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Metagenomic and satellite analyses of red snow in the Russian Arctic

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Cryophilic algae thrive in liquid water within snow and ice in alpine and polar regions worldwide. Blooms of these algae lowers albedo (reflection of sunlight), thereby altering melting patterns (Kohshima et al. 1993; Lutz et al. 2014; Thomas & Duval 1995) . Here metagenomic DNA analysis and satellite imaging were used to investigate red snow in Franz Josef Land in the Russian Arctic. Franz Josef Land red snow metagenomes confirmed that the communities are composed of the autotroph *Chlamydomonas nivalis* that is supporting a complex viral and heterotrophic bacterial community. Comparisons with white snow communities from other sites suggest that snow and ice are initially colonized by fungal-dominated communities and then succeeded by the more complex *C. nivalis*-heterotroph red snow. Satellite image analysis showed that red snow covers up to 80% of the surface of snow and ice fields in Franz Josef Land and globally. Together these results show that *C. nivalis* supports a local food web that is on the rise as temperatures warm, with potential widespread impacts on alpine and polar environments worldwide.

Metagenomic and Satellite Analyses of Red Snow in the Russian Arctic

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Keywords: red snow, snow, watermelon snow, Franz Josef Land, arctic, metagenomics, viruses, phage

1

9 Abstract

Cryophilic algae thrive in liquid water within snow and ice in alpine and polar regions 10 worldwide. Blooms of these algae lowers albedo (reflection of sunlight), thereby altering melting 11 patterns (Kohshima et al. 1993; Lutz et al. 2014; Thomas & Duval 1995). Here metagenomic 12 13 DNA analysis and satellite imaging were used to investigate red snow in Franz Josef Land in the 14 Russian Arctic. Franz Josef Land red snow metagenomes confirmed that the communities are composed of the autotroph *Chlamydomonas nivalis* that is supporting a complex viral and 15 16 heterotrophic bacterial community. Comparisons with white snow communities from other sites 17 suggest that snow and ice are initially colonized by fungal-dominated communities and then succeeded by the more complex C. nivalis-heterotroph red snow. Satellite image analysis showed 18 that red snow covers up to 80% of the surface of snow and ice fields in Franz Josef Land and 19 20 globally. Together these results show that C. nivalis supports a local food web that is on the rise as temperatures warm, with potential widespread impacts on alpine and polar environments 21 worldwide. 22

24 Introduction

Chlamydomonas nivalis is a unicellular snow alga that has been detected worldwide within the 25 upper snow layer in polar and alpine regions (Guiry et al. 2014) and is especially abundant in the 26 Arctic pack ice (Gradinger & Nurnberg 1996). In these harsh environments, C. nivalis has 27 adapted to intense UV exposure by producing astaxanthin, a UV-screening pigment that 28 produces a visible red hue in snow (Gorton & Vogelmann 2003; Williams et al. 2003). C. nivalis 29 spends most of its growth season in its red colored stage (Gorton & Vogelmann 2003; Stibal et 30 al. 2007; Williams et al. 2003); this coloration is visible across the snow/ice surface and can 31 reduce albedo to 40% [c.f., fresh snow albedo of 75% (Thomas & Duval 1995)]. The lower 32 albedo increases local temperature, promoting snow and ice melting and increasing the 33 34 abundance of *C. nivalis* (Thomas & Duval 1995). Through this positive feedback loop the abundance of C. nivalis amplifies snow and ice melting. C. nivalis may also contribute to CO2 by 35 fixing carbon. However, if there is a red snow associated heterotrophic viral and microbial 36 community, much of this newly fixed carbon may be released via respiration (Bardgett et al. 37

2008). *C. nivalis*-produced dissolved organic carbon (DOC) could also contribute to priming
when the melt water washes into the ocean (Geller 1986; Hamer & Marschner 2002; Madigan et
al. 1997; van Nugteren et al. 2009) and lead to increased CO₂ release.

