Eukaryotic microorganisms in cold environments: examples from Pyrenean glaciers

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INTRODUCTION

Small eukaryotes are probably the most abundant eukaryotes on Earth. They have been found in all extreme environments (Caron et al., 1999), addressing a wide range of temperatures from tropical oceans (Li et al., 1994) to polar sea ice (Kucby et al., 2011).

The discovery of cold-tolerant microorganisms in glaciated and permanently frozen environments has broadened the known range of environmental conditions which support microbial life. Although these microorganisms that inhabit permanently cold ecosystems (representing one of the largest biophere reserves on Earth) have been studied only for their ability to survive in such extreme conditions, recent studies have provided evidence that these habitats (deep sea, Polar Regions, mountain glaciers, etc.) can be colonized by both obligate and facultative psychrophilic microorganisms (Alcazar et al., 2010). Such ecosystems represent one of the last unexplored frontiers of ecology, and psychrophilic microbial populations sharing such habitats constitute an important part of cold-adapted biodiversity and play an essential role as nutrient cyclers and organic matter mineralizers.

It is difficult to characterize these organisms by simple observation with optical microscopy, and cultivation methods do not allow all the organisms to grow. Pigment and/or fatty acid analysis can provide some information on the structure and dynamics of the phototrophic and/or heterotrophic behavior of small eukaryotes, but the phylogenetic information supplied by these methods is limited (Lefranc et al., 2005). During the last decade, molecular techniques have greatly increased our knowledge by identifying the smallest organisms. Environmental studies of eukaryotic diversity based on polymerase chain reaction (PCR) amplification,
Viable bacterial communities have been observed beneath glaciers in the northern Apennines became the southernmost in Europe. If the present warming and associated environmental changes are predicted to have strong impacts on high-altitude ecosystems (IPvC, 2007), and the European mountains have already experienced an increase in temperature of 1–2°C during the twentieth century (Beniston et al., 1997; Dyck et al., 2012). A possible consequence of permafrost thawing, based upon predicted global warming scenarios, is that there may be an increase in microbial activity and an increase in active layer thickness (Gilichinsky and Wagen, 2006). Current atmospheric warming will have severe consequences for the structure and functioning of glacier ecosystems with changes that, in turn, may feed back on the global-scale composition of the atmosphere (Michelson et al., 2012; Wagner et al., 2012). Here we report the diversity and distribution of microbial eukaryotes in four Pyrenean glaciers studied by 18S rRNA gene libraries and addresses some interesting questions: (i) what is the effect of altitude and glacier area on the composition of the microbial community? (ii) taking into account that ice melting is more dramatic in lower glaciers, which are the environmental differences associated to ice melting that affect microbial community? (iii) is it possible to discriminate between the effect on microbial community of area/altitude and the effect of chemical parameters? (iv) is it possible to find a microbial community or species to be used as an indicator of glacier retreat?

MATERIALS AND METHODS

SAMPLE COLLECTION AND PROCESSING

Glacial ice samples were collected at four sites on the northern slope of Pyrenees: Aneto glacier (ANE), Maladeta glacier (MAL), Monte Perdido glacier (PER), and Lliterola glacier (LIT) in August, 2010 (Figure 1). These glaciers are located at altitudes of 3,404, 3,035, 3,355, and 2,740 m, and their surface are 64, 31, 32, and 1.5 ha, respectively (Rene, 2007; Arenillas et al., 2008). Ice samples were obtained by removing 20–30 cm of thick debris and cutting out a square block of 20 cm on a side. Three sampling replicates were collected from each glacier. Samples were wrapped in plastic bags and stored at −20°C until processing. Ice samples were processed by using a surface decontamination and melting procedure consistent with previous studies (Bidle et al., 2007). A section of a block of ice was removed from −20°C and soaked in ice-cold 95% ethanol for 1 min, followed by extensive rinsing with 0.2 μm-filtered MilliQ water, effectively ablating the exterior 3-cm shell of ice samples (corresponding to 30% of total ice volume). These procedures were effective at removing surface contamination from inner shell ice samples (Rogers et al., 2004; Christner et al., 2005). The decontaminated interior ice was thawed in a sterile plastic bag at 4°C and used for analyses. To control for laboratory contamination, 11 of MilliQ rinse water was frozen, thawed, filtered, and subjected to identical analytical procedures. All procedures were performed by using bleach-sterilized work areas, a UV-irradiated laminar flow hood, ethanol-sterilized tools, and sterilized gloves.
CHEMICAL ANALYSIS OF MELTWATER
Basic measurements of physical and chemical parameters of meltwater from various sites were made with a temperature-calibrated pH, conductivity, and salinity meter (WTW, Weilheim, Germany). Assays for dissolved inorganic nitrogen (NH$_4^+$, NO$_2^-$, and NO$_3^-$) were performed by ion chromatographic method using suppressed conductivity detection in a 861 Advance Compact IC system (Metrohm AG, Herisau, Switzerland). Chromatograms were recorded using the Metrohm IC Net 2.3 SR4 software. The system was run in the isocratic mode with the column at 45°C.

