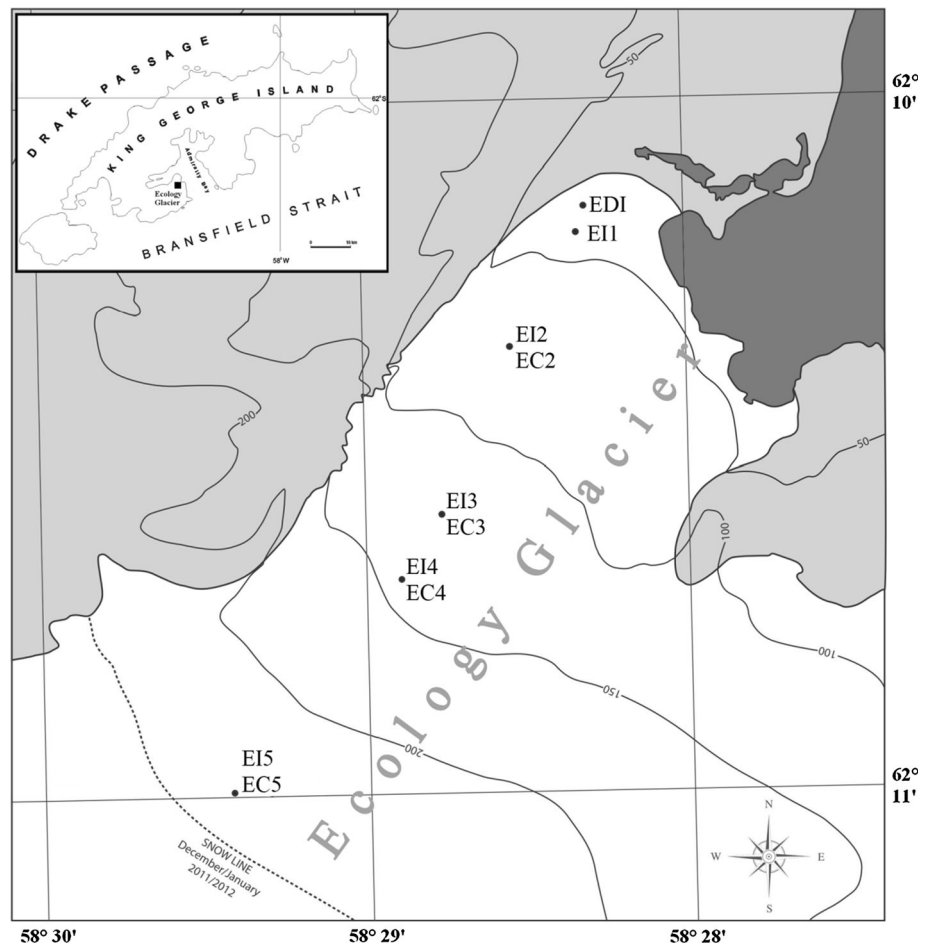
Measurement of ice and cryoconite components

Chloride, sulfate, nitrate, sodium, potassium, calcium, and magnesium ion concentrations were determined by high-pressure liquid chromatography (HPLC) in a Shimadzu Prominence modular HPLC model device. Phosphates, total phosphorus (after mineralization), total iron, and silicate contents were measured in a Shimadzu UV 1601 spectrophotometer, while nitrates were measured on an Epoll-Eco 20 spectrophotometer (Standard Methods 1980; Hermanowicz et al. 1999). Total organic carbon (TOC),

Table 1 Sampling points along transect from EDI ('dead ice') near the Ecology Glacier terminus to the highest part of the ablation zone (snow line)

Sample point	Latitude (S)	Longitude (W)	Distance from glacier terminus (m)	Elevation (m)	Description
EDI	62°10'09.3"	58°28'18.5"	0	12	Dead ice area, solid, blue glacier ice heavily covered with mineral debris. No cryoconite holes
EI1	62°10'11.6"	58°28'20.0"	79	35	Ablation zone, blue glacier ice heavily covered with mineral debris. No cryoconite holes
EI2; EC2	62°10'21.4"	58°28'32.6"	427	73	Ablation zone, castellated clear ice with mineral debris. Cryoconite holes present
EI3; EC3	62°10'35.8"	58°28'45.9"	914	125	Ablation zone, ice with very porous surface, minor mineral debris. Cryoconite holes present
EI4; EC4	62°10'41.4"	58°28'53.6"	1117	137	Ablation zone, ice with very porous surface, minor mineral debris. Cryoconite holes present
EI5; EC5	62°10'59.7"	58°29'25.6"	1841	235	Equilibrium line area, ice/firn with snow patches. Cryoconite holes present

EI, surface ice; EC, cryoconite holes

Fig. 1 Location of sampling points on Ecology Glacier surface

dissolved organic carbon (DOC), particulate organic carbon (POC), and total nitrogen contents were determined in a Shimadzu TOCV-CSH organic carbon analyzer with a

TNM-1 total nitrogen analyzer attachment. Particulate content in water was determined by filtering the sample through a combusted Whatman GF/C glass fiber filter

(nominal pore size, 1.2 μm) which was then dried at 105 °C; organic matter content was measured as that amount lost on ignition after combustion at 550 °C. Chlorophyll *a* and pheophytin concentrations were calculated after the Lorenzen equation after extraction in acetone (at 4 °C) and spectrophotometry. A DOC spectrum was determined in a quartz cuvette at 260 nm in a Shimadzu UV-1601 bichannel scanning spectrophotometer (Shinichi et al. 2004). The spectrum was corrected against a demineralized water background. SUVA (specific UV absorbance) was calculated as the relative content of aromatic matter ($=\text{Abs}_{260} \times 1000 \text{ DOC}^{-1}$).

Sample preparation for microbiology

Ice samples were melted in a refrigerator (4 °C) before processing for microbiological analyses. Cryoconite material was shaken gently on a universal shaker (Premed, model 327) (120 rpm, 20 min, 5 °C). Suspensions were then returned to the refrigerator for 10–20 min to allow larger particles to settle. Aliquots of 1, 0.5, and 0.1 ml were spread-plated on R2A agar (R2A) and eight-fold diluted R2A agar (R2A/8) (Biorcorp). Further, 300 ml of melted ice was filtered (47 mm polycarbonate filter, 0.2 μm pores), after which the filter was placed in a 50-ml sterile plastic tube and shaken with 20 ml of melted ice to concentrate the microbes by resuspension in an appropriate medium. The suspension was plated onto R2A agar, and eight-fold diluted R2A agar (R2A/8) as described above. The remainder was frozen for further analysis. Inoculated plates were incubated in darkness at 4 °C for 6 weeks. After 6 weeks incubation, cells were collected from R2A and R2A/8 plates by adding 2 \times 5 ml of sterile 0.9 % saline to the plate and suspending the colonies with a sterile plastic spreader (plate wash). The plate wash suspensions were collected into 15-ml sterile plastic tubes and homogenized by repeated pipetting (Ellis et al. 2003).