41

42 Materials and Methods

Analysis of satellite images: Remote sensing methods were used to estimate abundances
 of red snow at eleven locations around the world (see Supplementary Materials). Landsat
 satellite images were acquired from the USGS Earth Explorer site

(http://earthexplorer.usgs.gov/) and image analysis methods were adapted from Takeuchi et al. 46 (Takeuchi et al. 2006) as described in the Supplementary Materials. Red to green reflectance 47 band ratios with wavelengths 630 - 690 nanometers and 520 - 600 nanometers, respectively, 48 49 were used to detect red snow in the satellite images. The spectral reflectance of red snow shows that it has higher reflectance in the red band than in the green band, while the spectral reflectance 50 51 of snow and ice has higher reflectance in the green band than the red band (Takeuchi et al. 2006). 52 Therefore, red to green reflectance band ratios that are less than 1.0 are more likely to signify 53 white snow or ice while band ratios that are greater than 1.0 are more likely to signify red snow or ice. 54

55 ArcGIS version 10.2 was used to calculate the reflectance band ratios. Previous research indicates areas with reflectance band ratios > 1.02 are bright red when observed in the field 56 57 (Takeuchi et al. 2006). For this analysis, areas with reflectance band ratios greater than 1.0 were 58 considered to have a significant amount of red snow because such values have been shown to have an algal cell volume of 100 ml m⁻² (Takeuchi et al. 2006). Using the positive linear 59 correlation between algal cell volume biomass and reflectance band ratio, it was assumed that the 60 higher the reflectance band ratio, the higher the algal cell volume biomass. With this in mind, the 61 62 reflectance band ratios were divided into five categories for optimal visualization of various levels of concentrations of red snow (Supplementary Table 1 and Figure 1B). 63

Algal biomass: To estimate the algal biomass for each location, the surface area
belonging to each reflectance band ratio category was multiplied by the mean algal biomass of
that category. Although the extent of the area of interest is the same for all three images, they
have varying amounts of surface area where red snow can exist due to shifts in snow and ice
coverage. Therefore, in addition to the total algal biomass, the total area of snow coverage and

the percentage of the total area of snow that was covered with different abundances of algae were calculated. A pixel was categorized as snow if its normalized difference snow index (NDSI) was greater than 0.4 and, to mask out water, if its near-infrared reflectance value was greater than 0.11 (Sibandze et al. 2014). The number of pixels that meet these conditions was multiplied by the area of the pixel to get the total area of ground covered by snow/ice. To calculate the percentage of the total area of snow that is covered with algae, the total area with each algal abundance level was divided by the total area of snow coverage.

76 *Metagenomic sequencing:* Red snow samples were examined with microscopy to confirm the presence of C. nivalis based on morphology (Muller et al. 1998). Three red snow samples of 77 ~15 L were collected, melted, and passed through a 0.22 µm sterivex filter. Total DNA was 78 79 extracted in the field using the Soil DNA Isolation kit with a custom vacuum manifold (Norgen 80 BioTek Corp., cat# 26560). From the total DNA, a NexteraXT library kit was used to prepare DNA libraries for sequencing on the Illumina MiSeq. The libraries were named Nansen (135,749 81 82 reads), Greely 1 (86,932 reads) and Greely 2 (47,507 reads). Each metagenome was passed through the following quality control pipeline. PrinSeq was used to quality filter reads below 100 83 84 bp in length and below an average quality score of 25, and to remove duplicates and sequence tags (Schmieder & Edwards 2011b). Reads assigned as human were removed using DeconSeq 85 (Schmieder & Edwards 2011a). Post quality control, the Nansen library contained 121,455 reads, 86 Greely 1 contained 69,918 reads, and Greely 2 contained 40,344 reads. Seven publicly 87 88 accessible snow metagenomes from Svalbard glaciers (a.k.a., white snow throughout manuscript) sampled April through June were downloaded from MG-RAST and reads were quality filtered 89 using the same pipeline as the FJL red snow libraries (Maccario et al. 2014). The red snow and 90 white snow libraries were compared to the KEGG database to assign reads to KEGG pathways 91 92 (e-value $< 1 \times 10^{-5}$; >60% identity; >15 aa minimum alignment length). Estimations of taxonomic 93 composition of communities were based on translated comparisons to the non-redundant protein database M5NR (e-value $< 1 \times 10^{-5}$; >60% identity; >15 as minimum alignment length). The 94 dataset was normalized to ensure similar numbers of reads were used for each sample, and then 95 raw read counts were log transformed. Statistical differences between red snow and snow in the 96 97 numbers of reads assigned to KEGG pathway groups were identified by ANOVA. Multivariate statistics were performed in R using the vegan (Dixon 2003), clustsig and the stats packages. The 98 adonis function was used to compare metagenome compositions; vegdist was used to generate 99

distance matrices; *simprof* was used to cluster metagenomes based on similarity; and *prcomp*was used to perform Principal Component Analysis.