SCANNING ELECTRON MICROSCOPY
Samples (50 ml volume) for scanning electron microscopy (SEM) observation were filtered in the lab (0.22 μm Millipore filters), preserved in 2% glutaraldehyde, rinsed in 0.22 μm-filtered 50 mM phosphate buffer, and dehydrated using an ascending (30, 50, 70, 90, and 100%) series of ethanol. Filter disks were then air-dried overnight and mounted on SEM stubs with carbon pads and sputter coated with gold-palladium for 1.5 min at 15 mA voltage. Observation and imaging were examined using a Jeol 5600LV scanning electron microscope with an INCA Oxford auxiliary X-ray energy-dispersive spectroscopy microanalytical system.

DNA EXTRACTION AND PCR AMPLIFICATION
Approximately 400 ml of each frozen sample was melted at 4°C, and filtered through a 0.22-μm filter (Millipore). Community DNA was extracted using the GNOME BIO101 kit (MP Biomedicals, Illkirch, France) and purified with QIAquick PCR purification kit (QIagen, Hilden, Germany) according to manufacturers’ instructions. Extraction procedures were identical for all ice samples. The 18S rDNA genes from mixed microbial DNA were amplified by PCR. Near full-length 18S rDNA fragments were amplified by PCR using the eukaryotic-specific primers 1F (CTG GTT GTA T...).
CCT TGC CAG; Lefranc et al., 2005) and S02R (ACC AGA CCT GGC CTC C; Amann et al., 1990). PCR was carried out under the following conditions: 33 cycles (denaturation at 94°C for 30 s, annealing at 46°C for 30 s, extension at 72°C for 35 s), was preceded by 5 min denaturation at 94°C, and followed by 7 min extension at 72°C. PCR was optimized by both diluting the template and by increasing the number of thermal cycles.

To control for false-positive PCR signals, 1 1 of MiRep water was frozen, thawed, and subjected to the same DNA extraction procedure. This material was used as a template with the specific primers to test for contamination and PCR artifacts.

SEQUENCING AND PHYLOGENETIC ANALYSIS

18S rDNA PCR amplicons were cloned using TOPO TA Cloning Kit (Invitrogen, Carlsbad, CA, USA), and plasmid DNA (112 ANE clones, 35 MAL clones, 108 PER clones, and 44 LIT clones) was bidirectional sequenced with a 3730XL sequencer (Applied Biosystems). Chimeric sequences were identified with the CHIMERA-CHECK program (Ribosomal Database Project II, Michigan State University, East Lansing, MI, USA) and discarded.

Sequences were analyzed using BLAST at the NCBI database1. Representative sequences were aligned using the software ClustalW (version 2.0; Larkin et al., 2007), the results were corrected manually and alignment uncertainties were omitted in the phylogenetic analysis. Their phylogenetic relationship was analyzed using the software MEGA 5 (Tamura et al., 2011) and the ABB software (Ludwig et al., 2004) with parsimony, neighbor-joining, and maximum likelihood analyses. In all cases, general tree topology and clusters were stable, and reliability of the tree topologies was confirmed by bootstrap analysis using 1,000 replicate alignments. A consensus tree was generated.

Rarefaction analysis was performed using Analytic Rarefaction software (version 1.3)2, based on previous analytic solutions (Raup, 1975; Tipper, 1979).

STATISTICAL ANALYSIS

Statistical differences on the number of clones, number of operational taxonomic units (OTUs) and Shannon–Wiener index were studied by analysis of variance (ANOVA) test. Data of OTUs and clones are media values of three sampling replicates. Relationships between the number of taxa found in glaciers and environmental variables were analyzed by linear regression analysis (Pearson’s correlation coefficient r).

Abundance-based coverage estimator (ACE) and Chao1 (Chao and Jostman et al., 1995). For statistical analysis, Monte Carlo permutation tests with 500 permutations were used.

NUCLEOTIDE SEQUENCE ACCESSION NUMBERS

Sequences obtained in this study have been deposited in the EMBL sequence database under accession numbers JX196712 and JX456225 to JX456234.

RESULTS

GENERAL CHARACTERISTICS OF THE ICE SAMPLES AND CHEMICAL PROPERTIES

In this study, we assessed and compared the composition of eukaryotic microorganisms present in samples from four Pyrenean glaciers. ANE, MAL, PER, and LIT contained a broad size spectrum of particles and sand debris as the ice contained a layer of sand and organic matter of aeolian origin below the surface. Inorganic particles contributed to variations in chemical properties between the meltwater samples. Meltwater of ANE and MAL were pH 6.7 and 6, respectively, whereas those from PER and LIT were 4.8 and 5.7, possibly due to the chemical reactions of fine particles of, e.g., pyrite (Table 1). These fine particles were analyzed in a spectroscopy microanalytical system in order to exactly define whether the observed particles corresponded to microorganisms or inorganic material (Figures 1G, H). Spectrum 1 (Figure 1G) contains two major peaks corresponding to silicon and oxygen, demonstrating it is a clay mineral, moreover spectrum 2 (Figure 1H) shows a major peak of carbon. In addition, ice properties such as salinity, pH, NH3, NO2, and NO3 contents were determined (Table 1). The ice samples had overall low salinity and highly variable NO2 and NO3 contents ranging from 3.4 μM in ANE to 73.2 μM in LIT for NO3 and ranging from 4.5 μM in ANE to 103.5 μM in LIT for NO2. Further, NH3 presented a moderate variability between 2.1 μM in ANE and 6.6 μM in PER (Table 1). Generally, samples from higher glaciers showed lower amounts of ion concentrations, as salt solubility decreases in cold environments.