Microbial abundance

Sub-samples of melted ice were fixed with buffered formalin to a final concentration of 1 % for total counts (TC). Counts were determined in 5 ml of melted glacier ice and 1 ml of cryoconite water/sediment suspension by epifluorescence microscopy with 4,6-diamidino-2-phenylindole (DAPI) on black Nuclepore[®] polycarbonate 0.2- μm filters (Porter and Feig 1980), under a Nikon E-200 microscope fitted with a 100-W Hg lamp and 100 \times CFI 60 oil immersion objective, a digital DS-5Mc-U1 Cooled Camera Head, and a filter block of EX 330-380, DM 400, BA 420. Images of fields were analyzed in the Nikon NIS Elements BR 2.30 and MultiScan v.14.02. A minimum 400 cells in 20 fields per sample was counted automatically in the

image analysis system. Cell volumes were evaluated by image analysis (Sieracki et al. 1985; Świątecki 1997).

Photoautotrophs including cyanobacteria and photoautotrophic eukaryotes were counted under blue (450–490 nm, B-2A Nikon filter) and green (510–560 nm, G-2A Nikon filter) light (Putland and Rivkin 1999). Cell morphotype diversity was described through the frequency of cocci, rods, and curved forms in five volume classes (<0.1 ; 0.1–0.2; 0.2–0.5, 0.5–1.0; $>1 \mu\text{m}^3$), with a modified Shannon index (Gurienowich 1995; Nübel et al. 1999). Culturable microbial abundance was performed by CFU enumeration after 42 days incubation on R2A agar.

Denaturing gradient gel electrophoresis analysis

Water samples and cryoconite water–sediment mix (500 ml) were vacuum filtered through Millipore GTTP 0.2- μm pore-sized (47 mm diameter) white polycarbonate membranes in a sterile filter holder (Nalgene). Total DNA was extracted from the membranes in the UltraClean[™] Water DNA Isolation Kit (MoBio, Carlsbad, CA, USA) in accordance with the manufacturer's protocol. Total DNA from plate wash samples was extracted using Bacterial and Yeast Genomic DNA Purification Kit (EURx Ltd.) according to the manufacturer's protocol. DNA quality and yield were measured with a NanoVue[™] spectrophotometer (GE Healthcare Life Science, Germany). Extracted DNA was stored at $-20 \text{ }^\circ\text{C}$ until required. DNA molecular weight and quantity was checked by electrophoresis in a 0.8 % (w/v) agarose gel in 0.5 % TBE (Tris-borate-ethylenediamine-triacetate, pH 8.3) buffer, and stained with 0.9 $\mu\text{g ml}^{-1}$ ethidium bromide (Sambrook et al. 1989). A molecular size marker (1-kb ladder) was used as the reference.

Denaturing gradient gel electrophoresis (DGGE) analysis of the extracted DNA was performed with a D-Code Universal Mutation Detection System (Bio-Rad Laboratories, USA). One-microliter aliquots (~ 5 –10 ng in undiluted form) of each DNA sample was amplified in a 50- μl polymerase chain reaction mix (PCR) comprising 5 μl 10 \times buffer (Sigma Aldrich Co.), 6 μl of 25 mM MgCl_2 (Sigma Aldrich Co.), 1.2 μl of 20 mg ml^{-1} BSA (Sigma Aldrich Co.), 0.4 μl of 25 mM dNTP (Sigma Aldrich Co.), 0.5 μl of each primer (20 μM each), 0.2 μl of 5U μl^{-1} *Taq* DNA polymerase (Sigma Aldrich Co.), and 35.2 μl of PCR-grade water. Primers were F341 with a GC clamp (5'-CGC CCG CCG CGC CCC GCG CCC GTC CCG CCG CCC CCG CCC GCC TAC GGG AGG CAG CAG-3') complementary to position 341 to 357 (*Escherichia coli* numbering), and R907 (CCG TCA ATT CMT TTG AGT TT) complementary to positions 926–907 (Muyzer et al. 1993, 1998). Samples were loaded on a 6 % acrylamide gel with a denaturing gradient of 35–70 % (where 100 % denaturant is

7 M urea and 40 % formamide). Gels were run at 60 V for 17 h at 60 °C. Electrophoretic products were stained by gentle agitation of the gel for 30 min in 100 ml of $1 \times$ TAE containing 5 μ l 1:10,000 dilution of SYBR Gold nucleic acid stain (Invitrogen, Life Technologies, UK) in DMSO. DGGE banding patterns were visualized by UV transillumination and photographed in the Gel Doc 2000 gel documentation system (Bio-Rad Laboratories, USA). DGGE gel images were analyzed by ‘Quantity One’ software in the GelDoc gel documentation system (Bio-Rad Laboratories, USA). Gel bands were identified using GelCompar software to create a presence-absence matrix (Crump and Hobbie 2005). Each band represents a bacterial operational taxonomic unit (OTU). The presence or absence of a band in each line was converted to a binary matrix to access data for statistical analysis.

Functional diversity of total and cultivated microbiocenosis

Filter concentrated ice, cryoconite, and plate wash suspensions were adjusted to optical transmittance of 0.9 by the addition of sterile 0.9 % saline. Each suspension was then transferred aseptically to the Biolog Ecoplate (Biolog Inc., Hayward, CA, USA). Plates were incubated in darkness at 4 °C; color development was measured at 590 nm in a microplate reader (OmniLog), and cell respiration was measured kinetically by determining the colorimetric reduction of tetrazolium dye. Data were collected approximately twice weekly for 65 days. The Biolog Ecoplate assesses the ability of a mixed microbial community to utilize 31 carbon compounds presented as sole carbon sources. Before determining the absorbance in ‘OmniLog’ arbitrary units at different read times, samples were first blanked against the $T = 0$ reading and then against the plate’s respective control. Positive values were scored as the community’s ability to utilize a given carbon source. A community’s metabolic diversity was estimated from the number of substrates utilized. To compare the effect of a specific treatment, substrate utilization data were also subdivided into five categories representing different substrate types (carbohydrates, carboxylic and acetic acids, polymers, amino acids, and amines) (Weber and Legge 2009).

Statistics

Simple regression analysis between biological and environmental factors was carried out in STATISTICA v.9 (StatSoft). A principal component analysis (PCA) of physico-chemical and microbiological data as well as bacterial phylogenetic community structure was conducted in a statistical package for Windows, v. Canoco 4.5 (after ter Braak and Šmilauer 2002).

Results

Sampling site characteristics and environmental factors

Chlorophyll *a* contents varied in surface ice samples from 0.53 (EDI) to 3.53 μ g l^{-1} at EI4 (Table 2). A similarly increasing trend was evident in cryoconite holes, with 3.74 (EC2) to 52.18 μ g l^{-1} in EC5. Pheophytin and total chlorophyll levels in cryoconite holes also followed this pattern, showing the lowest values in EC2 (8.23 and 11.98 μ g l^{-1} , respectively), and highest in EC5 (123.26 and 175.43 μ g l^{-1} , respectively). The highest values in surface ice samples were detected in the dead-ice area (8.49 and 9.82 μ g l^{-1}). The highest amounts of total particulates were measured in the higher elevation points of the ablation zone for surface ice (EI4—769.20 mg dry wt. l^{-1}), and cryoconite holes (EC5—9213 mg dry wt. l^{-1}), comprising in all but one case (i.e., DIE) mostly organic material. pH ranged from 6.10 (EI3) to 7.15 (EI2) for ice samples, and 6.15 (EC2) to 7.00 (EC4) for cryoconite suspension.

