Biogeosciences Discuss., 9, 17475–17517, 2012 www.biogeosciences-discuss.net/9/17475/2012/ doi:10.5194/bgd-9-17475-2012 © Author(s) 2012. CC Attribution 3.0 License.



This discussion paper is/has been under review for the journal Biogeosciences (BG). Please refer to the corresponding final paper in BG if available.

Density and distribution of megafauna at the Håkon Mosby Mud Volcano (the Barents Sea) based on image analysis

E. Rybakova (Goroslavskaya)¹, S. Galkin¹, M. Bergmann², T. Soltwedel², and A. Gebruk¹

¹P. P. Shirshov Institute of Oceanology, Russian Academy of Sciences, Nakhimovsky Pr., 36, Moscow 117997, Russia ²Alfred Wegener Institute for Polar and Marine Research, Am Handelshafen 12, 27570, Bremerhaven, Germany

Received: 22 November 2012 – Accepted: 26 November 2012 – Published: 10 December 2012

Correspondence to: E. Rybakova (Goroslavskaya) (gorolen@mail.ru)

Published by Copernicus Publications on behalf of the European Geosciences Union.

iscussion Pa	BC 9, 17475–1	BGD 9, 17475–17517, 2012												
per Discussion	Megafau Håkon Mo Volc E. Ryb (Goroslavs	Megafauna at the Håkon Mosby Mud Volcano E. Rybakova (Goroslavskaya) et al.												
Paper	Title	Title Page												
—	Abstract	Introduction												
Disci	Conclusions	References												
ussion	Tables	Figures												
Pap	14	►I												
Φŗ	•	•												
	Back	Close												
iscussio	Full Scre	en / Esc												
on P	Printer-frier	dly Version												
aper	Interactive	Discussion												



Abstract

During a survey of the Håkon Mosby Mud Volcano (HMMV), located on the Bear Island Fan in the southwest Barents Sea at \sim 1250 m water depth, different habitats inside the volcano caldera and outside it were photographed using a towed camera platform, an

- Ocean Floor Observation System (OFOS). Three transects were performed across the caldera and one outside, in the background area, each transect was ~ 2 km in length. We compared the density, taxa richness and diversity of non-symbiotrophic megafauna in areas inside the volcano caldera with different bacterial mat and pogonophoran tube-worm cover. Significant variations in megafaunal composition, density and distribution
- ¹⁰ were found between considered areas. Total megafaunal density was highest in areas of dense pogonophoran populations (mean 52.9 ind. m⁻²) followed by areas of plain light-coloured sediment that were devoid of bacterial mats and tubeworms (mean 37.7 ind. m⁻²). The lowest densities were recorded in areas of dense bacterial mats (mean ≤ 1.4 ind. m⁻²). Five taxa contributed to most of the observed variation: the ophi-
- ¹⁵ uroid Ophiocten gracilis, lysianassid amphipods, the pycnogonid Nymphon macronix, the caprellid Metacaprella horrida and the fish Lycodes squamiventer. In agreement with previous studies, three zones within the HMMV caldera were distinguished, based on different habitats and megafaunal composition: "bacterial mats", "pogonophoran fields" and "plain light-coloured sediments". The zones were arranged almost concen-
- trically around the central part of the caldera that was devoid of visible megafauna. The total number of taxa showed little variation inside (24 spp.) and outside the caldera (26 spp.). The density, diversity and composition of megafauna varied substantially between plain light-coloured sediment areas inside the caldera and the HMMV background. Megafaunal density was lower in the background (mean 25.3 ind. m⁻²) compared to areas of plain light coloured sediments inside the caldera.
- ²⁵ pared to areas of plain light-coloured sediments inside the caldera.





1 Introduction

Mud volcanoes are formations created by geo-excreted fluids. To date, more than 1700 mud volcanoes have been identified, of which some 800 are located offshore (Dimitrov, 2002). It has been estimated that more than 10 000 mud volcanoes exist on continental shelves and abyssal plains (Ivanov et al., 1996; Milkov, 2000; Pimenov et al., 2000; Fleysher et al., 2001; Dimitrov, 2002; Kopf, 2002; Milkov et al., 2003). The processes leading to their formation include mud diaperism, tectonic compression, dehydration of clay minerals at depth or a rapid deposition of mass flows (Kohl and Roberts, 1994; Kopf, 2002; Huguen et al., 2004). Mud volcanoes are often associated with petroleum deposits, tectonic subduction zones and orogenic belts. They may form domes up to several kilometres in diameter and a height of several tens of meters above the adjacent seafloor (Vogt et al., 1999). Manifestations of mud volcanism are characterised by active seepages of fluids, gas plumes, enhanced heat flow, deposits

¹⁵ Sauter et al., 2006). They are considered a significant source contributing to the marine methane budget (Jerosch et al., 2007) and, as such, of greenhouse emissions to the hydrosphere and atmosphere (Milkov et al., 2003).

of gas hydrates and unique chemoautotrophic communities (Damm and Budeus, 2003;

The Håkon Mosby Mud Volcano (HMMV) was discovered in 1989 during a regional side-scan sonar mapping survey (Crane and Solheim, 1995; Vogt et al., 1997; Milkov et al., 1999). To date, the HMMV is the only active mud volcano known from high northern latitude beyond the polar circle. Centred at ca. 72°00.25′ N/14°43.50′ E, it is situated on the Norwegian continental slope in the SW Barents Sea (Fig. 1) at ca. 1250 m water depth with a maximum elevation of 10 m and covers an area of ca. 2 km² (Krupskaya et al., 2001).

A massive 6-km thick sediment pile, that started to accumulate 30–40 million years ago after the continental break-up into nowadays Greenland and Eurasia (Vogt et al., 1997; Hjelstuen et al., 1999), lies underneath the volcano. The formation of the volcano might have coincided with a submarine landslide during the late Pleistocene,





330 000–200 000 yr ago (Vogt et al., 1999). Today, fluids rise from a sediment depth of 2–3 km through a conduit below the HMMV (Vogt et al., 1999). The age of the carbonate structures inside the modern caldera has been estimated as 11 400 yr BP (Lein and Ivanov, 2009).

⁵ During the R/V *Håkon Mosby* cruise in 1995, the mud volcano was mapped by 3.5kHz profiling, sediment coring and heat-flow measurements (Vogt et al., 1997). A gravity core from the flank of the structure contained gas and gas hydrate. High heat flow values (337 m W m⁻²) were recorded. Together with observations of microbial methane (CH₄), hydrogen sulphide gas and CH₄ gas hydrate, it was inferred that the structure was an active gas seep on the seafloor (Vogt et al., 1997).

Subsequent investigations using deep-towed side-scan sonar, sediment coring, CTD and heat flow measurements, 8.8-kHz profiling and seafloor photography during the expedition of the R/V *Professor Logachev* in 1996, confirmed that the HMMV oozed mud and seeped gas. The heat flow in the central structure was very high, exceed-¹⁵ ing 1000 m W m⁻² (Eldholm et al., 1999). The warm mud volcano was coated with white substances and the occurrence of gas hydrates on the seafloor surface was suggested (Lein et al., 1998, 1999). This expedition brought the first evidence of the presence of high benthic microbial activity and symbiotrophic fauna (two species of pogonophorans) associated with the mud volcano caldera (Pimenov et al., 1999; 20 Smirnov, 2000).

The HMMV site was revisited in 1998 during the 40th cruise of the R/V Akademik *Mstislav Keldysh* with a joint team of scientists from Norway, Germany, the USA and Russia. Two *Mir* submersibles were used for six dives (Bogdanov et al., 1999). Detailed information about the geological features of mud volcano, gas hydrate distribution, bio-

geochemical processes in the sediment, water and biota were obtained. Biogeochemical processes related to the carbon and sulphur cycles at the HMMV, microbiological processes and the structure of bacterial mats were described (Lein et al., 2000; Pimenov et al., 2000). The general features of benthic communities were reported, based on camera observations and samples taken by submersibles (Gebruk et al.,





2003). It was shown that the megabenthic community is dominated by two species of symbiotrophic siboglinid tubeworms (pogonophorans): *Sclerolinum contortum*, later transferred to the genus *Archeolinum* (Smirnov, 2008), whose biomasses reached 435 g m⁻² (Gebruk et al., 2003) and *Oligobrachia haakonmosbiensis* (Smirnov, 2000), later re-identified as *O. webbi* (Meunier et al., 2010) reached 350 g m⁻² (Gebruk et al., 2003). Associations of benthic fauna with each pogonophoran species were described. The background community outside the caldera was dominated by the ophiuroids *Ophiocten gracilis* and *Ophiopleura borealis*. Pycnogonids (*Colossendeis proboscidea*), buccinid gastropods and asteroids were also present. The trophic relation-