102

103 **Results and Discussion**

Detection of red snow in a global sample of satellite images: Satellite images with 104 spectral reflectance data were used to approximate snow and ice cover, as well as red algae 105 abundance (Takeuchi 2009; Takeuchi et al. 2006) over several years in Franz Josef Land, as well 106 107 as eleven other regions of United States, Canada, Greenland, Norway, Austria, India, and New Zealand (Supplementary Figure 1). Red snow was detected at all eleven locations in all the years 108 (Figure 1A). The total area of snow and ice were lowest in the most recent year (2013, 2014 or 109 2015, depending on the location; Supplementary Figure 2; Greenland was the exception to this 110 111 trend). At least 50% of the total snow/ice area was covered with algae for the most recent year analyzed (Supplementary Figure 2; exception New Zealand and Franz Josef Land). In seven of 112 113 the locations, over 80% of the total snow and ice fields were covered in red algae in the most recent year analyzed (Supplementary Figure 2). 114

115 We performed a walking transect from sea level to the glacier on Nansen Island in Franz Josef Land in August 2013 (to be described in a separate manuscript). Therefore, this region was 116 117 targeted for more detailed analysis. Around and on Nansen, the total red snow algal biomass 118 increased by 124% from 1986 to 2002 and by 15% from 2002 to 2006, then decreased by 63%119 from 2006 to 2015 (Figure 1B). These changes in algal cover co-occurred with a total decline in the snow and ice cover (Figure 1B). Visual inspection of the snow and ice on Nansen Island in 120 August of 2013 confirmed the presence of red colored snow and microscopy of red snow 121 samples showed C. nivalis cells. Taken together, these results show that even as total snow and 122 123 ice cover declines, red snow cover is still highly prevalent or increasing both in Franz Josef Land 124 and other alpine/polar regions.

Microbes present in snow and red snow: For metagenomic sequencing, one and two red snow samples were taken from Nansen and Greeley Islands, respectively. Seven white snow metagenomes from Svalbard glaciers were also downloaded and analyzed for comparison (see Methods & Supplementary Materials). The genus-level taxonomic compositions of white snow and red snow were significantly different (ADONIS; F = 4.567; p = 0.007). When samples were clustered according to their taxonomical similarities, one red snow sample taken at Greely Island grouped with a Svalbard glacier sample; otherwise the red snow and white snow samples
clustered separately (Supplementary Figure 3). This indicates minimal overlap in microbial
composition at the genus level.

Community DNA sequences were further compared using multivariate analyses with the 134 top 10 most variable taxa (Supplementary Figure 4). The first two principal components 135 explained 70% of the between-sample variation in microbial community members. The first 136 principal component described red snow as having higher abundances of species from the 137 138 bacterial genera Pseudoalteromonas, Alteromonas, Vibrio, and Pedobacter, whereas white snow had higher abundances of species from the eukaryotic genera Aspergillus and Neurospora, as 139 well as the bacterial genera Nostoc, Bacillus and Spirosoma. Red snow had greater overall 140 abundances of Bacteria and viruses (Figure 2A) and a lower abundance of Eukaryotes (Figure 141 142 2A). The bacterial communities associated with red snow have also been observed in an alpine region (Thomas & Duval 1995) and are probably support by photosynthate from the C. nivalis. 143 144 Evidence also suggests that bacterial cells may physically attach to the outer mucilaginous coating of C. nivalis in red snow forming an arctic holobiont (Remias et al. 2005; Thomas & 145 146 Duval 1995).