18S rDNA GENE CLONE LIBRARIES

In order to understand the microbial eukaryotic populations of the ice samples, we amplified community DNA and constructed clone libraries with eukaryotic-specific 18S rDNA primers (Table 2). After removal of potential chimeric sequences and dereplication of identical sequences, phylogenetic analysis demonstrated that 31 different phylotypes could be identified (10 from 112 ANE clones, 7 from 35 MAL clones, 9 from 108 PER clones, and 5 from 44 LIT clones; Figure 2). Most of BLAST analysis of the 31 phylotypes (Table 2) revealed no identical clones to 18S rDNA sequences in the GenBank, to currently cataloged species. Overall, the ANE, MAL, and PER clone libraries showed a high diversity (Table 3, Figure 2) with sequences belonging to Viridiplantae, Stramenopiles, Fungi, Rhizaria, and Metazoa divisions.

COMMUNITY COMPOSITION AND STATISTICAL ANALYSIS: SPECIES RICHNESS, ABUNDANCE, AND DIVERSITY

The microbial communities in higher glaciers differed from the lower glacier communities. Regardless of the chemical composition of ice (Table 1), this difference was mainly due to the large numbers of Viridiplantae and Rhizaria among others in higher glaciers. There was also a marked decrease in the number of clones, number of OTUs and the Shannon index of diversity.
Table 1 | Chemical analysis of ice meltwater from ANE, MAL, PER, and LIT.

| Sample  | pH  | Salinity (ppt) | NH4 (μM) | NO2 (μM) | NO3 (μM) |
|---------|-----|----------------|----------|----------|----------|
| ANE-1   | 6.7 | 0.23           | 1.8      | 4.0      | 5.4      |
| ANE-2   | 6.7 | 0.27           | 2.3      | 3.1      | 3.9      |
| ANE-3   | 6.8 | 0.21           | 2.2      | 3.2      | 4.3      |
| Average (SD) | 6.7 (0.06) | 0.24 (0.03) | 2.1 (0.26) | 3.4 (0.49) | 4.5 (0.77) |
| MAL-1   | 6.5 | 0.20           | 4.0      | 4.8      | 6.5      |
| MAL-2   | 6.2 | 0.29           | 3.1      | 5.3      | 7.2      |
| MAL-3   | 5.3 | 0.23           | 4.3      | 4.9      | 7.0      |
| Average (SD) | 6.0 (0.51) | 0.24 (0.04) | 3.8 (0.51) | 5.0 (0.22) | 6.9 (0.29) |
| PER-1   | 4.9 | 0.15           | 75       | 3.5      | 6.1      |
| PER-2   | 5.0 | 0.21           | 5.9      | 4.1      | 3.9      |
| PER-3   | 4.6 | 0.19           | 6.4      | 3.8      | 4.2      |
| Average (SD) | 4.8 (0.21) | 0.18 (0.03) | 6.6 (0.62) | 3.8 (0.30) | 4.7 (0.19) |
| LIT-1   | 5.5 | 0.35           | 5.5      | 68.1     | 110.5    |
| LIT-2   | 5.9 | 0.28           | 6.8      | 76.3     | 101.6    |
| LIT-3   | 5.8 | 0.31           | 4.9      | 75.2     | 98.5     |
| Average (SD) | 5.7 (0.21) | 0.31 (0.06) | 5.7 (0.97) | 73.2 (4.45) | 103.5 (6.23) |

The highest microbial diversity was found in PER, followed by ANE, and the similarity between ANE and PER was significant (p < 0.004; Figure 5; Table 5). Other variables such as salinity and NH4 showed a negative correlation in Rhizaria (p = 0.0203) and Fungi (p = 0.0165), respectively. NO2− and NO3− also presented a negative correlation in the total number of OTUs (p = 0.0424 and 0.0432), respectively.

DISTRIBUTION OF TAXA AND PHYLOTYPES ACROSS ALL SAMPLES

All the sequences were affiliated to five phyla, Fungi, Metazoa, Rhizaria, Viridiplantae, and Stramenopiles, representing 9.7, 4.7, 36.5, 45.8, and 3.34%, respectively (Figure 6). The dominant taxa were represented in all samples with the exception of Metazoa in PER and Stramenopiles in ANE and MAL. The members of rare phyla (~1% of all classified sequences) included a member of the family Chrysophyceae in PER and a member of Viridiplantae in LIT samples (Table 2). The most abundant phylotype across all samples was a glacier algae in ANE and MAL samples, Chloromonas platystigma, representing 16% of all sequences. The most abundant phylotypes in PER and LIT were two uncultured Cercozoan, representing 6 and 5% of all sequences, respectively (Table 2).