During the expedition ARK XIX/3b of the German research icebreaker R/V *Po-larstern*, with the French remotely operated vehicle (ROV) *Victor 6000*, in 2003, the micro-topography of HMMV was determined, video observations of the seafloor were made and the sediment and water column was sampled (Klages et al., 2004). The ROV

- ¹⁵ data was used to estimate the area covered by bacterial mats and pogonophoran fields and their relationship to mud flow. High-resolution bathymetric data, still photographs and video mosaics were integrated in a Geographic Information System (GIS). This allowed a detailed analysis of spatial patterns and quantification of biogeochemical habitats by geostatistical methods (Jerosch et al., 2007; Decker and Olu, 2010; Port-
- nova et al., 2010). High concentrations of hydrogen sulphide and methane in the sediment at HMMV and other offshore mud volcanoes offer a specific habitat for microbiological and macrofaunal associations (Sibuet and Olu, 1998; Pimenov et al., 1999; Hinrichs and Boetius, 2002; Soltwedel et al., 2005; De Beer et al., 2006). Sulphide in these systems is often a product of anaerobic methane oxidation by a microbial
 consortium of sulphate-reducing Bacteria and methane oxidizing Archaea (Boetius et al., 2000). Anaerobic oxidation of methane in the sediment and aerobic oxidation in
- bacterial mats and reduces methane flux into the water column (Boetius et al., 2002; Hinrichs and Boetius, 2002; De Beer et al., 2006; Sauter et al., 2006). However, recent research suggests that high flow velocities of mud volcano water in the seafloor reduce





the efficiency of the natural gas filter by 40 % and that the majority of methane is consumed in the siboglinid tubeworm habitat and not in the centre (Niemann et al., 2006). Furthermore, a new group of methane-consuming Archaea that live in symbiosis with Bacteria and use sulphate to oxidize methane was discovered in sediment layers below the sulphur Bacteria (Niemann et al., 2006).

In previous studies three main habitat types have been identified in the HMMV: bacterial mats, pogonophoran fields and plain light-coloured sediments (Gebruk et al., 2003; Jerosch et al., 2007). Preliminary data were given on macro- and megafauna associated with these habitats (Gebruk et al., 2003) and on meiofauna (Soltwedel et al., 2005; Portnova et al., 2010). Based on a photographic survey of the HMMV with a

- ¹⁰ al., 2005; Portnova et al., 2010). Based on a photographic survey of the HMMV with a towed camera system, we compare composition, density and diversity of megafauna in habitats inside the volcano caldera with different density of seafloor coverage by bacterial mats and pogonophoran tubeworms, and in habitats with different combinations of bacterial mats and pogonophorans, to identify an effect of a habitat type on the
- structure of benthic communities. Also we compare composition, diversity and density of megafauna inside and outside the volcano caldera, to determine an effect of the mud-volcano environment on benthic communities and to identify possible specialists of the HMMV.

2 Material and methods

20 2.1 Photographic survey

5

25

A photographic survey was carried out during the expedition ARK-XVIII/1b of the research icebreaker R/V *Polarstern* in August 2002 to the HMMV. Different habitats inside the volcano caldera and outside were photographed using a towed camera system (Ocean Floor Observation System, OFOS). Four OFOS transects were carried out in total: three across the caldera (I–III) and one outside the caldera (IV) to the south of volcano (Fig. 2). The length of each transect was ca. 2 km (Table 1).





The OFOS was equipped with a still camera (Benthos Inc.) and a black and white video camera (Deep-Sea Power & Light), two high-intensity discharge (HID) lights of 250 W each, strobes (600 W s^{-1}) and three laser pointers. A triangle laser scale with 50 cm between lasers was used to determine the camera's footprint. The still camera was mounted to the frame in a vertical position to the seafloor. It was triggered auto-5 matically every 30 s. Additional manually triggered images were taken when features of particular interest occurred in the field of view of the video camera, which was aligned with the still camera to help the winch operator adapting to bottom topography (for more details see Soltwedel et al., 2009). The OFOS was towed at approximately 0.5 knots and a target altitude of 1.5 m. From earlier OFOS deployments, this altitude has proven 10 to be the optimal distance to the seafloor to achieve the best illumination and resolution of the images. The area of seafloor on the images ranged from 1 to 8 m² owing to ship's movement, in most cases it was 3-4 m². Start and end positions of OFOS transects (from GPS fixes) and water depths along transects (from echo soundings) were obtained from the ship's data acquisition and management system.

2.2 Image analysis and identification of megafauna

In total 1604 images were examined using a stereo microscope. Of these, 1045 images, taken in the soft-sediment environment and of satisfactory quality, were used for statistical analyses. Images with hard substrata (boulders, carbonate rocks and crust) were excluded. Among the analysed images, 894 were taken along the three tran-20 sects across the caldera and 151 images along the transect outside the volcano. All visible non-symbiotrophic megafauna were counted and identified to the lowest possible taxonomic level. Taxonomic identifications were made with the assistance of experts (see acknowledgements). We used the category "morphospecies" for megafauna identified only to high taxonomic level. Recurring biological objects that could not 25 be assigned to any taxa were indicated as "morphotypes". It is possible that some morphotypes (and morphospecies) include several species. Overall, 31 categories (29 species/morphospecies and 2 morphotypes of megafauna) were identified and



quantified and used in multivariate statistical analyses and estimations of megafaunal density (Table 2). Density was not determined for the two pogonophorans, Oligobrachia webbi and Archeolinum contortum. For these and for bacterial mats we estimated the degree of seafloor area coverage using the laser scale. Sediment colour was described

- ⁵ and used for habitat description. Identification and ground-truthing was aided by previously collected material and samples taken outside the caldera using an Agassiz trawl and inside the caldera using a box corer, ROV, trap lander and baited traps. For details on gears and deployments see Table 1 in the Supplement and Pimenov et al. (1999), Bergmann et al. (2009) and Hildebrandt et al. (2011). For a complete list of species
- found inside and outside the HMMV caldera see Table 2 in the Supplement. 10

Data analysis 2.3

15

The number of organisms on each image was converted to individuals per m² (densitv). Non-metric multi-dimensional scaling (MDS) plots were generated for each transect inside the caldera based on the Bray-Curtis similarity coefficient using the PRIMER v6 programme and v-transformed species density matrices. v-transformation of densities was used to buffer the influence of most abundant taxa.

Analysis of similarity (ANOSIM) was used to determine the degree of separation of images from areas with different coverage by bacterial mats or pogonophorans. Contribution of species to similarity of different groups of images was calculated using the

SIMPER subroutine of PRIMER v6 (Clarke and Gorley, 2006). ANOSIM and SIMPER 20 were used for comparing images taken on the plain light-coloured sediment inside the caldera and images outside the caldera.

Combined data from three OFOS transects over the caldera were used for the calculation of the total megafaunal density (±standard deviation) and the mean taxa densities (±standard deviation) in areas with different coverage by bacterial mats 25 or pogonophorans. The relative contributions of the most abundant taxa to the total density in different areas were calculated. Additionally the mean taxa densities were





estimated for areas with different combinations of bacterial mats and pogonophorans based on the OFOS transect III.

The following diversity indices were applied to combined data of the three OFOS transects to describe megafaunal assemblages: Pielou's evenness (J'), Shannon– ⁵ Wiener diversity (H') (log₂) and Simpson's diversity (1 – λ) in areas with different coverage by bacterial mats or pogonophorans. Similar calculations were performed for transect IV taken outside the caldera. Taxa richness in different areas was compared based on equal number of randomly selected images.

- We compared total megafaunal densities, mean taxa densities and diversity on images with different coverage by bacterial mats or pogonophorans overall using a Kruskal-Wallis test (Minitab 16 software). Pairwise comparisons of different areas were performed using the Mann-Whitney U-Test and Bonferroni corrections of p = 0.05/n, where *n* is number of comparisons (comparison of bacterial mat sites: n = 10, p = 0.005; comparison of pogonophoran sites: n = 6, p = 0.00833; comparison of area of plain light-coloured sediments and pogonophoran sites: n = 10, p = 0.005) (Minitab 16). Comparisons of diversities were done for the equal number of randomly selected images from each area because diversity indices are based on the taxa richness, which is affected by the number of images analysed. Similarly, images from outside the caldera were compared with images from the light-coloured sediments using
- ²⁰ the Mann-Whitney U-Test.

