Sea ice-pelagic-benthic links of bacterial diversity during the Arctic summer sea ice record minimum in 2012 Josephine Z. Rapp^{1,2*}, Mar Fernández-Méndez^{1,2}, Christina Bienhold^{1,2} and Antje Boetius^{1,2}

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Introduction

Microbial communities play an essential role in carbon and nutrient cycling not only at the seafloor but also in the sea ice and in the water column, contributing significantly to Arctic ecosystem functioning. Arctic sea-ice extent declined to a record minimum in summer 2012 and the observed rapid melting resulted in the sinking and widespread deposition of fresh ice algal aggregates of the centric diatom Melosira arctica to the deep-sea floor at 4400 m water depth (Boetius et al., 2013). Sediments with algal deposits showed elevated rates of oxygen consumption, indicating remineralization by bacteria, and evidencing a response of the entire ecosystem down to the deep sea to elevated carbon flux rates. Warming and its associated physical changes in the Arctic will also affect bacterial communities, but to understand the ecosystem consequences we lack baseline information on bacterial community composition and functions in different Arctic environments.

Figure 1: Cruise track of IceArc2012 (ARK-XXVII/3) from August to October, 2012. Red dots indicate the nine sampling sites, the star indicates station 7, for which Illumina sequences were obtained.





Conclusions

• Central Arctic sea ice, melt ponds, seawater, sediment and algal aggregates host distinct bacterial communities



• For melt ponds, community description differed from previous reports, potentially indicating community shifts

- High contribution of surface-derived bacterial cells to community composition in aggregate deposits in the deep sea, indicating a transport of cells from the surface to the deep sea by the rapidly deposited aggregates
- Increase in the export of sub-ice algae is expected, therefore the role of algal aggregates as transporters of bacteria to the deep sea may be underestimated

1. Bacterial community structure



Bacterial community structure clearly differs between sea ice, water column and deep sea environments.

2. Bacterial community overlap



85% of OTUs unique to individual environments, only 0.2% of OTUs ubiquitously shared

Aggregate deposits in the deep sea

shared relatively high percentages of

OTUs with communities from **melt**

ponds, sea ice, as well as with deep-

sea sediment



Similar patterns of environment-specific community structures conserved across different study sites

Figure 1: NMDS plot of bacterial community fingerprints. Each point represents the bacterial community of one sample. The distance between points reflects the relative differences in community structure between samples. Points within each group are connected to their group centroid with a spider diagram. The light pink ellipses show 95% dispersion of each group.

Figure 2: Percentage of OTUs shared between the investigated environments as detected by Illumina sequencing. Percentages are based on a Jaccard-distance-matrix of normalized data.

3. Bacterial community composition



Of 175 classified genera, only 3 were common to all environments (Rubritalea, Colwellia, *Pseudomonas*); **46 genera** were **unique** to one environment

Predominance of *Verrucomicrobia* in melt pond water is in **contrast** to previous reports. This could be a first indication for ongoing shifts in community structure due to environmental changes

Communities in aggregate deposits in the deep sea contained several surface-derived genera also observed in the ice environment (e.g. Octadecabacter, Glaciecola), indicating a transport of cells to the **deep sea**

Figure 3: Bacterial community composition based on relative sequence abundances of the five most abundant classes detected over all environments.

Materials & Methods



69 samples from nine **Central Arctic**

Molecular fingerprinting, subset of 9 i.e. Automated Ribosomal samples Intergenic Spacer Analysis (ARISA)

In-depth insights into community composition and identification

Illumina next generation sequencing of the V4-V6 region of the 16S rRNA gene



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stations

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