At the genus level, the comparison of the relative abundances revealed significant differences between glaciers. Chloromonas was the most abundant genus across all ice samples, representing 22% of all classified sequences in ANE and 4% in MAL. The distribution of the other dominant genera Raphidonema (4.6%), Heteromita (5.3%), Kolellia (5.4%), and Bodomorpha (4%) varied significantly between glaciers (p < 0.0001; Table 2).

DIFFERENCES IN COMMUNITY STRUCTURE BETWEEN GLACIERS

The relative abundances of dominant taxa varied between glaciers. Our clone libraries were clearly dominated by Viridiplantae in ANE.
Table 2 | Analysis of eukaryotic 18S rRNA clones retrieved from ANE, MAL, PER, and LIT ice samples.

| Number | Clone or group name | Max. no. of clones in glacier | Accession number | Closest sequence match from BLAST search | Similarity (%) | Taxon |
|--------|---------------------|-------------------------------|------------------|------------------------------------------|----------------|-------|
| 1      | ANE25 2 AF289921    | 10                            | JM191101         | Uncultured fungus sp. Kshps3-26          | 99             | Fungi; environmental samples |
| 2      | ANE28 2 AF144016    | 10                            | JM191101         | Chloromonas sp. 047-89                    | 99             | Viridiplantae; Chlorophyta; Chlorophyceae; Chlamydomonadales |
| 3      | ANE33 8 AJ067742    | 8                             | KM191101         | Uncultured chlorophyta alga, clone J53-W5-Un21 | 99             | Viridiplantae; Chlorophyta; environmental samples |
| 4      | ANE58 36 AF144011   | 8                             | KM191101         | Glacier alga: Chloromonas sp. f. pylatusigmata 020-99 | 99             | Viridiplantae; Chlorophyta; Chlorophyceae; Chlamydomonadales; Chlamydomonadaceae; Chloromonas |
| 5      | ANE59 8 AJ067731    | 8                             | KM191101         | Uncultured Chytridiomycete, clone WS 10-E15 | 99             | Fungi; Chytridiomycota; environmental samples |
| 6      | ANE66 8 AJ067718    | 8                             | KM191101         | Uncultured phototrophic eukaryote, clone RS 8-Un56-B | 92             | Viridiplantae |
| 7      | ANE67 4 AJ067714    | 4                             | KM191101         | Uncultured Cercozoan, clone BS 7-E06 | 96             | Rhizaria; Cercozoa; environmental samples |
| 8      | ANE71 2 AF164272    | 2                             | KM191101         | Raphidonema hardyi isolate AFTOL-ID 31 | 96             | Fungi; Chytridiomycota; Raphidonematales; Raphidonemataceae; Raphidonema |
| 9      | ANE72 4 AJ067712    | 4                             | KM191101         | Uncultured Cercozoan, clone BS 7-E06 | 96             | Rhizaria; Cercozoa; environmental samples |
| 10     | ANE79 8 AJ067712    | 8                             | KM191101         | Uncultured Cercozoan, clone BS 7-E06 | 96             | Rhizaria; Cercozoa; environmental samples |
| 11     | MAL1 1 AF289921     | 1                             | KM191101         | Uncultured fungus sp. Kshps3-26          | 99             | Fungi; environmental samples |
| 12     | MAL2 5 AF289921     | 5                             | KM191101         | Uncultured chlorophyta alga, clone J53-W5-Un21 | 99             | Viridiplantae; Chlorophyta; environmental samples |
| 13     | MAL3 12 AF144011    | 12                            | KM191101         | Glacier alga: Chloromonas cf. pylatusigmata 020-99 | 97             | Viridiplantae; Chlorophyta; Chlorophyceae; Chlamydomonadales; Chlamydomonadaceae; Chloromonas |
| 14     | MAL4 2 AJ067714     | 2                             | KM191101         | Uncultured Cercozoan, clone BS 7-E06 | 96             | Rhizaria; Cercozoa; environmental samples |
| 15     | MAL5 2 AJ067714     | 2                             | KM191101         | Uncultured Cercozoan, clone BS 7-E06 | 96             | Rhizaria; Cercozoa; environmental samples |
| 16     | MAL6 11 AM144014    | 10                            | KM191101         | Uncultured Cercozoan, clone WIFI144 | 98             | Rhizaria; Cercozoa |
| 17     | MAL7 2 AJ067729     | 2                             | KM191101         | Uncultured Chytridiomycete, clone WS 10-E02 | 99             | Fungi; Chytridiomycota; environmental samples |
| 18     | PER25 20 AM144014   | 20                            | KM191101         | Uncultured Cercozoan, clone WIFI144 | 98             | Viridiplantae; Chlorophyta; environmental samples |
| 19     | PER41 18 AF48477    | 18                            | KM191101         | Raphidonema nivalis                      | 98             | Viridiplantae; Streptophyta; Klebsormidiophyceae; Klebsormidiales; Elakatotrichaceae; Raphidonemata |
| 20     | PER44 16 AF48477    | 16                            | KM191101         | Heteromita globosa strain SCCAP-H251 | 99             | Rhizaria; Cercozoa; Cercomonadidae; Heteromita; Heteromitaraceae |
| 21     | PER46 4 AF395487    | 4                             | KM191101         | Chlamydomonas nivalis                    | 97             | Viridiplantae; Chlorophyta; Chlorophyceae; Chlamydomonadales; Chlamydomonadaceae; Chlamydomonas |
| Number | Clone or group name | Mean no. of clones in glacier | Accession number | Closest sequence match from BLAST search | Similarity (%) | Taxon |
|--------|---------------------|-------------------------------|------------------|------------------------------------------|---------------|-------|
| 22     | PER49               | 16                            | A3011569         | Koliella antarctica                      | 100           | Viridiplantae, Streptophyta, Klebsormidiales; Klebsormidiaceae; Koliella |
| 23     | PER50               | 20                            | AY95966          | Soil flagellate AND21                    | 99            | Phaeolalia; Cercozoa; Cercomonadida; Heteromitiidae |
| 24     | PER58               | 12                            | DQ211566         | Bodomorpha sp. HFC037                    | 97            | Phaeolalia; Cercozoa; Cercomonadida; Heteromitiidae, Bodomorpha |
| 25     | PER60               | 2                             | AY957435         | Uncultured chrysophyte, clone JU-ICE-Un-10 | 99            | Stramenopiles; Chrysophyceae; environmental samples |
| 26     | PER61               | 4                             | AY957629         | Uncultured Chrysidomyctaceae, clone V5-10-E02 | 98            | Fungi; Chytridiomycota; environmental samples |
| 27     | LIT1                | 8                             | HF913653         | Ardissonea formosa                       | 97            | Stramenopiles; Bacillariophyceae, Fragilariophyceae, Fragilariophycidae; Fragilariaceae, Fragilaria; Ardissonea |
| 28     | LIT2                | 10                            | AY101826         | Bensingtonia yamatoana                   | 99            | Fungi; Dikarya; Basidiomycota; Pucciniomycotina; Agaricostilbomycetes; Agaricostilbomycetes incertae sedis; mitosporic Agaricostilbomycetidae, Bensingtonia |
| 29     | LIT3                | 16                            | FN35463          | Uncultured eukaryote OTP210800382       | 98            | Phaeolalia; Cercozoa |
| 30     | LIT4                | 8                             | GQ501307         | Brachionus calyciflorus                  | 99            | Metazoa; Rotifera; Monogononta; Ploimida; Brachionidae; Brachionus |
| 31     | LIT5                | 2                             | DQ104080         | Uncultured Chlorophyta P200E-4          | 97            | Viridiplantae; Chlorophyta |
FIGURE 2 | Phylogenetic analysis of microbial community DNA in ice samples. Consensus phylogenetic tree derived from 18S rRNA gene sequence data showing the four groups of microbial eukaryotes found in Pyrenean glaciers. The distance corresponding to one base change per hundred nucleotide positions is indicated by the scale bar. Accession numbers for the sequences used to make this tree are given in Table 2. Names in capital letters (ANE, MAL, PER, and LIT) correspond to clones retrieved in this study.