3 Results

25

Twenty-four species/morphospecies were recorded from three OFOS transects taken across the HMMV caldera (transects I, II and III). Thirteen taxa were identified to species or genus, others were assigned to higher taxa (Table 2). The following higher taxa were present: Cnidaria (2 species), Nemertea (1), Polychaeta (1), tubeworms Siboglinidae (2), Gastropoda (1), Bivalvia (1), Pycnogonida (2), Crustacea (4), Ophi-uroidea (2), Crinoidea (1), Asteroidea (3), Holothuroidea (1), Echinoidea (1), and at





least two species of fish: *Lycodes squamiventer* and *Amblyraja hyperborea*. Samples taken for ground-truthing revealed 16 additional species (Table 2 in the Supplement). Echinoderms accounted for the majority of taxa (8 species/morphospecies). Variations in the mean density of selected taxa, area coverage by bacterial mats and pogonophorans and the sediment colour are shown on Fig. 8.

3.1 Megafauna in areas with different seafloor coverage by bacterial mats and pogonophorans

3.1.1 Bacterial mat sites

5

We used the following five categories for the seafloor area coverage by bacterial mats: 0%, 0–10%, 10–50%, 50–80% and 80–100%. ANOSIM revealed significant differences in megafaunal composition and density of images with no bacterial mats at all (0%) from images with bacterial mats present and did not reveal significant differences in megafaunal composition and density between images with different coverage of bacterial mats (Global R = 0.538, p = 0.001).

Figure 3a indicates two groups of images similar to groups revealed by ANOSIM. Some images with a coverage of 0–10 % by bacterial mats fell out of revealed groupings.

SIMPER indicated that five species contributed most to the separation of the mat coverage groups: *Ophiocten gracilis*, lysianassid amphipods, *Nymphon macronix, Metacaprella horrida* and *Lycodes squamiventer* (Fig. 4). The mean densities of these species in areas of different coverage by bacterial mats are presented in Table 4. The most notable differences in density were found in *O. gracilis* and *L. squamiventer. Ophiocten* densities decreased significantly from 37.9 ± 38.7 ind. m⁻² in the absence of mats to 1.3 ± 5.3 ind. m⁻² when bacterial mats (<10% coverage) appeared (M-W: W = 271840.0, p < 0.00001). By contrast, the density of *L. squamiventer* increased from 0.1 ± 0.3 to 1.2 ± 0.6 ind. m⁻² when bacterial mats were present (coverage > 80%). Fish densities on bacterial mats were significantly different between





bacterial mats > 80 % and all other bacterial mat categories (M-W test, p < 0.00001) with highest densities recorded in areas of 80 % bacterial coverage and lowest in areas 10 % coverage.

Total megafaunal densities sharply decreased from 43.7 ± 44.3 ind. m⁻² to 4.7 ± 8.0 ind. m⁻² when the coverage by bacterial mats increased from 0% to 10%. The Kruskal-Wallis test indicated significant differences in total megafaunal densities in areas of different coverage by bacterial mats (K-W: H = 417.90, p < 0.0001, DF = 4). The pairwise Mann-Whitney U-Test revealed significant differences in the total density of megafauna between images from areas with the following bacterial mat covrevealed significant differences between images from areas with bacterial mat coverage > 10% (p > 0.005).

Taxa richness gradually decreased with increasing coverage by bacterial mats: from 24 taxa to 3 (Table 3). Evenness and diversity were significantly higher in areas without ¹⁵ bacterial mats (J' = 0.3; H' = 0.4 and $1 - \lambda = 0.2$) and with bacterial mat coverage of up to 10 % (J' = 0.4; H' = 0.4 and $1 - \lambda = 0.2$) compared to areas with bacterial mat coverage exceeding 10 % ($J' \le 0.2$; $H' \le 0.2$ and $1 - \lambda \le 0.1$) (M-W: J': W = 8116.0-8563.5, $p \le 0.0006$; H': W = 8281-8734.5, p < 0.0001; $1 - \lambda : W = 8261.5-8672.0$, p < 0.0001). Evenness and diversity did not differ significantly in areas with bacterial mat < 10 % and in areas without bacterial mats (M-W: J': W = 29850.0, p = 0.8592; H' : W = 29057.5, p = 0.3034; $1 - \lambda : W = 29277.5$, p = 0.4279) (Table 3).

3.1.2 Pogonophoran sites

For the seafloor area coverage by pogonophoran tubeworms, we used the following four categories: 0%, 0–10%, 10–50% and 50–100%. ANOSIM indicated ²⁵ significant differences in megafaunal composition and density of images without pogonophorans (0%) from images with pogonophorans present (>0%) and of images with pogonophoran coverage < 10% from those with coverage 10–50%. Significant





differences were not observed between the categories 0-10% and 50-100%, 10-50% and 50-100% (Global R = 0.136, p = 0.001).

Figure 3b did not indicate any groupings of images based on a difference in the coverage by pogonophorans.

SIMPER revealed that five species contributed most to the separation of group-5 ings of images with different coverage by pogonophorans: O. gracilis, lysianassid amphipods, N. macronix, M. horrida and L. squamiventer (Fig. 4). The mean densities of these species in areas of different pogonophoran coverage are presented in Table 4. Most notable are differences shown by O. gracilis, lysianassid amphipods and *M. horrida*. Densities of these species increased with increasing coverage by 10 pogonophorans from <10% to >50%: O. gracilis densities from 11.9 ± 14.1 ind. m⁻² with <10% pogonophoran cover to 41.4 ± 62.2 ind. m⁻² with >50% pogonophoran A similar comparison with lysianassid amphipods showed an increase from 2.5 ± 3.5 to 8.9 ± 11.2 ind m⁻² under the same conditions and an increase in *M. horrida* from 0.1 ± 0.5 to 0.9 ± 1.9 ind. m⁻². Most pronounced were the differences in the 15 lysianassid amphipods and *M. horrida* (M-W: *W* = 25731.5, *p* < 0.00001; W=8977.0, p=0.0004 respectively. Significant differences in Ophiocten densities were found only between areas with pogonophoran coverage of 10-50 % and >50 %: 6.5 ± 19.9 and 41.4 \pm 62.2 ind. m⁻² respectively (M-W: W = 26805.0, p < 0.001). Significant differences in Ophiocten densities were not found between areas with pogonophoran 20 coverage < 10 % and > 50 %: 11.9 ± 14.1 and 41.4 ± 62.2 ind. m⁻² respectively (M-W: W = 9625.0, p = 0.0134).

Total megafaunal densities increased from 16.7 ± 15.8 ind. m⁻² to 14.6 ± 28.7 ind. m⁻² with increasing pogonophoran coverage from <10% to 10– ²⁵ 50% and total megafaunal densities increased to 52.9 ± 71.5 ind. m⁻² with increasing pogonophoran coverage from 10–50% to >50%. There were significant differences in total megafaunal densities between areas with different coverage by pogonophorans (K-W: H = 20.81, p < 0.0001, DF = 3). Pairwise comparisons revealed significant





differences in the total megafaunal densities only between images with 10–50 % and >50 % of coverage by tubeworms (M-W: W = 27782.5, p < 0.0014).

There were no significant differences in the taxa richness of different areas when equal numbers of images were considered. Taxa richness was 17–18 (Table 3). ⁵ Evenness and diversity were not significantly higher in areas with coverage by pogonophorans of < 10 % (J' = 0.4; H' = 0.5 and $1 - \lambda = 0.3$) and >50 % (J' = 0.5; H' =0.6 and $1 - \lambda = 0.3$) (M-W: J': W = 7841.5, p = 0.0184; H': W = 7808.5, p = 0.0243; $1 - \lambda$: W = 7911.5, p = 0.0099) (Table 3).

3.1.3 Area without bacterial mats and pogonophorans (plain light-coloured sediments)

10

ANOSIM revealed significant differences in the species composition of images taken from light-coloured sediments and pogonophoran settlements (Global R = 0.315, p = 0.001). SIMPER indicated two species contributing most to the separation of images from these areas: *O. gracilis* and lysianassid amphipods. The mean ¹⁵ *Ophiocten* densities were significantly lower on the plain light-coloured sediment $(35.7 \pm 21.7 \text{ ind. m}^{-2})$ than in areas with a coverage by pogonophorans exceeding 50 % (M-W: W = 108228.5, p < 0.00001), but higher than in areas with a coverage of 0– 50 % (M-W: W = 92895.0-100705.5, p < 0.00001). The mean lysianassid densities were significantly lower on light-coloured sediments than in areas with pogonophorans ²⁰ (M-W: p < 0.00001) (Table 4).

Mean megafaunal densities were significantly lower on light-coloured sediments $(37.7 \pm 22.3 \text{ ind. m}^{-2})$ compared with areas exceeding 50 % pogonophoran coverage (M-W: W = 107801.5, p = 0.0008), but significantly higher than in areas of 0–50 % pogonophoran coverage (M-W: W = 100773.0-93744.0, p < 0.00001).

Taxa richness was slightly higher on light-coloured sediments than in areas with pogonophorans (20 compared to 17–18). Evenness and diversity of light-coloured sediments were significantly lower (J' = 0.2; H' = 0.2 and $1 - \lambda = 0.1$) than those inhabited by pogonophorans (M-W: p < 0.00001) (Table 3).