The metagenomes were also used to verify the presence of *Chlamydomonas* in snow samples (Supplementary Figure 5). Of the sequence reads assigned to Eukaryotes, the proportion of reads assigned to the *Chlamydomonas*-containing phylum Chlorophyta was higher in red snow than white snow (Figure 2B). Conversely, the proportion of reads assigned to the fungal phylum Ascomycota was higher in white snow (Figure 2B).

Functional capabilities of microbial communities in red snow and snow: The 152 metagenomes were also analyzed for potential metabolic functions. The functions encoded by 153 154 the red and white snow samples clustered into 8 significant groups, with the red snow samples 155 from Greely and Nansen Islands forming a significant cluster (Supplementary Figure 6). Four white snow samples formed a cluster and the remaining snow samples clustered individually. 156 Multivariate analysis of the top 10 most variable functions showed that the first two principal 157 components explained 82% of the variation in the abundances of functional pathways 158 159 (Supplementary Figure 7). The first component (70% of the variation) showed that the red snow had higher abundances of genes involved in membrane transport, carbohydrate metabolism, 160 nucleotide and amino acid synthesis/degradation, and energy metabolism. White snow 161

communities were shifted toward cell growth and death, folding sorting and degradation,
 transcription, transport and catabolism pathways and pathways annotated as important in
 neurodegenerative diseases (i.e., mitochondrial functions in Eukaryotes).

In order to examine whether microbial communities in red snow encoded completely 165 different functional capabilities from those in snow, the numbers of reads assigned to all KEGG 166 pathways were compared using a matrix of Bray-Curtis dissimilarities. Overall the abundances 167 of KEGG pathways were not significantly different between red snow and white snow 168 (ADONIS; F = 2.135; p = 0.12). However, separate analyses that compared individual pathways 169 between red snow and white snow identified several pathways as significantly different, 170 including pathways related to sugar biosynthesis and metabolism and energy metabolism. Red 171 snow communities had higher abundances of genes that encode lipopolysaccharide biosynthesis 172 173 and peptidoglycan biosynthesis. Red snow also had a higher proportion of reads assigned to oxidative phosphorylation, methane metabolism, carbon fixation in photosynthetic organisms 174 175 and carbon fixation pathways (Figure 3B). White snow had higher relative abundances of genes that encode glycan biosynthesis and related pathways such as GPI-anchor biosynthesis, other 176 177 types of O-glycan biosynthesis and various types of N-glycan biosynthesis (Figure 3A). Taken together, these results support the hypothesis that red snow communities include primary 178 179 producers with a large, heterotrophic community including viruses. In contrast, white snow communities appear to be dominated by fungi, maybe eating refractory organic carbon delivered 180 181 with the snow (Clarke & Noone 1985; Rosen et al. 1981; Thevenon et al. 2009).

182

183 Conclusions

Microbiology of snow and ice fields has a long history, including a reference to red snow 184 185 by Aristotle. However, until now we have not had the tools to determine the full extent and 186 makeup of these communities. Here we use a combination of satellite and metagenomic approaches to show that red snow covers up to 80% of the examined ice and snow fields. 187 Metagenomics of red snow from Franz Josef Land, one of the most remote polar land masses in 188 world, show that these communities support a full food web ranging from algae to heterotrophic 189 190 microbes to viruses. Because of the reduced albedo associated with these communities, red snow creates a positive feedback loop that increases its abundance while simultaneously melting ice 191 and snow (Figure 4). In addition to the direct effects on sunlight absorbance, the heterotrophic 192

- 193 activity (including viral lysis) will increase local temperatures. Together, these effects may
- 194 significantly increase ice and snow melting in the Barents Sea region that is already one of the
- 195 fastest-warming regions on earth. Projections for global red snow coverage and its influence on
- 196 warming patterns should be investigated further.
- 197

205

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207 References

- 208
- Bardgett RD, Freeman C, and Ostle NJ. 2008. Microbial contributions to climate change through
 carbon cycle feedbacks. *Isme Journal* 2:805-814. DOI 10.1038/ismej.2008.58
- Clarke AD, and Noone KJ. 1985. Soot in the Arctic Snowpack a Cause for Perturbations in
 Radiative-Transfer. *Atmospheric Environment* 19:2045-2053. Doi 10.1016/0004 6981(85)90113-1
- Dixon P. 2003. VEGAN, a package of R functions for community ecology. *Journal of Vegetation Science* 14:927-930. DOI 10.1111/j.1654-1103.2003.tb02228.x