Surface ice TOC, DOC, and POC levels peaked at EI4 (4.90 mg l^{-1}), EI3 (3.76 mg l^{-1}), and EI2 (3.78 mg l^{-1}) respectively; the lowest values were recorded noted at EI5 (1.23 mg l^{-1}), EDI (0.38 mg l^{-1}), and EI3 (1.10 mg l^{-1}). TOC in cryoconite holes ranged from 3.10 to 9.18 mg l^{-1} , and was composed mostly of DOC (EC3–EC5). The aromatic character of DOC (SUVA) was greatest at points closest to the glacier’s terminus (EDI -17.94 L mgC $^{-1}$ m $^{-1}$) for surface ice, and EC2 (9.92 L mgC $^{-1}$ m $^{-1}$) for cryoconite holes. Low amounts of inorganic dissolved nitrogen compounds were found along the transect, with highest levels of 0.09 mg l^{-1} for NO $_2^-$ and 0.05 mg l^{-1} for NH $_4^+$. Nitrogen sources were mostly organic, and varied between 0.15 (EDI, EI3) to 0.45 mg l^{-1} (EI5) for ice, and 0.16 (EC2) to 3.20 mg l^{-1} (EC5) for cryoconite samples. Total phosphorus amounts were low, ranging from 0.03 to 0.07 mg l^{-1} for ice samples and 0.07 to 0.21 mg l^{-1} for cryoconite suspension. In points near the snow line (EI4, EI5; EC4, EC5), phosphorus sources were mostly organic, while at near the glacier terminus (EDI, EI1) they were inorganic. Sodium and potassium levels varied along the transect, reaching highest values in EI3 (3.11 mg l^{-1}) and EI2 (0.34 mg l^{-1}) for ice samples, and EC3 (3.99 and 0.91 mg l^{-1} , respectively) for cryoconite holes. The concentrations of other elements are shown in Table 2.

Microbial abundance and diversity

Total counts in ice samples ranged from 8.1×10^3 (EI1) to 1.01×10^5 (EI4) cells ml $^{-1}$, while that in cryoconite holes varied between 1.34×10^5 (EC2) and 1.96×10^6 (EC4) cells ml $^{-1}$ (Table 3). CFU counts on R2A agar ranged from

Table 2 Values of chlorophyll and pheophytin concentrations, particulates, pH, carbon

Sampling points	EDI	EI 1	EI 2	EI 3	EI 4	EI 5	Mean	EC 2	EC 3	EC 4	EC 5	Mean
Chlorophyll a ($\mu\text{g/l}$)	1.34	0.53	0.75	0.96	3.53	2.57	1.61	3.74	6.15	9.84	52.18	17.98
Pheophytin ($\mu\text{g/l}$)	8.49	1.49	1.35	2.93	4.48	4.02	3.79	8.23	11.63	10.45	123.26	38.39
Total chlorophyll ($\mu\text{g/l}$)	9.82	2.02	2.10	3.89	8.01	6.59	5.41	11.98	17.78	20.28	175.43	56.37
Total seston (mg dry wt./l)	27.50	54.00	144.80	42.00	769.20	515.60	258.85	514.67	135.00	1305.60	9213.00	2792.07
Mineral seston (mg dry wt./l)	17.50	12.80	17.20	12.00	56.80	40.00	26.05	48.00	32.00	107.20	535.00	180.55
Organic seston (mg dry wt./l)	10.00	41.20	127.60	30.00	712.40	475.60	232.80	466.67	103.00	1198.40	8678.00	2611.52
pH	6.96	7.00	7.15	6.10	6.66	6.65	6.75	6.15	6.98	7.00	6.74	6.72
TOC (mg/l)	2.35	2.33	4.47	4.85	4.90	1.23	3.36	5.96	9.18	3.10	3.95	5.55
DOC (mg/l)	0.38	1.22	0.69	3.76	2.25	0.92	1.54	0.79	5.94	1.91	2.42	2.77
POC (mg/l)	1.97	1.11	3.78	1.10	2.65	0.30	1.82	5.18	3.25	1.19	1.53	2.79
SUVA	17.94	8.14	10.74	1.65	2.04	5.42	7.66	9.92	4.95	2.41	4.30	5.40
NH_4^+ (mg/l)	0.01	nd	0.05	0.01	0.01	0.01	0.02	0.01	nd	nd	0.01	0.01
NO_2^- (mg/l)	nd	0.09	nd	nd	nd	nd	0.02	nd	nd	nd	nd	–
NO_3^- (mg/l)	nd	nd	nd	nd	nd	nd	–	nd	nd	nd	nd	–
Total nitrogen (mg/l)	0.17	0.29	0.28	0.16	0.18	0.46	0.26	0.18	0.44	0.34	3.21	1.04
Organic nitrogen (mg/l)	0.15	0.20	0.23	0.15	0.17	0.45	0.23	0.16	0.44	0.33	3.20	1.03
PO_4^{3-} (mg/l)	0.06	0.02	0.02	0.02	0.02	0.02	0.03	0.02	0.02	0.02	0.01	0.02
Total phosphorus (mg/l)	0.06	0.03	0.04	0.03	0.06	0.07	0.05	0.07	0.13	0.21	0.20	0.15
Organic phosphorus (mg/l)	nd	0.01	0.02	0.01	0.04	0.04	0.02	0.05	0.11	0.19	0.19	0.14
Na^+ (mg/l)	2.53	2.34	1.56	3.11	1.82	2.09	2.24	3.81	3.99	3.21	3.70	3.68
K^+ (mg/l)	0.30	0.09	0.34	0.18	0.25	0.13	0.22	0.45	0.91	0.22	0.28	0.47
Ca^{2+} (mg/l)	1.12	0.55	0.28	0.87	0.63	0.33	0.63	0.41	0.26	0.51	0.61	0.45
Mg^{2+} (mg/l)	4.97	0.79	1.39	4.45	3.58	2.07	2.88	3.46	2.62	5.28	7.59	4.73
Cl^- (mg/l)	0.67	4.41	1.07	1.43	0.42	0.67	1.45	3.30	2.82	2.20	2.64	2.74
SO_4^{2-} (mg/l)	0.63	0.41	0.08	0.25	0.17	0.14	0.28	0.41	0.75	0.37	0.49	0.51
SiO_2 (mg/l)	0.98	0.51	0.63	0.47	1.61	0.59	0.80	0.68	0.35	1.00	2.18	1.05
Total iron (Fe) (mg/l)	0.14	0.01	0.29	0.18	0.05	0.07	0.12	0.06	0.08	0.70	3.43	1.06
CaCO_3 (mg/l)	12.39	1.97	3.48	11.11	8.93	5.15	7.17	8.64	6.53	13.16	18.93	11.82

TOC, total organic carbon; DOC, dissolved organic carbon; POC, particulate organic carbon; nitrogen, phosphorus, and other mineral contents in samples used for microbiological analysis; EDI, EI1–5, ice samples; EC2–5, cryoconite suspension

2.14×10^3 to 1.08×10^4 ml^{-1} . The CFU/TC ratio was highest in ice samples near the glacier edge, reaching 60.87 % (EI1). The percentage of culturable cells was much lower in cryoconite holes, with a maximum of 2.03 %.