3.1.4 Combination of areas with bacterial mats and pogonophorans

The mean density of selected taxa in areas with different combination of bacterial mats and tubeworms was evaluated based on the OFOS transect III. Three variations of seafloor coverage were considered both for bacterial mats and tubeworms: 0%, 0– 50%, 50–100%.

Relatively high densities of megafauna (16-98 ind. m⁻²) were observed in areas without bacterial mats. Highest densities were recorded in areas devoid of bacterial mats but with dense settlements of tubeworms (50–100 %). Areas devoid of bacterial mats and pogonophorans were characterised by high ophiuroid densities (33.1 ind. m⁻²) but relatively low densities of other taxa (0.8 ind. m⁻²). Areas devoid of bacterial mats but with dense coverage by tubeworms (>50 %) haboured the highest ophiuroid densities (77.7 ind. m⁻²) and other taxa (21.2 ind. m⁻²) (Fig. 5).

Relative contributions of the most abundant taxa to the total density in areas with different coverage by bacterial mats and tubeworms are shown on Fig. 6. Fish accounted

¹⁵ for 86% of the total density $(0.8 \pm 0.6 \text{ ind. m}^{-2})$ in areas with extensive bacterial mats (>50%). By contrast, the relative contribution of fish was 12% $(0.2 \pm 0.4 \text{ ind. m}^{-2})$ in areas of extensive pogonophoran settlements (>50%) and 6% $(0.1 \pm 0.3 \text{ ind. m}^{-2})$ in areas without bacterial mats and tubeworms. The relative contribution of ophiuroids increased from 0.8% $(0.1 \pm 0.4 \text{ ind. m}^{-2})$ in areas with extensive bacterial mats to ²⁰ 39% $(40.2 \pm 62.3 \text{ ind. m}^{-2})$ in areas of extensive pogonophoran fields and up to 87% $(34.4 \pm 21.6 \text{ ind. m}^{-2})$ in areas without bacterial mats and tubeworms (on light-coloured sediments). The highest relative contribution of lysianassids (26%) and caprellids (0.9– 2%) was found in the pogonophoran area.

3.2 Comparison of megafauna from three zones inside the volcano caldera

Inside the HMMV caldera the following three main habitats can be distinguished: bacterial mats, pogonophoran settlements and light-coloured sediments (Fig. 7). The habitats are arranged in zones around the volcano centre in a more or less concentric





17489

particular habitat type.

densities varied from 0 to 115.6 ind. m⁻² between individual images (mean 35.7 ± 21.7 ind. m⁻²). Some of the megafauna (mainly relatively large mobile crustaceans and rays) were present throughout the caldera with no clear association to any 25

individual images (mean 24.9 ± 48.1 ind. m⁻²). 15 Zone III The zone of plain light-coloured sediments devoid of bacterial mats and

density of 0.6 ind. m⁻² in this zone.

5

10

20

contortum and Oligobrachia webbi is located further away from the centre of the caldera. Most of the megafauna is absent from this zone, whereas the density of pycnogonids, N. macronix, caprellids, M. horrida, lysianassid amphipods and bivalve molluscs clearly increased on siboglinid fields. Total megafaunal densities in this zone ranged from 0.1 to 271.2 ind. m⁻² between individual images (mean 34.0 ± 56.0 ind. m⁻²). Ophiuroid densities ranged from 0 to 248 ind. m⁻² between

mats and pogonophoran colonies may co-occur in different proportions (Fig. 8). Zone I A zone of bacterial mats is most clearly distinguished around the caldera centre. The zone is characterised by extensive bacterial mats on grey or yellowish sediment. Megafaunal densities ranged from 0.1 to 52.7 ind. m⁻² between individ-

pattern, however, boundaries between zones are not always clear because bacterial

Discussion Paper Megafauna at the ual images (mean 3.3 ± 6.4 ind. m⁻²). The fish Lycodes squamiventer reached a Håkon Mosby Mud Volcano Discussion Paper Zone II The zone of settlements of siboglinid tubeworms (pogonophorans) Archeolinum E. Rybakova (Goroslavskava) et al. Title Page Abstract Introduction **Discussion** Paper Conclusions References **Tables Figures** pogonophorans forms the periphery of the caldera. This zone was dominated by the ophiuroid O. gracilis. Most of the megafauna taxa, first of all the echinoderms, **|**◀ were abundant in this zone. Total megafaunal densities ranged from 0.1 to 116.3 ind. m^{-2} between individual images (mean 37.7 ± 22.3 ind. m^{-2}). Ophiuroid Back Close **Discussion** Paper Full Screen / Esc **Printer-friendly Version** Interactive Discussion



9, 17475-17517, 2012

The highest density of non-symbiotrophic megafauna was recorded in the zone of light-coloured sediments, followed by the zone of pogonophoran settlements. The lowest densities were recorded from the bacterial mat zone (Fig. 9).

3.3 Comparison of the volcano caldera and the background area

OFOS IV transect outside the caldera revealed twenty-four The species/morphospecies and two morphotypes of megafauna. Eleven taxa were identified either to species or genus level, others were assigned to higher taxa: Porifera (2), Cnidaria (1), Nemertea (2), Polychaeta (2), Gastropoda (1), Bivalvia (1), Pycnogonida (2), Crustacea (4), Ophiuroidea (2), Crinoidea (1), Asteroidea (4), Echinoidea (1) and at least one fish species. Additionally two morphotypes (undeter-10 mined sedentary animals) were recorded. Echinoderms (8 species/morphospecies) and sedentary animals (5 species/morphospecies/morphotypes) accounted for the majority of taxa (Table 2). Ten additional species were found in trawl samples (Table 2 in the Supplement). However, this number cannot be considered representative since the trawl net was damaged by several large rocks during trawling. Mean 15 megafaunal densities outside the caldera were 25.3 ± 21.0 ind m⁻² ranging from 1.9 to 84.0 ind. m^{-2} .

Six taxa accounted for more than 95% of the total density: *O. gracilis*, sabellid polychaetes, the sponge *Stylocordyla borealis*, lysianassid amphipods, unidentified sedentary animals (morphotype 2) and bivelves. *Ophiocten gracilis* was the most abundant

tary animals (morphotype 2) and bivalves. Ophiocten gracilis was the most abundant species (65.7%).

ANOSIM revealed significant differences in megafaunal composition and density on images taken inside the caldera on light-coloured sediments devoid of bacterial mats and pogonophorans and images outside the caldera (Global R = 0.378, p = 0.001) in-

²⁵ dicating significant differences. SIMPER showed that the following three species contributed most to this separation of images: *O. gracilis*, sabellid polychaetes and *S. borealis*.





The composition of the dominant taxa in the background area differed from that inside the caldera. Only two taxa, *O. gracilis* and lysianassid amphipods, were abundant in both areas (Table 7). In the background area, other abundant taxa included the sabellids, *S. borealis* and unidentified sedentary organisms absent or found in low numbers

⁵ inside the caldera. The mean density of *O. gracilis* was significantly lower in the background area $(22.3 \pm 21.8 \text{ ind. m}^2)$ compared to the zone of light-coloured sediments inside the caldera $(35.7 \pm 21.7 \text{ ind. m}^2)$ (M-W: W = 29635.0, p = 0.00001). Similarly the mean density of lysianassid amphipods was significantly lower in the background $(0.3 \pm 0.7 \text{ ind. m}^2)$ compared to the zone of light-coloured sediments inside the caldera $(0.5 \pm 1.6 \text{ ind. m}^2)$ (M-W: W = 91731.5, p = 0.0236) (Table 4).

Mean total megafaunal density was significantly lower in the background $(25.3 \pm 21.0 \text{ ind. m}^{-2})$ compared to the area of light-coloured sediments inside the caldera $(37.7 \pm 22.3 \text{ ind. m}^{-2})$ (M-W: W = 104087.0, $\rho < 0.00001$).

- In total, 24 taxa were found inside the volcano caldera and 26 outside on the OFOS ¹⁵ images. In both areas 19 taxa were present (Table 2). However, sedentary biota, such as sponges, cnidarians and some unidentifiable morphotypes were only recorded in the background area. Based on the equal number of randomly selected images, taxa richness was higher outside the caldera (26) than in the area of light-coloured sediments inside the caldera (21).
- Evenness and diversity were significantly higher in the background than on the lightcoloured sediments (J' = 0.4 vs. J' = 0.2; H' = 0.6 vs. H' = 0.2 and $1 - \lambda = 0.3$ vs. $1 - \lambda = 0.1$) (M-W: W = 29113.0, p = 0.00001; W = 29899.0, p = 0.00001; W = 29660.5, p = 0.00001 respectively) (Table 3).