...and MAL where they represented nearly 80%. Rhizaria, especially represented by the Cercozoa in PER (Figure 6) and sequences affiliating to Rhizaria were also the most abundant in LIT library.

Principal components analysis (PCA; analysis no. 1) based on the relative abundances of the microbial phyla confirmed that microbial communities in glaciers were quite different (Figure 7). MAL contained the only samples that shared similar composition with those from ANE and PER. ANE and PER contained only one common species (a soil flagellate belonging to Cercozoa), and LIT appears to be the most different.

Canonical correspondence analysis with all environmental variables was used to estimate the proportion of the community variability attributable to variability in the environment, which was estimated in several runs, each with a single variable. The eigenvalues corresponding to the four ordination axes were used to characterize the results of particular analysis (Table 6). CCA diagrams show the interrelationships between microbial communities and environmental variables that were observed in the four glaciers (Figure 8). The OTUs and sampling points mutually portray the dominant patterns in community composition to the extent that these could be elucidated by the selected variables (Jongman et al., 1995). The length of an arrow representing an environmental variable was considered to be equal to the rate of change in the score as inferred from Figure 8, hence a measure of how much the microorganism distribution differ along that variable.

In the analysis no. 2, the CCA produced four axes which accounted for 100% of the total variance in abundances of microbial OTUs among the glaciers. Figure 8 shows a biplot diagram of OTUs, glaciers, and environmental variables. The forward selection of variables demonstrated that the relationship between microbial communities and altitude (p = 0.02), area (p = 0.02), pH (p = 0.02), NO_2^- (p = 0.046), and NO_3^- (p = 0.048) were significant. The CANOCO program excluded NH_4^+ and salinity because they exhibited negligible variance. Subsequent analysis also demonstrated that altitude, area, and pH can explain the total community variability (analyses no. 3 and 4), meanwhile NO_2^- and NO_3^- are not so relevant (analysis no. 5, 6, and 7).