Average cell volume determined by EFM was higher in cryoconite samples, with a maximum of $0.06 \mu\text{m}^3$ compared to ice samples (max. $0.03 \mu\text{m}^3$). The morphotype diversity index (Shannon index) exhibited the highest values in ice samples nearest the snow line (0.73–EI5), and lowest at the glacier terminus (0.34–EI1, 0.35–EDI). In cryoconite holes, this value was lowest and highest in neighboring sampling sites (0.43–EC3; 0.94–EC4). The highest and lowest dividing cell counts (DC) followed the TC trend, albeit 10-fold lower. In ice and cryoconite samples, the highest DC/TC ratios were noted in points nearest the snow line (20.1 %—EI5, 31.6 %—EC5); the lowest DC/TC were at the adjacent site (1.7 %—EI4, 5.4 %—EC4).

Photoautotroph abundance in ice samples (PHAC) followed the pattern of TC, with the highest numbers at EI4 (9.13×10^3 cells ml^{-1}) and lowest at the edge of the glacier (ED1— 9.5×10^2 cells ml^{-1}). In cryoconite holes, high PHAC numbers were noted in the samples nearest the glacier terminus (4.72×10^4 cells ml^{-1} —EC2) and near the snow line (3.94×10^4 cells ml^{-1} —EC5), dropping slightly in between (1.72×10^4 cells ml^{-1} —EC4). The average length of photoautotrophic cells (PHAC av. l.) per ice sample varied from 3.18 to 15.27 μm , and increased gradually with distance from the glacier edge. Photoautotrophic cell length in cryoconite holes ranged from 5.05 (EC3) to 12.06 μm (EC2).

Phylogenetic community diversity in terms of operational taxonomic units (OTU—DGGE bands) was lower on average in the ice samples, with a maximum of 12 (EI4). OTU numbers in cryoconite holes varied from 11 to 19. In the R2A medium plate wash, a max. of 7 and 9 OTUs were

Table 3 Values of microbiological parameters of native and cultivated microbes

Sampling points	EDI	EI 1	EI 2	EI 3	EI 4	EI 5	Mean	EC 2	EC 3	EC 4	EC 5	Mean
TC ($\times 10^4$ ml ⁻¹)	0.83 ± 0.13	0.81 ± 0.29	3.10 ± 0.52	2.21 ± 0.15	10.1 ± 1.49	2.59 ± 0.36	3.27	13.6 ± 4.32	53.3 ± 11.6	196 ± 16.3	16.0 ± 3.94	69.73
CFU _{R2A} ($\times 10^3$ ml ⁻¹)	3.99	4.92 ± 2.72	3.46 ± 1.72	2.14 ± 1.37	3.98 ± 0.41	2.46 ± 1.10	3.49	2.41 ± 0.43	10.8 ± 1.14	5.77 ± 2.39	3.01 ± 0.98	5.50
CFU _{R2A} /TC (%)	48.07	60.87	11.15	9.69	3.92	9.50	23.87	1.77	2.03	0.29	1.88	1.49
PHAC ($\times 10^3$ ml ⁻¹)	0.95	0.97	3.95	3.89	9.13	6.08	4.16	47.20	17.40	17.20	39.50	30.33
PHAC av.1 (µm)	3.18	7.88	8.44	13.37	15.27	13.07	10.20	12.60	5.05	11.66	10.29	9.90
V (µm ³)	0.02	0.021	0.018	0.038	0.031	0.034	0.03	0.067	0.025	0.067	0.039	0.05
Shannon index	0.359	0.348	0.496	0.569	0.662	0.736	0.53	0.768	0.431	0.94	0.593	0.68
DC ($\times 10^3$ ml ⁻¹)	0.95	0.87	1.3	1.3	1.74	5.21	1.90	13.60	36.10	107.00	50.60	51.83
DC/TC (%)	11.3	10.7	4.2	5.9	1.7	20.1	8.98	10.0	6.8	5.4	31.6	13.45
PHAC/TC (%)	11.4	11.9	12.7	17.6	9.0	23.4	14.33	34.7	3.2	0.8	24.6	15.83
DGGE _{NATIVE} (OTUs)	10	5	8	11	12	10	9.33	16	11	19	11	14.25
DGGE _{R2A} (OTUs)	4	3	3	6	5	7	4.67	9	8	4	8	7.25
DGGE _{R2A/8} (OTUs)	6	6	6	6	9	6	6.50	7	6	7	7	6.75
EcoPlate _{NATIVE} (number of positive responses)	2	3	6	1	15	6	5.50	26	18	13	18	18.75
EcoPlate _{R2A} (number of positive responses)	0	14	16	3	20	22	12.50	17	16	20	23	19.00
EcoPlate _{R2A/8} (number of positive responses)	20	21	11	26	23	25	21.00	22	21	22	24	22.25

TC, total microbial count; CFU_{R2A}, colony-forming units on R2A agar; PHAC, photoautotrophic cell count; PHAC av. 1., average length of photoautotrophic cells, morphotype diversity index (Shannon index); V, cell volume; DC, dividing cell count; DGGE_{NATIVE/R2A/R2A/8}, phylogenetic diversity of native and cultivated samples given in operational taxonomic units (OTUs); EcoPlate_{NATIVE/R2A/R2A/8}, functional diversity given in positive response numbers on Biolog Ecoplates