4 Discussion

The Håkon Mosby Mud Volcano has received considerable attention over the last decade as one of the target areas of ambitious EU-funded programmes such as HER-MES and HERMIONE and various nationally funded projects. However, studies on





17492

Two species of siboglinid tubeworms (perviate pogonophorans), *Archeolinum contortum* and *Oligobrachia webbi*, dominated inside the caldera. Both genera are common at depths of the Norwegian-Greenland Basin (NGB) and are usually associated with reducing biotopes. They were found at methane seeps at 1230 m depth on the Vestnesa

macro- and megafaunal communities inhabiting the HMMV are still very limited and our knowledge of the volcano background community remains poor. We were able to identify 40 taxa in total (including species, "morphospecies" and

"morphotypes") based on images and samples taken inside the volcano caldera. Ge-

- ⁵ bruk et al. (2003) reported 46 benthic taxa from this area. A large portion of the taxa listed by these authors included small and infaunal forms, such as polychaetes and amphipods that are not easily detected on seafloor photographs and were missing in our records. Based on the combined data of Gebruk et al. (2003) and our present study, the species inventory of the HMMV comprises at least 71 taxa. In the present study we
 added at least 25 species/morphospecies to the inventory: Cnidaria (4), Nemertea (1).
- Annelida (4), Sipunculida (1), Cephalorhyncha (1), Bivalvia (2), Crustacea (3), Echinodermata (6), Pisces (3) (Table 2 in the Supplement).

Taxa richness did not differ noticeably inside and outside the caldera: 40 vs. 36, although the number of species outside the caldera could be higher if three transects

- (as inside the caldera) had been taken or the sampling effort had been more intense. A photographic survey at the HAUSGARTEN observatory west of Svalbard (~ 1300 m depth) revealed lower taxa richness (12 taxa/morphotypes) (Soltwedel et al., 2009). Thus, Vinogradova et al. (1996) reported 200 benthic taxa based on 22 trawls at the Russian nuclear submarine *Komsomolets* wreck site at 1700 m depth in the Bear Is-
- ²⁰ land area, approximately 120 miles north of the HMMV on the continental slope. The benthic community at this site (based on the biomass) was termed the "*Bathybiaster-Ophiocten*-Amphipoda-Oweniidae" assemblage. Similar to the HMMV, benthos at the *Komsomolets* site was dominated by echinoderms (primarily asteroids and ophiuroids), amphipods and polychaetes. In our study, in trawl catches outside the caldera ophi-²⁵ uroids and asteroids were also dominant.





Ridge (Gebruk et al., 2003) and on the Norwegian slope at the Nyegga site at ~ 720 m depth (Krylova et al., 2011). *Archeolinum contortum* was also abundant at the recently discovered deep-sea hydrothermal vent site "Loki's Castle" at a depth of 2350 m at the junction of the Mohn and Knipovich Ridges (Pedersen et al., 2010). The species

- O. webbi was reported from the shallower hot vent site on the Mohn Ridge north of Jan Mayen (500–750 m water depth; Schander et al., 2010). Although specialists of reducing habitats, both siboglinid species seem to be opportunistic: they occur in methane seep and hydrothermal vent habitats, on sunken wood or other decomposing organic material (Gebruk et al., 2003).
- ¹⁰ In the HMMV area both pogonophorans are restricted to the volcano and the nearest surrounding (Gebruk et al., 2003). Populations of the more abundant *A. contortum* cover thousands of square meters of the seabed. The geo-referenced video mosaics and detailed spatial analyses by Jerosch et al. (2007) illustrate particularly dense populations of these tubeworms in the so-called "hummocky periphery" of the caldera and
- estimated the total area covered by tubeworms at 275 958 m², by dense bacterial mats (density > 50 %) at 31 435 m² and the central caldera part at 115 165 m². Our data indicate that the highest densities of *A. contortum* are in areas with bacterial mat coverages of less than 50 % of the sea floor. Our results based on the analysis of 894 images from HMMV disagree with conclusion of Niemann et al. (2006) that sulphide-oxidizing mats and tubeworm colonies are mutually exclusive. A great number of images showed both
- biota.

According to our data, the density and proportions of taxonomic groups noticeably varied between habitats inside and outside the caldera. The megafaunal species observed along OFOS transects could be grouped according to their occurrence in differ-

²⁵ ent habitats. Many species abundant outside the caldera and at the inactive periphery of the volcano were either completely absent or very rare in the pogonophoran and bacterial mat zones. Among such species were echinoderms (echinoids, starfish, crinoids and holothurians), polychaetes, sponges and nemertea. An exception was the ophiuroid *O. gracilis,* which was quite abundant in the dense settlements of pogonophorans





(>50%) without bacterial mats. The mean density of *O. gracilis* in this habitat was higher than in the background: 63.6 ± 67.8 ind. m⁻² and 22.3 ± 21.8 ind. m⁻², respectively. A photographic survey at the HAUSGARTEN observatory west of Svalbard (~ 1300 m depth), conducted also on the ARK XIX/3 cruise, revealed slightly lower densities of *O. gracilis* (16.8 ± 3.7 ind. m⁻²) and an aggregated occurrence (Soltwedel et al., 2009). *Ophiocten gracilis* occurs in a wide range of habitats; this species appears to be very adaptive and opportunistic. In hydrothermal vent habitats on the Mid-Atlantic Ridge ophiuroids (*Ophiocten acies*) reached densities as high as 80 ind. m⁻² (Gebruk et al., 2000) in the Logatchev area.

Some megafaunal species at HMMV were clearly associated with pogonophorans. Among them were the caprellid amphipod *Metacaprella horrida*, the pycnogonid *Nymphon macronix* and lysianassid amphipods. The density of burrowing bivalve molluscs also increased on the pogonophoran fields. Total megafaunal densities of pogonophoran fields would increase dramatically if the tubeworms themselves were added to our calculations. However, on the seafloor images it is almost impossible to determine even approximate numbers of tubes, especially when they are curled as in *A. contortum*. Also many tubes in dense populations are empty (A. Gebruk, personal

observation, 1998).
Overall, total megafaunal densities clearly increased on the pogonophoran fields
(>50% coverage) compared to the background habitat (25.3±21.0 ind. m⁻²) and dropped to minimum values on the bacterial mats (>50%). The limited number of megafauna species able to live in a "moderate" reducing environment are apparently capable of taking advantage of low competition in partitioning of resources and reach high population densities. Models based on temperature profiles indicate lower

fluid flow rates (0.4 m yr⁻¹) in the pogonophoran fields compared to those of the centre or bacterial mat zone (Kaul et al., 2006). The increased habitat complexity provided by pogonophorans together with a more stable environment may result in higher megafaunal densities. Furthermore, Niemann et al. (2006) concluded that microbe—invertebrate symbioses have an advantage over free-living microbial populations as





they can engineer their environment to increase access to both electron donors and acceptors by special migratory behaviour, mining and pumping.

The bacterial mat zone was characterized by the lowest megafaunal density and species richness. The opposite trend was shown for the density of meiofauna at the

- 5 HMMV it was the highest in the bacterial mat zone compared to pogonophoran fields and the background (Portnova et al., 2010). In our study, megafaunal density decreased sharply even at low bacterial coverage (<10%). The only species with higher densities on bacterial mats was the fish *Lycodes squamiventer*. This species had significantly higher growth rates at the HMMV compared with conspecifics from HAUS-OADTEN equals to compare the density of the second states of the second states.
- ¹⁰ GARTEN caught at comparable depth (Hildebrandt et al., 2011) indicating favourable local growth conditions at the mud volcano. Furthermore, stable isotope analyses showed that the δ^{13} C and δ^{15} N in *L. squamiventer* muscle tissues from the HMMV were significantly lower than those from HAUSGARTEN indicating that the fish from HMMV relied on food linked to the methane-based food web (Gebruk et al., 2003;
- Bergmann et al., 2009, and unpublished data). Zoarcidae is a fish family often encountered at hydrothermal vents and cold seeps, with some specialised species that are likely to have adapted to the toxic environment (Desbruyères et al., 2006).

The density of megafauna in our study was relatively high. For example, on the mud volcanoes in the eastern Mediterranean the density of non-symbiotrophic megafauna were two or three orders of magnitude lower (Olu-Le Roy et al., 2004). However the

were two or three orders of magnitude lower (Olu-Le Roy et al., 2004). How depth of these volcanoes is greater, approximately from 1700 to 2000 m.