In summary, significant differences of the community structure between the four glaciers were visible. The comparison of relative abundances at the level of phyla also revealed significant differences (Figure 6; Table 4). In general, Viridiplantae, Fungi,
Table 3 | Diversity of small eukaryotes in clone libraries from glaciers.

| Taxon         | Mean no. of OTUs (clones) in glacier |
|---------------|-------------------------------------|
|               | ANE   | MAL  | PER  | LIT  |
| Viridiplantae  | 4 (80) | 2 (17) | 3 (38) | 1 (2) |
| Streptophyta   | 2 (34) |        |        |      |
| Chlorophyta    | 3 (72) | 2 (17) | 1 (4)  | 1 (2) |
| Environmental samples | 1 (8) |        |        |      |
| Stramenopiles  | 1 (2)  |        |        | 1 (8) |
| Chrysophyceae  |        |        | 1 (2)  |      |
| Bacillariophyta| 1 (8)  |        |        |      |
| Fungi          | 3 (12) | 2 (3)  | 1 (4)  | 1 (110) |
| Chytridomycota | 2 (10) | 1 (1)  | 1 (4)  |      |
| Dikarya        |        |        |        | 1 (110) |
| Environmental samples | 1 (2) | 1 (1)  |        |      |
| Rhizaria       | 2 (12) | 2 (13) | 4 (68) | 1 (16) |
| Cercozoa       | 2 (12) | 2 (13) | 4 (68) | 1 (16) |
| Metazoa        | 1 (4)  | 1 (2)  |        | 1 (8) |
| Rotifera       | 1 (4)  | 1 (2)  |        | 1 (8) |

Table 4 | Number of clones, number of OTUs, and diversity index for the samples from Pyrenean glaciers.

| Glacier | Mean no. of clones | Mean no. of OTUs | Diversity index H′ |
|---------|--------------------|------------------|--------------------|
| ANE     | 112                | 10               | 1.832              |
| MAL     | 35                 | 7                | 1.649              |
| PER     | 108                | 9                | 2.017              |
| LIT     | 44                 | 5                | 1.565              |
| $\rho^2$| 0.9905***          | 0.9688**         | 0.9955***          |
| Multiple comparison | ANE vs. PER*** | ANE vs. PER*** | ANE vs. PER*** |
|         | PER vs. LIT***     | PER vs. LIT***   | PER vs. LIT***    |
|         | MAL vs. PER***     | MAL vs. PER***   | MAL vs. PER***    |
|         | MAL vs. ANE***     | MAL vs. ANE***   | MAL vs. ANE***    |
|         | LIT vs. MAL***     | LIT vs. MAL***   | LIT vs. MAL***    |

Statistical differences were studied by ANOVA on the number of clones, number of OTUs and Shannon–Wiener index (**, p ≤ 0.001; ***, p ≤ 0.0001). Statistical significance was achieved by Newman-Keuls post-test (NS, not significant; *, p ≤ 0.05; **, p ≤ 0.01; ***, p ≤ 0.001).

and Rhizaria were positively influenced by altitude, area, and pH of glaciers, and negatively by salinity and NO$_2^−$ and NO$_3^−$ contents. These environmental characteristics were dominant in higher glaciers (Figure 8). On the contrary, samples collected at low altitudes, in LIT, contained a majority of Stramenopiles and Metazoa, and were positively influenced by salinity and NH$_4^+$, NO$_2^−$, and NO$_3^−$ contents (Figure 8).

DISCUSSION

Among the organisms that have successfully colonized extreme cold environments, a variety of survival mechanisms have been exploited. Microbial activity in ice is restricted to small amounts of unfrozen water inside the permafrost soil or the ice, and to brine channels. While there are sparse communities of lichens, mosses, and soil microorganisms, the limited availability of liquid water curtails biological activity for most of the year (Laybourn-Parry, 2002). Glaciers have truncated food chains with no animals or plants and a dominance of protozoa, bacteria, fungi, and microalgae. These microorganisms may be trapped during ice formation and remain inactive and frozen, but also active microorganisms live within the ice, being subjected to strong physical and chemical constraints. Contrary to what one might suppose, many of the microorganisms in glaciers do not cease to function in the winter months. Bacteria continue to grow all year, showing cycles that appear to be related to the availability of dissolved organic carbon.

![Rarefaction curves determined for the different 18S rRNA gene clones.](image)

![Richness estimates of Pyrenean glaciers at a genetic distance of 3%.](image)
They provide potential energy for a spectrum of heterotrophic and mixotrophic protozoans (Heath, 1988; Bell and Laybourn-Parry, 1999).

In cold environments, organisms are confronted by continuous low temperatures as well as a nutrient limitation. When compared to other known microorganisms, psychrophiles possess many unique qualities and molecular mechanisms that allow their adaptation to cold environments (Alcazar et al., 2010). In order to maintain activity in winter, microorganisms adopt one or more of a variety of strategies that enable them to enter the summer with actively growing populations. In this regard, some bacteria have been found to contain polyunsaturated fatty acids in their plasma membranes, which generally do not occur in other organisms. Further, some of them use enzymes that continue to function at near freezing ambient temperatures. And finally, they are able to produce proteins that are stable at cold temperatures (García-Descalzo et al., 2011). Among the Protozoa, many of the most successful species survive the winter in an active state by using endogenous energy reserves or employing nutritional versatility. Mixotrophy is also an important nutritional strategy (Laybourn-Parry, 2002). It involves a combination of autotrophy and heterotrophy in varying degrees. Some protozoa are forced to sustain a mixotrophic strategy and cannot survive by photosynthesis alone. The dependence on ingesting bacteria varies seasonally. One argument suggests that it is a means of acquiring inorganic nutrients for photosynthesis during phases of limitation (Nygaard and Tobiesen, 1993). Other researchers contend that it is a means of supplementing the carbon budget (Jones et al., 1993).