Gorton HL, and Vogelmann TC. 2003. Ultraviolet radiation and the snow alga Chlamydomonas
 nivalis (Bauer) Wille. *Photochemistry and Photobiology* 77:608-615. Doi 10.1562/0031 8655(2003)077<0608:Uratsa>2.0.Co;2

Gradinger R, and Nurnberg D. 1996. Snow algal communities on Arctic pack ice floes dominated by *Chlamydomonas nivalis* (Bauer) Wille. *Proceedings of the National Institute of Polar Research Symposium on Polar Biology* 9:35-43.

Guiry MD, Guiry GM, Morrison L, Rindi F, Valenzuela Miranda S, Mathieson AC, Parker BC,
Langangen A, John DM, Barbara I, Carter CF, Kuipers P, and Garbary DJ. 2014. AlgaeBase:
an on-line resource for Algae. *Cryptogamie Algologie* 35:105-115. DOI
10.7872/crya.v35.iss2.2014.105

Hamer U, and Marschner B. 2002. Priming effects of sugars, amino acids, organic acids and
catechol on the mineralization of lignin and peat. *Journal of Plant Nutrition and Soil Science- Zeitschrift Fur Pflanzenernahrung Und Bodenkunde* 165:261-268. Unsp [P88/4b]

231 Doi 10.1002/1522-2624(200206)165:3<261::Aid-Jpln261>3.0.Co;2-I

Kohshima S, Seko K, and Yoshimura Y. 1993. Biotic Acceleration of Glacier Melting in Yala
 Glacier, Langtang Region, Nepal Himalaya. *Snow and Glacier Hydrology*:309-316.

Lutz S, Anesio AM, Villar SEJ, and Benning LG. 2014. Variations of algal communities cause
 darkening of a Greenland glacier. *Fems Microbiology Ecology* 89:402-414. Doi
 10.1111/1574-6941.12351

- 237 Maccario L, Vogel TM, and Larose C. 2014. Potential drivers of microbial community structure
- and function in Arctic spring snow. *Frontiers in Microbiology* 5. ARTN 413
- 239 DOI 10.3389/fmicb.2014.00413
- Madigan MT, Martinko JM, and Parker J. 1997. *Brock Biology of Microorganisms*: Prentice Hall, Upper Saddle River, NJ.

221

Geller A. 1986. Comparison of Mechanisms Enhancing Biodegradability of Refractory Lake
 Water Constituents. *Limnology and Oceanography* 31:755-764.

Muller T, Bleiss W, Martin CD, Rogaschewski S, and Fuhr G. 1998. Snow algae from northwest
 Svalbard: their identification, distribution, pigment and nutrient content. *Polar Biology* 20:14 DOI 10.1007/s003000050272

Remias D, Lutz-Meindl U, and Lutz C. 2005. Photosynthesis, pigments and ultrastructure of the
 alpine snow alga Chlamydomonas nivalis. *European Journal of Phycology* 40:259-268. Doi
 10.1080/09670260500202148

- Rosen H, Novakov T, and Bodhaine BA. 1981. Soot in the Arctic. *Atmospheric Environment* 15:1371-1374. Doi 10.1016/0004-6981(81)90343-7
- Schmieder R, and Edwards R. 2011a. Fast Identification and Removal of Sequence
 Contamination from Genomic and Metagenomic Datasets. *Plos One* 6. ARTN e17288
 DOI 10.1371/journal.pone.0017288
- Schmieder R, and Edwards R. 2011b. Quality control and preprocessing of metagenomic
 datasets. *Bioinformatics* 27:863-864. DOI 10.1093/bioinformatics/btr026

Sibandze P, Mhangara P, Odindi J, and Kganyago M. 2014. A comparison of Normalised Difference Snow Index (NDSI) and Normalized Difference Principal Component Snow Index (NDPCSI) techniques in distinguishing snow from related land cover types. . *South African Journal of Geomatics* 3:197-209.