The three habitats (zones) distinguished at the HMMV are arranged in a more or less concentric pattern: the central zone, around it pogonophoran fields and the lightsediment zone in the periphery. The latter two zones harbour a complicated mosaic

of bacterial mats. Such a concentric structure of a habitat and community pattern is more typical of hydrothermal vents, centred around hot fluid emissions (Galkin, 2002). For example, Hessler and Kaharl (1995) distinguished "the vent opening, the nearfield and the periphery". This obvious simplification points at pronounced gradients of environmental parameters that are very characteristic of hydrothermal vents and





other habitats with focused emission of heat and reduced compounds, one of such habitats being mud volcanoes. At typical cold methane seep sites the pattern of fauna distribution is usually more heterogeneous and is first of all controlled by fluid flow (Sahling et al., 2002; Levin, 2005).

- ⁵ The central zone of the volcano appears to be devoid of megafauna most likely because of the instability of the environment. Indeed, Feseker et al. (2008) reported an exceptionally high level of mud volcanic activity with temperature gradients of more than 40 °C m⁻¹ within the upper half meter of the sediments close to the geometric centre of the caldera, frequent mud eruptions of small volume and limited local extent.
- The variety of reducing habitats known at present in the Norwegian-Greenland Basin (NGB) includes two "cold" methane seep sites – Nyegga at 710–730 m depth on the Norwegian slope in the Storrega slide area (Ivanov et al., 2007; Ivanov et al., 2010) and the methane seep area at 1230 m depth on the Vestnesa Ridge (Gebruk et al., 2003), which includes the relatively hot and active mud volcano (HMMV), a shallow-water hy-
- ¹⁵ drothermal vent north of Iceland, Kolbenisey (~90 m depth) (Fricke et al., 1989), two upper bathyal hydrothermal vents, Grimsey (400 m) north of Iceland (Hannington et al., 2001) (no biological investigations yet) and the area north of Jan Mayen on the Mohn Ridge with the fields Trollveggen (500–560 m), Galionella Garden (~615 m), Soria Moria I and II (700–750 m) (Schander et al., 2010), and one recently discovered
- hydrothermal vent site, Loki's Castle, at 2350 m depth, at the junction of the Mohn and Knipovich Ridges (Pedersen et al., 2010). According to published data, most of the macro- and megafauna of methane seeps and hydrothermal vents in the NGB, including the HMMV, is not specialised to these habitats, it comprises primarily regular background fauna (with exception of two species of pogonophoran tubeworms,
- opportunists of reducing environments). This feature is unusual: in other areas, reducing habitats are dominated by highly specialized megafaunal species restricted in their distribution to a specific reducing habitat or rarely to several of them. In the NGB the only species clearly obligate to methane seeps is the bivalve *Isorropodon nyeggaensis* (Krylova et al., 2011), representative of the family Vesicomyidae, one of the iconic





taxa of reducing environments (Levin, 2005; Desbruyères et al., 2006). The lack of vent/seep specialists in the NGB may result from a young age of fauna: the recent deep-sea benthic fauna north from the Faroe-Iceland Ridge shows signs of origin after the end of the last glacial period (Golikov, 1985; Svararsson et al., 1993).

⁵ Supplementary material related to this article is available online at: http://www.biogeosciences-discuss.net/9/17475/2012/ bgd-9-17475-2012-supplement.pdf.

Acknowledgements. We are grateful to the Captain, the officers and the crew of R/V Polarstern for their support during the ARK-XVIII/1b, ARK-XX/1, ARK-XXII/1b expeditions. We also thank
 and the teams of the remotely operated vehicles "Victor 6000" and "Quest 4000" for their support in obtaining samples for ground-truthing. This work was partially supported by HERMES – Hotspot Ecosystem Research on the Margins of European Seas (contract number GOCE-CT-2005-511234), funded by the European Commission's Framework Six Programme and HERMIONE – Hotspot Ecosystem Research and Man's Impact on European Seas (contract number 226354) funded by the European Commission's Framework Seven Programme. Financial support also was provided by the Grant 12-05-31066 of RFBR (Russian Federation) to ER and by the Grant 8664 of Minnauki (Russian Federation) to AG. Our special thanks to experts

- for their help in taxonomic identifications: K. Tabachnik and O. Tendal (Porifera), N. Sanamyan (Actiniaria and Ascidiacea), E. Rodriguez (Actiniaria), N. Budaeva and S. Gromisz (Polychaeta),
 M. Kedra (Sipuncula), E. Krylova (Bivalvia), A. Warén (Mollusca), A. Raiskiy and R. Bramber (Pycnogonida), G. Vinogradov and J.-M. Weslawski (Amphipoda), A. Brandt (Isopoda),
 J. Guerrero (Tanaidacea), A. Mironov (Echinoidea), D. Piepenburg (Ophiuroidea, Asteroidea),
 A. Rogacheva (Holothuroidea) and P. R. Möller (Fish). The authors thank Paul Dando for the correction of the style of English and useful comments on the manuscript. We are indebted to
- anonymous reviewers for improvements to an earlier draft of the manuscript. This is publication
 no. 31571 of the Alfred Wegener Institute for Polar and Marine Research.





References

20

- Bergmann, M., Dannheim, J., Bauerfeind, E., and Klages, M.: Trophic relationships along a bathymetric gradient at the deep-sea observatory HAUSGARTEN, Deep-Sea Res. Pt. I, 56, 408–424, 2009.
- ⁵ Boetius, A., Ravenshlag, K., Schubert, C. J., Rickert, D., Widdel, F., Gieseke, A., Amann, R., Jorgensen, B. B., Witte, U., and Pfannkuche, O.: A marine microbial consortium apparently mediating anaerobic oxidation of methane, Nature, 407, 623–626, 2000.
 - Boetius, A., Jorgensen, B. B., Amann, R., Henriet, J. P., Hinrichs, K. U., Lochte, K., MacGregor, B. J., and Voodrouw, G.: Microbial systems in sedimentary environments of continental mar-
- gin, in: Ocean Margin Systems, edited by: Wefer, G., Billett, D., Hebbeln, D., Joergensen, B.
 B., Schlüter, M., and van Weering, T. C. E., Springer Verlag, Berlin-Heidelberg, Germany, 479–495, 2002.
 - Bogdanov, Y. A., Sagalevich, A. M., Vogt, P. R., Mienert, Y., Sundvor, E., Crane, K., Lein, A. Y., Egorov, A. V., Peresypkin, V. I., Cherkashev, G. A., Gebruk, A. V., Ginsburg, G. D., and
- Voitov, D. V.: The Håkon Mosby Mud Volkano in the Norwegian Sea: results of comprehensive investigations with submersibles, Okeanologia, 39, 412–419, 1999 (in Russian).
 - Clarke, K. R. and Gorley, R. N.: PRIMER v6: user manual/tutorial, PRIMER-E Ltd, Plymouth, 2006.
 - Crane, K. and Solheim, A. (Eds.): Sea floor atlas of the northern Norwegian-Greenland Sea, 137, Norsk Polarinstitutt Meddelelser, Oslo, Norway, 172 pp., 1995.
- Damm, E. and Budeus, G.: Fate of vent-derived methane in seawater above the Håkon Mosby Mud Volcano (Norwegian Sea), Mar. Chem., 82, 1–11, 2003.
 - De Beer, D., Sauter, E., Niemann, H., Witte, U., Schlüter, M., and Boetius, A.: In situ fluxes and zonation of microbial activity in surface sediments of the Håkon Mosby Mud Volcano, Limnol.
- ²⁵ Oceanogr., 51, 1315–1331, 2006.
- Decker, C. and Olu, K.: Does macrofaunal nutrition vary among habitats at the Hakon Mosby mud volcano?, Cah. Biol. Mar., 51, 361–367, 2010.
 - Desbruyères, D., Segonzac, M., and Bright, M. (Eds.): Handbook of Deep-sea Hydrothermal Vent Fauna, Second completely revised edition, Denisia, 18, 544 pp., 2006.
- ³⁰ Dimitrov, L. I: Mud volcanoes the most important pathway for degassing deeply buried sediments, Earth-Sci. Rev., 59, 49–76, 2002.



- Discussion Paper Eldholm, O., Sundvor, E., Vogt, P. R., Hielstuen, B. O., Crane, K., Nilsen, A. K., and Gladczenko, T. P.: SW Barents Sea continental margin heat flow and Håkon Mosby Mud Volcano, Geo-Feseker, T., Foucher, J.-P., and Harmegnies, F.: Fluid flow or mud eruptions? Sediment temperature distributions on Håkon Mosby mud volcano, SW Barents Sea slope, Mar. Geol., 247, Fleysher, P., Orsi, T. H., Richardson, M. D., and Anderson, A. L.: Distribution of free gas in marine sediments: a global overview, Geo-Mar. Lett., 21, 103-122, 2001. Fricke, H., Giere, O., Stetter, K., Alfredsson, G. A., Kristjansson, J. K., Stoffers, P., and Svavars-
- son, J.: Hydrothermal vent communities at the shallow subpolar mid-Atlantic ridge, Mar. Biol., 10 102. 425-429. 1989.