According to our results, the taxonomic affiliation of the eukaryotic sequences associated to the samples from higher Pyrenean glaciers was markedly different from that of lower glacier. Ice in samples from higher glaciers was characterized by a clear dominance of Viridiplantae, fundamentally Chloromonas, and Rhizaria (Figures 5 and 8) and these glaciers were quite similar in terms of relative abundance of phyla with typical glacier protist lineages, most notably Chlorophyta, Streptophyta, and Cercozoa (Cameron et al., 2012). Generally the less thawed areas, located in ANE, present a majority of Viridiplantae. Probably, ice maintains microorganisms isolated in clusters where they must survive as photosynthetic and primary producers. However, frequently thawed areas host a majority of Rhizaria that are heterotroph. In this case, microorganisms are able to move into the meltwater, reaching their preys. It is interesting to comment that although PER conserves an intermediate extension of ice, it presents a majority of Rhizaria, what could be due to the high level of fragmentation observed in the mass of ice which originates freeze-thaw cycles.

In this study, the dominant taxa were present in all samples, and corresponded roughly with those reported in other studies regarding protist community composition in ice samples (Bachy et al.,...
Table 5 | Correlation analysis between taxa of small eukaryotes from glaciers and environmental variables.

| Taxon       | Altitude | Area | Salinity | pH  | NH₄⁺ | NO₂⁻ | NO₃⁻ |
|-------------|----------|------|----------|-----|------|------|------|
| **Fungi**   |          |      |          |     |      |      |      |
| r           | 0.4981   | 0.8541 | −0.04913 | 0.8843 | −0.9670 | −0.4966 | −0.4940 |
| p           | 0.5019   | 0.1459 | 0.4754   | 0.1157 | 0.0165* | 0.5034 | 0.5060 |
| **Metazoa** |          |      |          |     |      |      |      |
| r           | −0.4777  | 0.0033 | 0.7839   | 0.8467 | −0.6172 | 0.4673 | 0.4697 |
| p           | 0.5223   | 0.9967 | 0.1086   | 0.1533 | 0.1914 | 0.5327 | 0.5303 |
| **Rhizaria**|          |      |          |     |      |      |      |
| r           | 0.7167   | 0.3152 | −0.9594  | −0.6392 | 0.3679 | −0.7460 | −0.7480 |
| p           | 0.2833   | 0.6848 | 0.0203*  | 0.3608 | 0.3161 | 0.2540 | 0.2520 |
| **Viridiplantae** |     |      |          |     |      |      |      |
| r           | 0.9654   | 0.9532 | −0.6558  | 0.2951 | −0.5976 | −0.9144 | −0.9132 |
| p           | 0.0346*  | 0.0468* | 0.1721  | 0.7049 | 0.2012 | 0.0856 | 0.0888 |
| **Stramenopiles** |     |      |          |     |      |      |      |
| r           | −0.3209  | −0.6954 | 0.05431  | −0.8066 | 0.8018 | 0.4118 | 0.4099 |
| p           | 0.6791   | 0.3046 | 0.4728   | 0.1934 | 0.0991 | 0.5882 | 0.5901 |
| **Total**   |          |      |          |     |      |      |      |
| r           | 0.9926   | 0.9223 | −0.7566  | 0.1718 | −0.4871 | −0.9576 | −0.9588 |
| p           | 0.0074** | 0.0777 | 0.1217   | 0.8282 | 0.2564 | 0.0424* | 0.0432* |

Statistical differences were studied by Pearson’s r (*p ≤ 0.05; **p ≤ 0.01).

In general, our knowledge about microbial eukaryotic communities in glaciers is quite limited. There are not many similar reports to compare, but overall abundance and biodiversity in our samples appear to be low, which may be due to the structure of ice, that keeps the microorganisms isolated. LIT is the only one with representatives of all phyla, since its deglaciation may allow greater exchange of macroorganisms. Samples were collected in summer, when iced areas were minimal, and LIT virtually becomes a lake.