Stibal M, Elster J, Sabacka M, and Kastovska K. 2007. Seasonal and diel changes in
 photosynthetic activity of the snow alga Chlamydomonas nivalis (Chlorophyceae) from
 Svalbard determined by pulse amplitude modulation fluorometry. *Fems Microbiology Ecology* 59:265-273. DOI 10.1111/j.1574-6941.2006.00264.x

Takeuchi N. 2009. Temporal and spatial variations in spectral reflectance and characteristics of
 surface dust on Gulkana Glacier, Alaska Range. *Journal of Glaciology* 55:701-709.

Takeuchi N, Dial R, Kohshima S, Segawa T, and Uetake J. 2006. Spatial distribution and
 abundance of red snow algae on the Harding Icefield, Alaska derived from a satellite image.
 Geophysical Research Letters 33. Artn L21502

268 Doi 10.1029/2006gl027819

- Thevenon F, Anselmetti FS, Bernasconi SM, and Schwikowski M. 2009. Mineral dust and
 elemental black carbon records from an Alpine ice core (Colle Gnifetti glacier) over the last
- 271 millennium. Journal of Geophysical Research-Atmospheres 114. Artn D17102
- 272 10.1029/2008jd011490
- Thomas WH, and Duval B. 1995. Sierra-Nevada, California, USA, Snow Algae Snow Albedo
 Changes, Algal Bacterial Interrelationships, and Ultraviolet-Radiation Effects. *Arctic and Alpine Research* 27:389-399. Doi 10.2307/1552032

van Nugteren P, Moodley L, Brummer GJ, Heip CHR, Herman PMJ, and Middelburg JJ. 2009.
 Seafloor ecosystem functioning: the importance of organic matter priming. *Marine Biology* 156:2277-2287. 10.1007/s00227-009-1255-5

- 279 Williams WE, Gorton HL, and Vogelmann TC. 2003. Surface gas-exchange processes of snow
- algae. Proceedings of the National Academy of Sciences of the United States of America
- 281 100:562-566. DOI 10.1073/pnas.0235560100

282

284 Figure Legends

285

Figure 1. (A) A time series comparison of the percentage of total snow or ice that is covered with algae at selected alpine and polar regions throughout the world, according to data derived from satellite images. (B) A time series comparison of the total area of snow and sea ice, total algal biomass, and percentage of total snow that is covered with algae within the map extent near Nansen Island, Franz Josef Land, for years 1986, 2002, 2006 and 2015. The colored time series shows spatial distribution maps of algal densities of the Nansen Island area in Franz Josef Land.

Figure 2. (A) Abundances of viruses, Bacteria and Eukaryotes in samples from red snow and snow communities. The y-axis shows abundances after normalizing and standardizing raw read counts to ensure cross-sample comparisons are valid. (B) Bar plots showing abundances of two Eukaryotic phyla found in red snow and snow communities. Chlorophyta is the phylum that contains the genus *Chlamydomonas*.

Figure 3. (A) Bar plots showing functional pathways that were statistically significantly
different in abundance between red snow and snow. (B) Bar plots depicting energy metabolism
pathways and their abundances in red snow and snow.

Figure 4. Model of *C. nivalis* in white and red snows. The left panel shows several microbial communities that are found in white snow. Sunlight promotes astaxanthin expression in *C. nivalis*, turning the snow to red and promoting community metabolism shifts through stimulation of heterotrophic metabolism. The *C. nivalis* blooms, albedo is decreased and local snow and ice melts at a faster rate (right panel).