Mar. Lett., 19, 29–37, 1999.

194-207, 2008.

5

- Galkin, S. V.: Hydrothermal vent communities of the World Ocean. Structure, typology, biogeography, GEOS, Moscow, Russia, 200 pp., 2002.
- Gebruk, A. V., Chevaldonne, P., Shank, T., Lutz, R. A., and Vrijenhoek, R. C.: Deep-sea hydrothermal vent communities of the Logatchev area (14°45' N, Mid-Atlantic Ridge): diverse 15 biotopes and high biomass, J. Mar. Biol. Assoc. UK, 80, 383-394, 2000.
 - Gebruk, A. V., Krylova, E. M., Lein, A. Y., Vinogradov, G. M., Anderson, E., Pimenov, N. V., Cherkashev, G. A., and Crane, C.: Methane seep community of the Håkon Mosby Mud Volcano (the Norwegian Sea): composition and trophic aspects, Sarsia, 88, 394-403, 2003.
- Golikov, A. N.: Stages and frontiers of the Cenozoic. Analysis of geological time and evolu-20 tion of marine gastropod fauna, Proceedings of the Zoological Institute, USSR Academy of Sciences, 130, 31–52, 1985 (in Russian).
 - Hannington, M., Herzig, P., Stoffers, P., Scholten, J., Botz, R., Garbe-Schönberg, D., Jonasson, I. R., Roest, W., and Shipboard Scientific Party: First observations of high-temperature sub-
- marine vents and massive anhydrite deposits off the north coast of Iceland, Mar. Geol., 177, 25 199-220, 2001.
 - Hessler, R. R. and Kaharl, V. A.: The deep-sea hydrothermal vent community: an overview, in: seafloor hydrothermal systems: physical, chemical, biological, and geochemical interactions, edited by: Humphris, S. E., Zierenberg, R. A., Mullineaux, L. S., and Thomson, E., American Geophysical Union, Washington, 72-84, 1995.
- 30 Hildebrandt, N., Bergmann, M., and Knust, R.: Longevity and growth efficiency of two deepdwelling Arctic zoarcids and comparison with eight other zoarcid species from different climatic regions, Polar Biol., 34, 1523-1533, 2011.





- Hinrichs, K. U. and Boetius, A.: The anaerobic oxidation of the methane: new insights in microbial ecology and biogeochemistry, in: Ocean Margin Systems, edited by: Wefer, G., Billet, D., Hebbeln, D., Jorgensen, B. B., Schlüter, M., and Weering, T. C. E. V., Springer Verlag, Berlin-Heidelberg, Germany, 457–477, 2002.
- ⁵ Hjelstuen, B. O., Eldholm, O., Faleide, J. I., and Vogt, P. R.: Regional setting of Håkon Mosby Mud Volcano, SW Barents Sea margin, Geo-Mar. Lett., 19, 22–28, 1999.
 - Huguen, C., Mascle, J., Chaumillon, E., Kopf, A. J., Woodside, J., and Zitter, T.: Structural setting and tectonic control of mud volcanoes from the Central Mediterranean Ridge (Eastern Mediterranean), Mar. Geol., 209, 245–263, 2004.
- ¹⁰ Ivanov, M. K., Limonov, A. F., and Van Weering, T.: Comparative characteristics of the Black Sea and Mediterranean Ridge mud volcanoes, Mar. Geol., 132, 253–271, 1996.
 - Ivanov, M., Blinova, V., Kozlova, E., Westbrook, G. K., Mazzini, A., Minshull, T. A., and Nouzé, H.: First sampling of gas hydrate from the Vøring Plateau, Eos T. Am. Geophys. Un., 88, 209–216, 2007.
- Ivanov, M., Mazzini, A., Blinova, V., Kozlova, E., Laberg, J.-S., Matveeva, T., Taviani, M., and Kaskov, N. Seep mounds on the Southern Vøring Plateau (offshore Norway), Mar. Petrol. Geol., 27, 235–1261, 2010.
 - Jerosch, K., Schlüter, M., Foucher, J.-P., Allais, A.-G., Klages, M., and Edy, C.: Spatial distribution of mud flows, chemoautotrophic communities and biogeochemical habitats at Håkon Mosby Mud Volcano, Mar. Geol., 243, 1–17, 2007.

20

Kaul, N., Foucher, J. P., and Heesemann, M.: Estimating mud expulsion rates from temperature measurements on Håkon Mosby Mud Volcano, SW Barents Sea, Mar. Geol., 229, 1–14, 2006.

Klages, M., Thiede, J., and Foucher, J.-P.: The expedition ARKTISXIX/3 of the research vessel

- ²⁵ "Polarstern" in 2003, Reports on Polar and Marine Research, Alfred Wegener Institute for Polar and Marine Research, Bremerhaven, Germany, 488, 346 pp., 2004.
 - Kohl, B. and Roberts, H. H.: Fossil foraminifera from four active mud volcanoes in the Gulf of Mexico, Geo-Mar. Lett., 14, 126–134, 1994.
- Kopf, A. J.: Significance of mud volcanism, Rev. Geophys., 40, 1005, doi:10.1029/2000RG000093, 2002.
 - Krupskaya, V. V., Andreeva, I. A., Sergeeva, E. I., Cherkashev, G. A., and Ivanov M. K.: The Håkon Mosby Mud Volcano (Norwegian Sea): features of a structure and deposit composition, in: Experience of systemic oceanological exploration in Arctic regions, edited





by: Lisitsin, A. P. and Vinogradov, M. E., Nauchnyi Mir, Moscow, Russia, 492-502, 2001 (in Russian).

- Krylova, E. M., Gebruk, A. V., Portnova, D. A., Todt, C., and Haflidason, H.: New species of the genus Isorropodon (Bivalvia: Vesicomyidae: Pliocardiinae) from cold methane seeps at
- Nyegga (Norwegian Sea, Voring Plateau, Storrega slide), J. Mar. Biol. Assoc. UK, 91, 1135-5 1144, 2011.

Lein, A. Y. and Ivanov, M. V. (Eds.): Biogeochemical cycle of methane in the ocean, Nauka, Moscow, Russia, 576 pp., 2009.

Lein, A. Y., Vogt, P. R., Crane, K., Egorov, A., Pimenov, N. V., Savvichev, A. S., Ginsburg, G.

- D., Ivanov, G. I., Cherkashev, G. A., and Ivanov, M. V.: Geochemical pecularities of gas-10 bearing (CH_4) deposits of the submarine mud volcano in the Norwegian Sea, Geohymiya, 3, 230-249, 1998 (in Russian).
 - Lein, A. Y., Vogt, P. R., Crane, K., Egorov, A., and Ivanov, M.: Chemical and isotopic evidence for the nature of the fluid in CH₄ - containing sediments of the Håkon Mosby Mud Volcano,
- Geo-Mar. Lett., 19, 76-83, 1999. 15

25

30

- Lein, A. Y., Pimenov, N. V., Savvichev, A. S., Pavlova, G. A., Vogt, P. R., Bogdanov, Y. A., Sagalevich, A. M., and Ivanov, M. V.: Methane as a source of organic matter and the hydrocarbon acid of carbonates at a cold seep in the Norwegian Sea, Geohymiya, 3, 268-281, 2000 (in Russian).
- Levin, L. A.: Ecology of cold-seep sediments: interactions of fauna with flow, chemistry and 20 microbes, Oceanogr. Mar. Biol., 43, 1-46, 2005.
 - Meunier, C., Andersen, A. C., Bruneaux, M., Le Guen, D., Terrier, P., Leize-Wagner, E., and Zal, F.: Structural characterization of hemoglobins from Monilifera and Frenulata tubeworms (Siboglinids): First discovery of giant hexagonal-bilayer hemoglobin in the former "Pogonophora" group, Comp. Biochem. Phys. A., 155, 41-48, 2010.
 - Milkov, A. V.: Worldwide distribution of submarine mud volcanoes and associated gas hydrates, Mar. Geol., 167, 29-42, 2000.

Milkov, A. V., Vogt, P. R., and Cherkashev, G. A.: Sea-floor terrains of Håkon Mosby Mud Volcano as surveyed by deep-tow video and still photography, Geo-Mar. Lett., 19, 143-149, 1999.

Milkov, A. V., Sassen, R., Apanasovich, T. V., and Dadashev, F. G.: Global gas flux from mud volcanoes: a significant source of fossil methane in the atmosphere and the ocean, Geophys. Res. Lett., 30, 1037, doi:10.1029/2002GL016358, 2003.