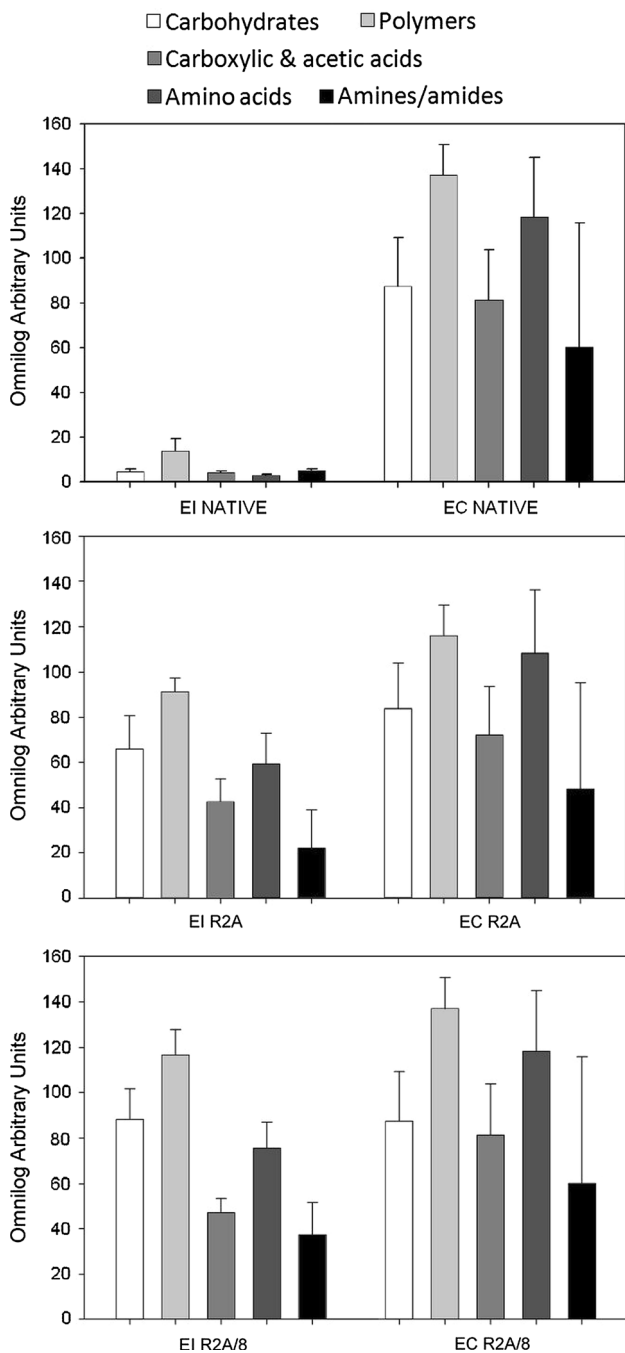


Fig. 2 Well color development calculated from Omnilog Arbitrary Unit values of substrate utilization on Biolog EcoPlates by glacial microbial communities. Substrates were divided into five categories: carbohydrates ($n = 10$), polymers ($n = 4$), carboxylic and acetic acids ($n = 9$), amino acids ($n = 6$), and amines/amides ($n = 2$). *EI*, surface ice samples ($n = 6$); *EC*, cryoconite samples ($n = 4$); *NATIVE*, samples applied directly to EcoPlates; *R2A*, samples incubated on full strength R2A agar (plate wash); *R2A/8*, samples incubated on 8-fold diluted R2A agar (plate wash)

determined in the ice and cryoconite samples, respectively. The eight-fold diluted R2A medium yielded similar results. Functional diversity in Biolog EcoPlates was low in the ice

samples, ranging from 1 positive response (*EI3*) to 15 (*EI4*), and relatively high in cryoconite holes (13—*EC4*, 26—*EC2*). Diluted R2A was superior to full strength R2A in recovering the functional potential of cultivable microbiota on the glacier (Table 3).

Substrate-utilizing activities were very low in surface ice samples compared to the cryoconite samples, and the cultivable fraction of the microbial community (Fig. 2). Polymers were the most actively metabolized group in native ice samples, specifically Tween 40 and Tween 80. D-cellobiose, D-malic acid, pyruvic acid methyl ester, α -D-lactose, glycogen, glucose-1-phosphate, putrescine, and itaconic acid were also frequently utilized (data not shown). The native cryoconite microbial community displayed a broad range of metabolic activities, with amino acids being the preferred substrate type. Compounds such as itaconic acid, L-asparagine, D-mannitol, L-serine, D-galactonic acid γ -lactone, L-arginine, pyruvic acid methyl ester, putrescine, N-acetyl-D-glucosamine, D-malic acid, and Tween 40 were the most frequently used substrates. Responses in the EcoPlate of the R2A plate wash-obtained surface ice microbiota (*EI R2A*) displayed a pattern similar to that of the respective native samples, but were much more intensely developed. Polymers remained the favorable carbon source, but carbohydrates and amino acids were also popular. The most marked color changes were seen in wells containing D-mannitol, D-cellobiose, α -D-lactose, L-asparagine, glycogen, Tween 40, itaconic acid, α -cyclodextrin, or N-acetyl-D-glucosamine. Cryoconite microbes cultivated on R2A agar also showed higher tetrazolium dye conversion rates than did the native samples, with polymers being the group they utilized most frequently, followed by amino acids and carbohydrates. D-mannitol, L-asparagine, itaconic acid, L-arginine, L-serine, α -cyclodextrin, Tween 40, α -D-lactose, D-galacturonic acid, and D-cellobiose were the preferred carbon sources. Plate wash from diluted R2A medium yielded the highest functional diversity; patterns of substrate utilization were the same in the full strength R2A plate wash, but the tetrazolium dye conversion rates were substantially higher (Fig. 2).

Statistics

The PCA of bacterial phylogenetic community structure in surface ice and cryoconite holes was based on the occurrence of DGGE bands (OTUs) in each of the sampling points (Fig. 3); vectors highlight the position and relative determining strength of the bands in the analysis. A phylogenetically diverse and unique community was detected in cryoconite holes *EC2* and *EC4*. A poor microbial community diversity was detected in surface ice samples

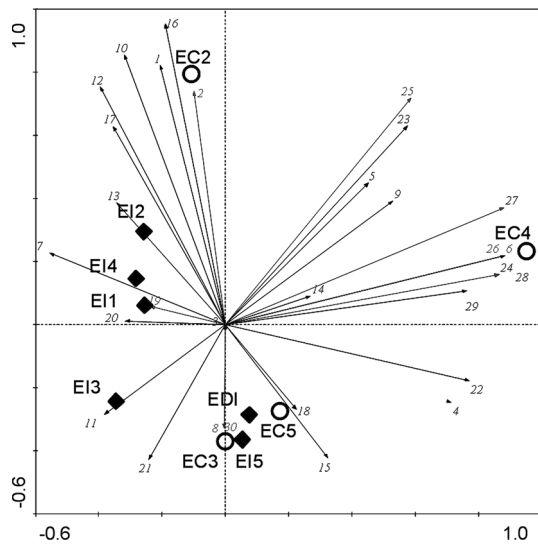


Fig. 3 Principal component analysis of bacterial phylogenetic community structure (based on OTUs relations) in surface ice (black diamonds) and cryoconite samples (circles)

(EI1–EI4), defined by a limited number of bands. A similar phylogenetic community structure was also revealed in samples EC3, EC5, EDI, and EI5, with EDI being the dead-ice samples and EI5 the snow line samples, 1.8 km apart.

A simple regression analysis between certain biological and environmental factors has been conducted. Physical, chemical, or biological factors that showed significant ($p < 0.05$) correlation have been displayed in Table 4 for

surface ice and Table 5 for cryoconite holes. Distance from the glacier edge positively correlated with the organic phosphorus concentration ($p < 0.05$) photoautotroph cell average length ($p < 0.05$) and Shannon index ($p < 0.01$) in surface ice samples (Table 4) and with chlorophyll *a*, nitrate, organic and total nitrogen, and seston contents in cryoconite holes ($p < 0.05$) (Table 5). Total microbial count (TC) positively correlated in surface ice samples with functional diversity ($p < 0.01$), nitrate, seston, SiO₂ and chlorophyll *a* ($p < 0.05$) levels. In cryoconite samples, it displayed a negative correlation with Na and Cl ions ($p < 0.05$). The phototroph percentage correlated positively ($p < 0.05$) with organic nitrogen levels in surface ice, in cryoconite samples with functional diversity and relative content of aromatic matter (SUVA) ($p < 0.01$) but negatively with pH ($p < 0.01$). In surface samples, the dividing to total cell ratio displayed negative correlations with total organic carbon ($p < 0.01$) and nitrate concentrations ($p < 0.05$) but a positive one with organic nitrogen content ($p < 0.05$). The CFU/TC ratio showed only a negative correlation with the distance from glacier edge ($p < 0.05$) in surface ice but several positive [with POC ($p < 0.01$); SUVA and Cl⁻ ($p < 0.05$)] as well as negative [with Ptot, Porg ($p < 0.01$); NO₃⁻ ($p < 0.05$)] in cryoconite hole samples. The phylogenetic diversity by DGGE correlated positively with Mg²⁺ and CaCO₃ contents in the surface ice samples, while in cryoconite samples only with morphotype Shannon index. The functional diversity by the EcoPlate assay displayed positive correlations with seston ($p < 0.01$), chlorophyll *a*, SiO₂ and