308 309

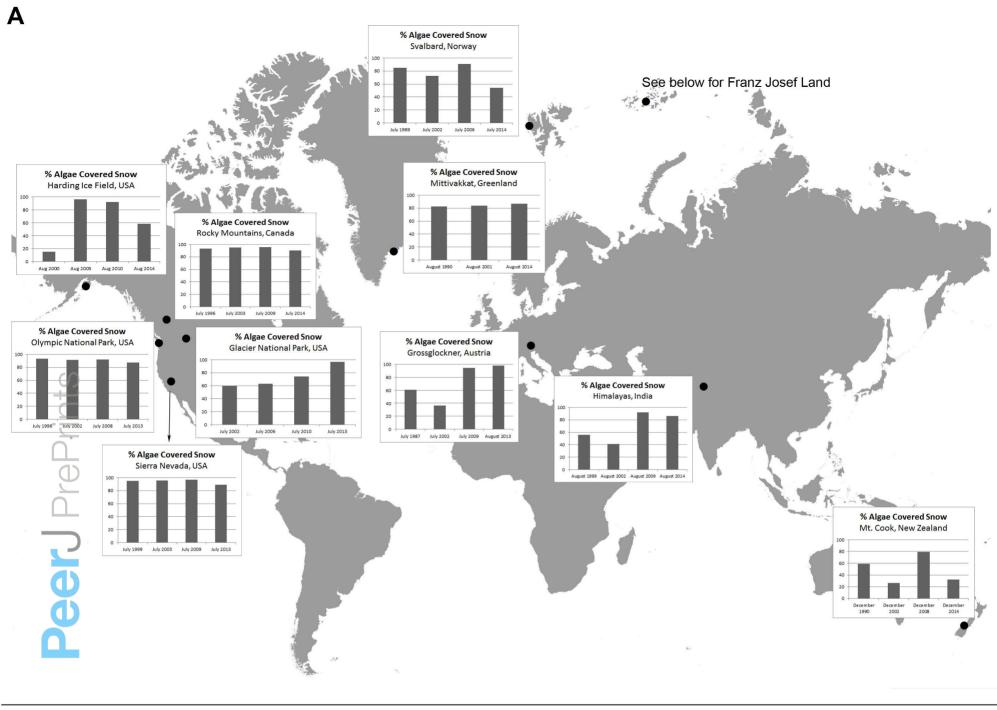
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298

Figure 1(on next page)

Changes in red snow through time

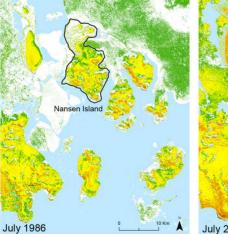
(A) A time series comparison of the percentage of total snow or ice that is covered with algae at selected alpine and polar regions throughout the world, according to data derived from satellite images. (B) A time series comparison of the total area of snow and sea ice, total algal biomass, and percentage of total snow that is covered with algae within the map extent near Nansen Island, Franz Josef Land, for years 1986, 2002, 2006 and 2015. The colored time series shows spatial distribution maps of algal densities of the Nansen Island area in Franz Josef Land.

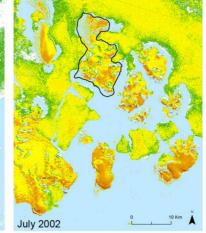


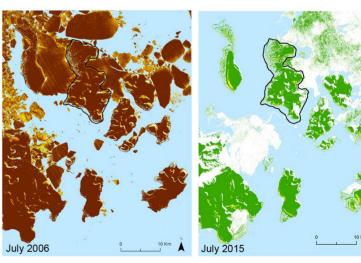




White snow or ice Low (100 - 168.75) Medium (168.75 - 237.5) High (237.5 - 300) Very high (> 300)