The retreat of glaciers is related to their geographical features such as altitude and area, while other environmental variables such as pH is dependent of mineral salts solubility, which in turn increases with temperature. Moreover, pH may depend on the chemical composition of the soil. We cannot determine whether pH has a direct or indirect effect on community composition, as a number of ice properties (e.g., salinity) are directly or indirectly related to pH. Thus, the effect of a number of different factors is...
Table 6 | Summary of correspondence analysis and eigenvalues (λ).

| No. of | Type of | Environmental variables | Environmental covariables | $\lambda_1$ | $\lambda_2$ | $\lambda_3$ | $\lambda_4$ |
|--------|---------|-------------------------|--------------------------|------------|------------|------------|------------|
| 1 | PCA | – | – | 0.612 | 0.310 | 0.074 | 0.003 |
| 2 | CCA | Altitude, area, NO$_2^-$, NO$_3^-$, pH, salinity | – | 1.000 | 0.798 | 0.200 | 0.003 |
| 3 | CCA | Altitude, area | – | 0.835 | 0.780 | 0.383 | 0.006 |
| 4 | CCA | Altitude, area, pH | – | 1.000 | 0.798 | 0.200 | 0.003 |
| 5 | CCA | Altitude, area, pH | – | 0.842 | 0.094 | 0.323 | 0.005 |
| 6 | CCA | Altitude, area, NO$_2^-$ | – | 0.956 | 0.798 | 0.198 | 0.007 |
| 7 | CCA | Altitude, area, NO$_2^-$ | – | 0.798 | 0.198 | 0.007 | 0.005 |

FIGURE 8 | Canonical correspondence analysis. Ordination diagram based on CCA, with respect to seven quantitative variables. The axes are scaled in standard deviation units. Eigenvalues for the axes are detailed in Table 6. The diagram displays triangles that represent phyla, circles representing sampling sites, and arrows that symbolize environmental variables.

reflected by ice pH and these factors may also drive community composition. Further, PER is placed in a calcareous massif, while the other three glaciers are located in a granite massif (Arenillas et al., 2008). However, in terms of biodiversity, the three highest glaciers are similar, regardless of their geochemical characteristics.

Climate change is one of the most important problems that concern modern society. Until now, scarce data are available to evaluate the environmental impact on living organisms due to the climate change. It is necessary to develop a rigorous investigation before undertaking any intervention, and that is why accurate data are needed about possible responses from living organisms to climate change. It is important to know how microbial patterns are being altered and how these changes are affecting to climate, as these microorganisms are essential components of the microbial food webs and are often dominant primary producers. Although the full range of ecological implications remains poorly understood, changes at the base of food webs necessarily entail consequences for higher trophic levels, while modifying the biochemical cycling of major elements including, but not limited to carbon, nitrogen, phosphorous, and silicon (Hobbs et al., 2010). The evaluation of the effect of global warming at the microbial community level is a difficult task because of the many mutually dependent response variables, non-linear responses, etc. These difficulties can be partially overcome using CCA, which enables an evaluation of the influence of the environment on the composition of the community and provides a distribution-free Monte Carlo test of significance (Jongman et al., 1995). The analysis of protist diversity in Pyrenean glaciers revealed statistically significant differences in protist diversity and community structure between four glaciers melted to different extent. The analysis of influences of ice chemical properties on protist community structure revealed that pH had the strongest effect on protist community structure of the analyzed ice properties. NO$_2^-$ and NO$_3^-$ contents appear to have a minor impact on psychrophilic protist community structure and diversity. In this survey, the correlations between ice melting and community composition were obvious.

The effect of altitude and glacier area on the composition of the microbial community is essential, as demonstrated by CCA analysis (Figure 8). This analysis also demonstrated that it is effectively possible to discriminate between the effect on microbial community of area/altitude and the effect of chemical parameters. Chemical composition differences appear associated to ice melting in lower glaciers, in which NO$_2^-$ and NO$_3^-$ contents are correlated to community composition.

Further work is needed to evaluate whether our observations can be generalized to other glacial regions. The retreat of mountain glaciers subsequent to ongoing climate change has been documented extensively in the last years (Méter, 1984; Gómez et al., 2003; Paul et al., 2014; Zemp et al., 2006; Citterio et al., 2007). Several studies developed analytical models to forecast the retreating trend of glaciers in the future. They predicted that a consistent loss (or even the complete extinction) of most of the ice masses will be observed by the end of this century in which case should the current climate trend continue, and concluded that small glaciers of southern Europe will be among the most reliable witnesses of global warming (Oerlemans, 1997; Zuo and Oerlemans, 1997; Zemp et al., 2006). Accordingly, such southern glaciers can be considered important for studying climate and environmental changes occurring in the Mediterranean region.
Future studies will be useful to better understand the impact of global warming on microbial communities that behave as bio- sensors. In our results, two main groups act as indicators of glacier retreat. Stramenopiles and Metazoa, meanwhile Viridiplan- teae is the dominant group in the less melted glaciers. There are still many unanswered questions relating to the physiology and biochemistry of surviving the cold and adaptation to warming, particularly as there is a huge biotechnological potential in such psychrophiles.

 Canonical correspondence analysis (CCA) was performed to analyze the relationships between the observed taxa in Table 2 and environmental variables in Table 1. Seven environmental variables were taken into account. Two of them, altitude and glacier area are related to the glacier degree of conservation, as ice melting is less dramatic in higher glaciers, which maintain wider areas of ice. The rest of environmental variables, pH, salinity, NH₄⁺, NO₃⁻, and NO₂⁻ are associated to the ice chemical composition. To separate the sources of variability in the microbial community various combinations with different environmental variables were applied (Table 6).

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