Table 4 Correlation coefficients (* $p < 0.05$, ** $p < 0.01$) between ice samples' biological parameters

Surface ice	Distance	TC	PHAC/TC	PHAC av.l	DC/TC	Shannon index	DGGE	EcoPlate
PHAC av.l	0.814*	0.669	0.317	1.000	-0.188	0.860*	0.504	0.565
CFU/TC	-0.756*	-0.618	-0.324	-0.773*	0.271	-0.868*	-0.704	-0.515
Shannon index	0.974**	0.579	0.545	0.860*	0.119	1.000	0.644	0.552
EcoPlate	0.416	0.950**	-0.343	0.565	-0.387	0.552	0.408	1.000
Chlorophyll <i>a</i>	0.686	0.808*	0.034	0.613	0.006	0.746	0.696	0.825*
Total seston	0.715	0.858*	0.028	0.705	-0.057	0.784*	0.529	0.913**
Mineral seston	0.661	0.853*	-0.028	0.626	-0.045	0.732	0.575	0.906**
TOC	-0.076	0.555	-0.464	0.392	-0.943**	0.117	0.379	0.337
SUVA	-0.674	-0.537	-0.281	-0.958**	0.229	-0.699	-0.336	-0.404
NO ₃ ⁻	0.231	0.778*	-0.338	0.647	-0.841*	0.419	0.574	0.578
Organic nitrogen	0.701	-0.116	0.787*	0.257	0.771*	0.581	-0.047	0.075
Organic phosphorus	0.804*	0.701	0.248	0.716	0.111	0.838*	0.342	0.810*
Mg ²⁺	-0.009	0.134	-0.103	-0.025	-0.183	0.045	0.766*	-0.084
SiO ₂	0.085	0.837*	-0.599	0.224	-0.452	0.229	0.565	0.818*
CaCO ₃	-0.009	0.134	-0.103	-0.025	-0.183	0.045	0.766*	-0.084

TC, total microbial count; CFU/TC, colony-forming units on R2A agar to total microbial count ratio; PHAC, photoautotrophic cell count; PHAC Av. L., photoautotrophic cell average length; PHAC/TC, photoautotrophic cell count to total microbial count ratio; DC/TC, ratio of dividing cell count to total count, morphotype diversity index (Shannon index); DGGE_{NATIVE}, phylogenetic diversity of native samples by operational taxonomic units (OTUs); EcoPlate, functional diversity of native samples by positive response numbers on Biolog Ecoplates) and distance from glacier terminus and chosen biological parameters and non-biological factors

Table 5 Correlation coefficients ($*p < 0.05$, $**p < 0.01$) between cryoconite hole biological parameters

Cryoconite holes	Distance	TC	CFU/TC	PHAC/TC	PHAC av.l	Shannon index	DGGE	EcoPlate
PHAC/TC	-0.531	-0.713	0.745	1.000	0.432	0.012	-0.009	0.977**
DGGE	-0.417	0.662	-0.157	-0.009	0.691	0.953*	1.000	-0.153
Chlorophyll <i>a</i>	0.915*	-0.014	-0.587	-0.145	0.077	-0.225	-0.502	-0.198
Total seston	0.890*	-0.015	-0.574	-0.089	0.152	-0.170	-0.456	-0.155
DOC	0.120	-0.197	0.087	-0.512	-0.995**	-0.789	-0.635	-0.322
POC	-0.823	-0.758	0.978**	0.854	0.050	-0.174	-0.037	0.917*
SUVA	-0.678	-0.767	0.879*	0.972**	0.269	-0.081	-0.033	0.989**
NO ₃ ⁻	0.899*	0.537	-0.914*	-0.450	0.312	0.254	-0.005	-0.574
Total nitrogen	0.895*	-0.103	-0.518	-0.109	0.012	-0.313	-0.580	-0.145
Organic nitrogen	0.896*	-0.102	-0.520	-0.112	0.009	-0.315	-0.582	-0.147
Total phosphorus	0.830	0.754	-0.981**	-0.843	-0.036	0.179	0.037	-0.909*
Organic phosphorus	0.857	0.729	-0.985**	-0.812	-0.010	0.173	0.015	-0.883*
Na ⁺	-0.233	-0.946*	0.740	0.467	-0.582	-0.862	-0.780	0.632
Cl ⁻	-0.592	-0.905*	0.918*	0.913*	0.030	-0.341	-0.282	0.977**

TC, total microbial count; CFU/TC, colony-forming units on R2A agar to total microbial count ratio; PHAC, photoautotrophic cell count; PHAC/TC, photoautotrophic cell count to total microbial count ratio morphotype diversity index (Shannon index), DGGE, phylogenetic diversity of native samples by taxonomic operational units (OTUs); EcoPlate, functional diversity of native samples by positive response numbers on Biolog EcoPlates) and distance from glacier terminus and chosen biological parameters and non-biological factors

Porg ($p < 0.05$) in surface ice. This parameter in cryoconite samples correlated positively with SUVA, Cl⁻ ($p < 0.01$) and POC contents ($p < 0.05$) but negatively with pH ($p < 0.01$) and Porg and Ptot contents ($p < 0.05$).

The CFU/TC ratio displayed several negative correlations with biological and chemical factors in surface ice, *inter alia* with the phylogenetic and functional diversity ($p < 0.05$) (Table 4). The statistical principal component analysis (PCA) of physico-chemical data showed significant differences between the sampling points, with a clear distinction between surface ice and cryoconite holes, caused mainly by the different concentration of nutrients and suspended solids in both types of samples (Fig. 4). The PCA of microbiological data indicated an even more pronounced distinction of these habitats. Microbiocenoses of surface ice samples seem to form one cluster, defined mainly by the CFU/TC ratio, whereas cryoconite hole microbiota differ greatly from one melt hole to another (Fig. 5).

Discussion

Microbial community development on a transect across the ablation zone of Ecology Glacier was investigated, from the glacier's terminus to the snow line (Table 1). The transect has been established in anticipation of there being microbial community structure changes as the underlying glacial ice is exposed by melting snow. Chemical

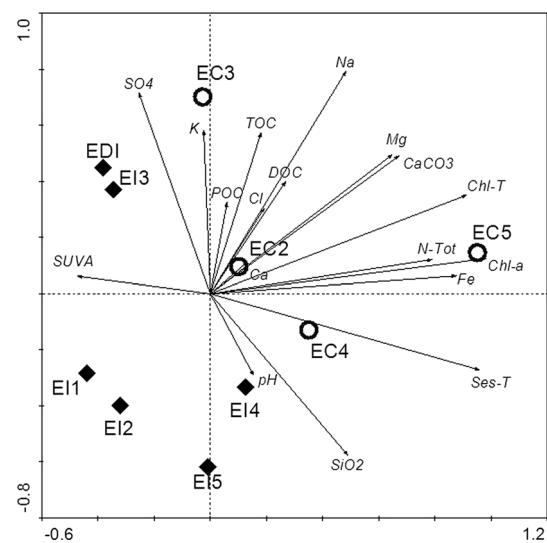


Fig. 4 Principal component analysis of physico-chemical data in surface ice (black diamonds) and cryoconite samples (circles). TOC, total organic carbon; DOC, dissolved organic carbon; POC, particulate organic carbon; Ntot, total nitrogen; SO₄, sulphate conc.; K, potassium conc.; Cl, chloride conc.; Na, sodium conc.; Mg, magnesium conc.; CaCO₃, calcium carbonate conc.; Chl-T, total chlorophyll conc.; Chl-a, chlorophyll *a* conc.; Fe, total iron conc.; Ses-T, total seston conc.; SiO₂, silica conc.; SUVA, aromatic content of DOC

composition, bacterial phylogenetic and microbial functional diversity, and abundance were each determined in surface ice and cryoconite holes. Our data describe microbial supraglacial habitats and a comparison of direct and culturable approaches.