July 2015

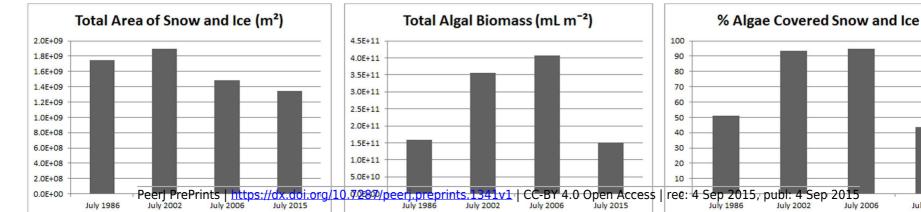
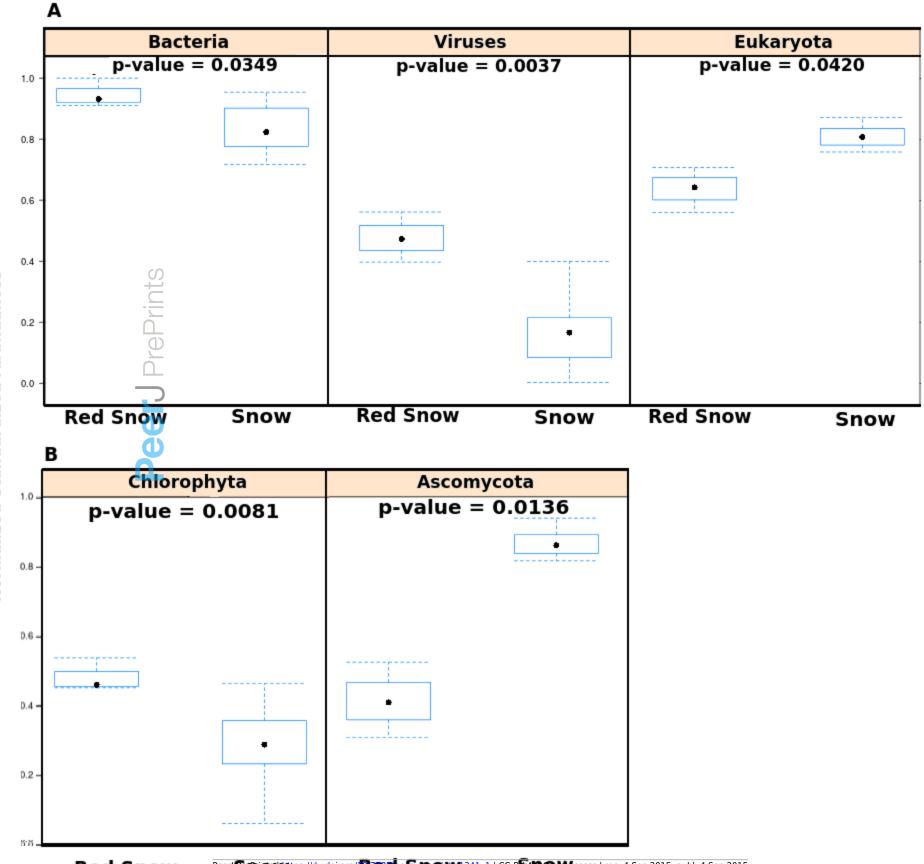


Figure 2(on next page)

Abundances of microbes in red snow and white snow samples

(A) Abundances of viruses, Bacteria and Eukaryotes in samples from red snow and snow communities. The y-axis shows abundances after normalizing and standardizing raw read counts to ensure cross-sample comparisons are valid. (B) Bar plots showing abundances of two Eukaryotic phyla found in red snow and snow communities. Chlorophyta is the phylum that contains the genus *Chlamydomonas*.



Normalized Standardized Abundances

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Figure 3(on next page)

Functional pathways in red snow and white snow

(A) Bar plots showing functional pathways that were statistically significantly different in abundance between red snow and snow. (B) Bar plots depicting energy metabolism pathways and their abundances in red snow and snow.

GPI-anchor biosynthesis Types of O-glycan biosynthesis Peptidoglycan biosynthesis Types of N-glycan biosynthesis Lipopolysaccharide biosynthesis p-value = 0.023 p-value = 0.0005 p-value = 0.017 p-value = 0.003 p-value = 0.006 1.0 0.8 0.6 Standardized Abundances . nts ٠ -----1 В Oxidative phosphorylation Carbon fixation pathways Carbon fixation in Methane metabolism Sulfur metabolism photosynthetic organisms in prokaryotes p-value = 0.045 p-value = 0.0110 p-value = 0.0082 p-value = 0.0846 p-value = 0.0011 Normalized 0.8 0.6 -----_____ 0.4 -0.2 -D.0 -Red Snow PeerJ PrePS now dx.doi.org/10.7 Redinsmow1v1 | CC-BY 4S Prowss | rec: 4 Sep Redins Prow **Red Snow Red Snow** Snow Snow Snow

Figure 4(on next page)

Model of red snow microbiology

Model of *C. nivalis* in white and red snows. The left panel shows several microbial communities that are found in white snow. Sunlight promotes astaxanthin expression in *C. nivalis*, turning the snow to red and promoting community metabolism shifts through stimulation of heterotrophic metabolism. The *C. nivalis* blooms, albedo is decreased and local snow and ice melts at a faster rate (right panel).

SNOW

RED SNOW