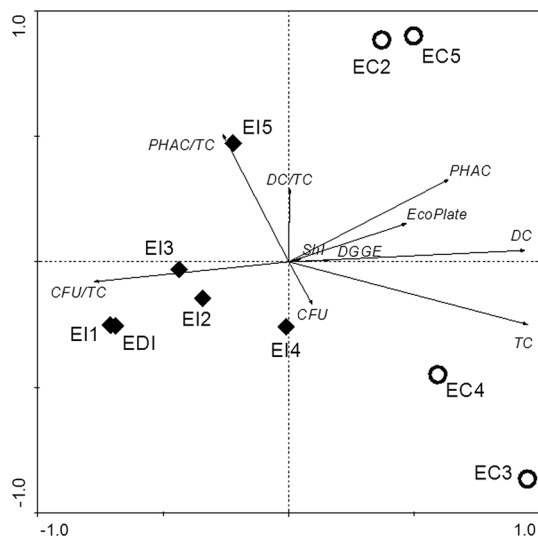


Fig. 5 Principal component analysis of microbiological data in surface ice (black diamonds) and cryoconite samples (circles). PHAC, photoautotrophic cell count; TC, total microbial count; CFU/TC, ratio of colony-forming units on R2A agar to total count; PHAC/TC, photoautotrophic cell count to total microbial count ratio; DC/TC, ratio of dividing cell count to total count, *ShI*, morphotype diversity index (Shannon index); DGGE, phylogenetic diversity of native samples by operational taxonomic units (OTUs); CFU, colony-forming units on R2A agar; *EcoPlate*, functional diversity of native samples by positive response numbers on Biolog Ecoplates; DC, dividing cell count

Physico-chemical and biological data

Chlorophyll *a* content, DOC, inorganic nitrogen compounds, and reactive phosphorus concentrations were in line with published values for Antarctic and high Arctic glaciers (Porazinska et al. 2004; Foreman et al. 2007; Telling et al. 2012; Zarsky et al. 2013). Anion and cation concentrations were at the same level or slightly lower than those published for Antarctic glaciers, with the exception of Mg^{2+} which were up to 10× higher in Ecology Glacier (Welch et al. 1996; Foreman et al. 2007).

Our TC data were in line with those previously reported for Ecology Glacier surface ice, and those for Canada Glacier, Antarctica (Foreman et al. 2007; Zdanowski et al. 2013). Stibal et al. (2012) and Singh et al. (2014) reported higher microbial abundance in cryoconite holes at Svalbard and Greenland glaciers than we have done here. However, these authors' counts were based on per gram of sediment; Stibal et al. (2012) also presented photoautotroph counts of $10\text{--}10^2$ lower than the total count, also consistent with our observations. The little published data on supraglacial CFU numbers that are available are in line with our findings (Christner et al. 2003; Zdanowski et al. 2013; Singh et al. 2014).

Surface ice and cryoconite hole microbiocenosis comparison

Cryoconite holes have been praised to be the hot spots of microbial life on glaciers (Edwards et al. 2013), yet a comprehensive comparison of cryoconite hole and surface ice habitats is scarce. Our study shows that microbial counts were indeed on average 20× higher in cryoconite holes than in the surrounding surface ice, although the carbon fraction amounts were not that different, which implies that the carbon concentration is not a limiting factor. This is consistent with the findings of Bagshaw et al. (2013) on McMurdo Dry Valley Glaciers. Total nitrogen, phosphorus, and iron levels were however considerably higher (on average 4, 3, and 8 times, respectively). The higher macro and micronutrient levels may be attributed to physical and chemical weathering of the cryoconite sediment, but also to microbial activity like nitrogen fixation (Telling et al. 2011) and bioleaching of insoluble mineral compounds (Frey et al. 2010), especially iron containing minerals (Bashan et al. 2013). The slightly elevated PO_4^{3-} and Ca^{2+} concentrations in surface ice compared to cryoconite holes may be attributable to the lower demand for those nutrients of the less numerous microbiota in surface ice with weathering processes still present. The PHAC/TC ratio was the same in cryoconite hole and surface ice, yet the chlorophyll levels were on average 10 times higher in cryoconite samples. Considering that cell volume in cryoconite holes was approximately double that in surface ice, this could suggest that this habitat supports bigger photoautotroph cells containing more chlorophyll. The bacterial OTU numbers were not much higher in cryoconite holes, but the phylogenetic structure of those communities (based on DGGE band migration patterns and clustered by the PCA) differed greatly. However, the similarities in the phylogenetic community structure of the surface ice sample EI5 and cryoconite sample EC5 gathered at the verge of the snow line could suggest the same source of influence, presumably the melting snow. The more diverse bacterial phylogenetic structure from one cryoconite hole to another compared to surface ice and the higher metabolic activity given as a spectrum of assimilated carbon sources implies a rapid development of a more complex trophic food web and a more diverse nutrient resource since the melthole formation. An evolution on a chemical level throughout the summer season in cryoconite holes was observed by Bagshaw et al. (2007) on McMurdo Dry Valley Glaciers.

Spatial variation issue

Many researchers investigating glacial microbiota have collected samples along a transect like ours (Mindl et al. 2007; Uetake et al. 2010; Edwards et al. 2011; Zarsky et al.

2013). However, few have explained the spatial variation of chemical and biological components measured. Moreover, none regarded the ablation zone as an annual chronosequence of ice exposure from beneath the melting snow, although some mentioned the receding snow line during the summer season (Telling et al. 2011; Stibal et al. 2012). Simple regression analysis showed several positive correlations with distance from the glacier edge in surface ice samples and cryoconite holes. A strong correlation ($p < 0.01$) between distance from glacier terminus and the morphotype Shannon index in surface ice was observed; there was a weaker correlation ($p < 0.05$) between distance from glacier terminus and chlorophyll *a*, nitrogen, and seston concentrations in cryoconite holes. These observations reflect the finding of Stibal et al. (2012) for a Greenland glacier, and Takeuchi (2001) for algal biomass in an Alaskan glacier. Relatively high amounts of dissolved organic nitrogen and phosphorus at the snow line (EI5) could also be attributed to either nutrient excretion or cell lysis of the snow-derived microbiota. Takeuchi et al. (1998) observed a replacement of the snow-specialists by ice-specialist algae at the snow line of a Himalayan Glacier, suggesting cell degradation of the former group. The average length of photoautotrophic cells also increased with distance from the glacier terminus. This might be attributable to more frequent sightings of eukaryotic cells (large, spheroid, single cells, exhibiting characteristic autofluorescence in green-filter light) in the upper parts of the ablation zone—an observation also made by Uetake et al. (2010). Takeuchi (2001) suggested that filamentous and colonial cyanobacteria have an advantage in lower parts of the ablation zone by not being washed from the ice by meltwater.

Considering that the ablation zone represents a time scale, some interesting features do emerge. For example, a substantial spike in microbial total count was noted in both surface ice and cryoconite holes at sampling point 4. This was preceded by or coincided with high photoautotroph counts and chlorophyll contents as mentioned above. Such a phenomenon has also been reported for cryoconite holes at Werenskiold Glacier (Svalbard, Arctic) through the whole ablation season (Stibal et al. 2008b). TC counts in surface ice correlated with high functional diversity and chlorophyll levels. This may indicate a potential role of photoautotrophs in promoting microbial growth, again either via organic carbon excretion or cell lysis. In this respect, Medina-Sánchez and Villar-Argiz (2006) noted substantial DOC excretion by algae in a boreal lake in response to UV irradiation and P limitation. Similarly, Myklestad (1995) observed that P and N limitation caused eukaryotic photoautotrophs (especially diatoms) to excrete polysaccharides and free amino acids. Such responses could explain the marked affinity of supraglacial communities for polymers and amino acids. High TC and

phylogenetic community diversity (number of OTUs) at the verge of the snow line observed here have also been seen on the verge of the snow in the ablation zone of Alaska's Gulkana Glacier (Segawa et al. 2010).

Microbial counts (TC) fell at point 3 (EI3, EC3), while the phylogenetic structure was similar to points at the snow line (EI5, EC5). A potential mechanism that might give rise to this is viral lysis, given that DOC concentrations peaked in the surface ice and cryoconite holes at this point in the transect. Bellas et al. (2013) described high viral abundance in cryoconite holes on an Arctic glacier, and concluded that most of the DOC released in these habitats derived from viral-induced mortality.

Significant environmental controls

A consistent parameter here whose benefit to the microbial community was reflected in elevated abundance and diversity was particulate organic and inorganic material (seston, SiO_2 or CaCO_3). This was apparent, especially in surface ice where this material was labile. The amount of particulate material correlated positively with microbial abundance and functional diversity, whereas CaCO_3 concentration corresponded with bacterial phylogenetic diversity. The beneficial role of particulates for microbial activity have been previously recorded for polar aquatic habitats (Junge et al. 2004).

Nitrogen and phosphorus seem to be the major limiting factors in the supraglacial environment of Ecology Glacier, as amounts of organic nitrogen in surface ice correlated with the DC/TC and PHAC/TC ratios (despite a strong negative correlation with TOC of the former). Functional diversity on the other hand correlated positively with organic phosphorus levels. Total and organic phosphorus emerged as the only strong correlation (with TC), when conducting the simple regression analysis for all the ice and cryoconite samples together (data not shown). Phosphorus and nitrogen deficits as factors limiting microbial growth and activity in supraglacial environments have been reported before for Arctic glaciers (Stibal et al. 2008a; Telling et al. 2011).

SUVA measurements indicate mainly humic and fulvic acid abundance (Weishaar et al. 2003). Their presence has been connected with decreased albedo of cryoconite sediment, higher water temperatures, and a subsequent increase in microbial activity (Stibal et al. 2006; Edwards et al. 2011). Our data suggest that the PHAC/TC ratio and tetrazolium reducing activity in Ecoplates increased in cryoconite samples in which SUVA measurements were elevated.

Culturable glacial microbes

Cultivation techniques have been dismissed as flawed and of little use in obtaining data on microbial communities in

environmental settings (Wagner et al. 1993; Ranjard et al. 2000). However, Sigler and Zeyer (2004) proposed cultivable microorganisms to be the opportunistic part of the community, thus making their contribution to the total count an indicator of the stability of conditions in microbial habitats. Thus, a pattern can be determined along the transect defined here, with several correlations emerging between the CFU/TC ratio and other factors. The CFU/TC ratio displayed a negative correlation with distance from glacier edge implying a decrease in habitat stability towards the terminus. Factors that positively correlated with microbial abundance (Ptot, Porg, nitrate) correlated negatively with the CFU/TC ratio, indicating their positive influence on the non-cultivable part of the microbial community, and at least according to Sigler and Zeyer (2004), showing where unstable and unfavorable conditions prevailed where the ratio was the highest (EDI, EI1). This part of the ablation zone was characterized by high mineral content, low microbial numbers and activity, and a high phylogenetic diversity. Wind-transported mineral debris at glacial termini have been observed, and are believed to be a nutrient source (Hodson et al. 2008; Uetake et al. 2010). High OTU and TC numbers have also been noted at an Alaskan glacier's edge (Segawa et al. 2010), while Stibal et al. (2012) described low microbial numbers and activities in a comparable area on a Greenland glacier. Considering the glacier edge borders three very different ecosystems (supraglacial, subglacial, and proglacial), the extant microbial community structure might be greatly influenced by allochthonous microorganisms that contribute to elevated phylogenetic diversity but fail to cope metabolically with the prevailing conditions (Zdanowski et al. 2013). The R2A/8 plate wash showed that metabolic potential of this community can be 'unlocked' in the right conditions. Interestingly, the functional diversity and the CFU/TC ratio shared many correlation (positive and negative) with environmental factors in cryoconite holes yet there was no correlation between the two. This implies, that the same factors that decrease the non-opportunist fraction abundance also widen the communities' metabolic spectrum of carbon source assimilation.

R2A and the eight-fold diluted R2A recovered cultivable bacteria diversity well, with >50 % on R2A and >60 % on R2A/8 bands identical to those in native samples. The plate wash also revealed the functional potential of the supraglacial community, especially those in the surface ice, since native samples responded only weakly. Wiebe et al. (1992) explained this phenomenon, stating that the heterotrophic microbiota in cold aquatic habitats have reduced substrate affinities and thus require higher nutrient concentrations. However, the plate wash from the diluted medium has proven more adequate for glacial microbiota, suggesting that nutrient concentrations need to be adjusted to reflect that of the environment.

Conclusions

Passage of the snow line and subsequent exposure of the glacial ice surface to ablation and allochthonous influences changes environmental conditions on Ecology Glacier. These changes caused the emergence of different microbial habitats, of which surface ice and cryoconite holes are most prominent, yet very different in terms of quantity and quality of the microbiocenosis' structure. The spatial variability of both of these habitats indicates that the melting snow was a major source of nutrients and living cells to the glacial surface. A cascade of events triggered by this "snow augmentation" shaped the supraglacial communities towards the glacial terminus, indicating their high adaptive potential. The enriching effect was, however, only temporary. The cryoconite hole and surface ice communities responded similarly to snow melt and exposure to allochronic inputs, despite their differences in microbial numbers and diversity. Nitrogen and phosphorus amounts were the main factors responsible for limiting microbial abundance and activity. We hope our findings will contribute to the still sparse knowledge of these rapidly disappearing habitats and provide insight into microbe-environment interactions.

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