- Niemann, H., Lösekann, T., De Beer, D., Elvert, M., Nadalig, T., Knittel, K., Amann, R., Sauter, E. J., Schlüter, M., Klages, M., Foucher, J. P., and Boetius, A.: Novel microbial communities of the Håkon Mosby mud volcano and their role as a methane sink, Nature, 443, 854–858, 2006.
- ⁵ Olu-Le Roy, K., Sibuet, M., Fiala-Médioni, A., Gofas, S., Salas, C., Mariotti, A., Foucher, J. P., and Woodside, J.: Cold seep communities in the deep eastern Mediterranean Sea: composition, symbiosis and spatial distribution on mud volcanoes, Deep-Sea Res. Pt. I, 51, 1915– 1936, 2004.

Pedersen, R. B., Rapp, H. T., Thorseth, I. H., Lilley, M. D., Barriga, F. J. A. S., Baumberger,

- ¹⁰ T., Flesland, K., Fonseca, R., Früh-Green, G. L., and Jorgensen, S. L.: Discovery of a black smoker vent field and vent fauna at the Arctic Mid-Ocean Ridge, Nat. Commun., 1, 1–6, 2010.
 - Pimenov, N. V., Savvichev, A. S., Rusanov, I., Lein, A. Y., Gebruk, A. V., and Moskalev, L. I.: Microbial processes of carbon cycles at the base of the food chain of Håkon Mosby Mud Volcano benthic community. Geo-Mar. Lett., 19, 89–96, 1999.
- Pimenov, N. V., Savvichev, A. S., Rusanov, I., Lein, A. Y., and Ivanov, M. V.: Microbiological processes of carbon and sulfur cycles at cold methane seeps in the North Atlantic, Microbiologiya, 69, 831–843, 2000 (in Russian).

15

20

Portnova, D., Mokievsky, V., and Soltwedel, T.: Nematode species distribution patterns at the Håkon Mosby Mud Volcano (Norwegian Sea), Mar. Ecol.-Evol. Persp., 31, 24–41, 2011.

Sahling, H., Rickert, D., Lee, R. V., Linke, P., and Suess, E.: Macrophaunal community structure and sulphide flux at has hydrate deposits from the Cascadia convergent margin, NE Pacific, Mar. Ecol.-Prog. Ser., 231, 121–138, 2002.

Sauter, E. J., Muyakshin, S. I., Charlou, J.-L., Shlüter, M., Boetius, A., Jerosch, K., Damm, E.,

- Foucher, J.-P., and Klages, M.: Methane discharge from a submarine mud volcano into the upper water column by gas hydrate-coated methane bubbles, Earth Planet. Sc. Lett., 243, 354–365, 2006.
 - Schander, C., Rapp, H. T., Bakken, T., Berge, J., Cochrane, S., Kongsrud, J. A., Oug, E., Byrkjedal, I., Cedhagen, T., Fosshagen, A., Gebruk, A., Larsen, K., Obst, M., Pleijel, F., Stöhr,
- ³⁰ S., Todt, C., Warén, A., Hadler-Jacobsen, S., Keuning, R., Mikkelsen, N. T., Petersen, K. H., Torseth, I., and Pedersen, R. B.: The fauna of hydrothermal vents on the Mohn Ridge (North Atlantic), Mar. Biol. Res., 6, 155–171, 2010.





- Sibuet, M. and Olu, K.: Biogeography, biodiversity and fluid dependence of deep-sea cold-seep communities at active and passive margins, Deep-Sea Res. Pt. II, 45, 517–567, 1998.
- Smirnov, R. V.: Two new species of pogonophoran from the Arctic mud volcano off northwestern Norway, Sarsia, 85, 141–150, 2000.
- Smirnov, R. V.: Morphological characters and classification of the subclass Monilifera (Pogonophora), the problem of evolution of bridle in pogonophora, Russ. J. Mar. Biol., 34, 404–413, 2008 (in Russian).
 - Soltwedel, T., Portnova, D., Kolar, I., Mokievsky, V., and Schewe, I.: The small-sized benthic biota of the Håkon Mosby Mud Volcano (SW Barents Sea slope), J. Marine Syst., 55, 271–290, 2005.
- 10 290, 2005
 - Soltwedel, T., Jaeckisch, N., Ritter, N., Hasemann, C., Bergmann, M., and Klages, M.: Bathymetric patterns of megafaunal assemblages from the arctic deep-sea observatory HAUS-GARTEN, Deep-Sea Res. Pt. I, 56, 1856–1872, 2009.
 - Svararsson, J., Strömberg, J.-O., and Brattegard, T.: The deep-sea asselote (Isopoda, Crus-
- tacea) fauna of the northern seas: species composition, distributional patterns and origin, J. Biogeogr., 20, 537–555, 1993.
 - Vinogradova, N. G., Turpaeva, E. P., Moskalev, L. I., Galkin, S. V., and Bagirov, N. E.: Species composition of the bottom fauna in the area of study, leading and common forms and modifications of communities, in: Oceanographic research and underwater technical operations at
- the site of the nuclear submarine Komsomolets wreck, edited by: Vinogradov, M. E., Sagalevitch, A. M., and Khetagurov, S. V., Nauka, Moscow, Russia, 202–206, 1996 (in Russian).
 Vogt, P. R., Cherkashev, G. A., Ginsburg, G. D., Ivanov, G., Milkov, A. V., Crane, K., Lein, A. Y., Sundvor, E., Pimenov, N., and Egorov, A.: Håkon Mosby Mud Volcano provides unusual example of venting, EOS T. Am. Geophys. Un., 78, 556–557, 1997.
- Vogt, P. R., Gardner, J., and Crane, K.: The Norwegian-Barents-Svalbard (NBS) continental margin: introducing a natural laboratory of mass wasting, hydrates and ascent of sediment pore water and methane, Geo-Mar. Lett., 19, 2–21, 1999.

	BC	D													
	9, 17475–1	7517, 2012													
-	Megafau Håkon Mo Volc	na at the osby Mud ano													
	E. Rybakova (Goroslavskaya) et al.														
)															
	Title Page														
-	Abstract	Introduction													
2	Conclusions	References													
	Tables	Figures													
J	14	►I.													
	•	•													
	Back	Close													
	Full Scre	en / Esc													
J	Printer-frien	dly Version													
	Interactive	Discussion													

Discussion Pape

Jiscussion Pape

Discussion Paper

Discussion Paper



	B 9, 17475–	GD 17517, 2012														
	Megafa Håkon M Vol E. Ry	Megafauna at the Håkon Mosby Mud Volcano E. Rybakova (Goroslavskava) et al.														
	(Goroslav	skaya) et al.														
	Abstract	Introduction														
	Conclusions	References														
0000	Tables	Figures														
	I	►I.														
-	•	•														
2	Back	Close														
0100	Full Sc	reen / Esc														
	Printer-frie	endly Version														
DODr	Interactive	e Discussion														



Table 1. Details of OFOS transects. Transects I, II and III – inside the caldera, transects IVa and b – reference site.

Gear	Transect/ Deployment	Date, August 2002		Posi North-	ition, East (°)		Survey time (min)	Water (r	depth n)	Total No. of images	No. of images analysed	Total area covered by images
			Start		End			Start	End			(m ²)
OFOS	1	21	72.01 14.44		71.60	14.44	105	1261	1268	401	256	1003
OFOS	II	21	72.00	14.41	72.00 14.45		130	1284	1254	391	233	1042
OFOS	111	22/23	72.00	14.42	71.60	14.46	94 1278 1254		1254	562	405	1480
OFOS	IVa	21	71.56	71.56 14.39		14.39	31	1329	1308	36	24	93
	IVb	21	71.57	71.57 14.40		14.39	160	1325	1313	196	127	389

 Table 2. Megafauna and other categories recorded on OFOS transects.

High taxon/object	Species/morphospecies/ morphotype/category	HMMV	Background
Porifera	cf. <i>Stylocordyla borealis</i> Grey form (?Demospongia)		++++
Cnidaria			
Anthozoa	Purple form (cf. Edwardsiida)	+	
	Small pinkish (cf. Hormathiidae)	+	
	Undetermined		+
Nemertea	cf. Nipponnemertes pulchra		+
	nemertini-like worm	+	+
Annelida			
Polychaeta	Oligobrachia webbi	+	
	Archeolinum contortum	+	
	cf. Polynoidae		+
	cf. Sabellidae (tubes)	+	+
Mollusca			
Gastropoda	cf. Buccinidae	+	+
Bivalvia	Whitish burrowing form	+	+
Arthropoda			
Pycnogonida	cf. Nymphon macronix	+	+
	cf. Collossendeis proboscidea	+	+
Crustacea,			
Amphipoda	cf. Metacaprella horrida	+	+
	cf. Lysianassidae	+	+
?lsopoda	Small whitish form	+	+
Decapoda	Bentho-pelagic redish form	+	+

Diechleeion Da	BC 9, 17475–1	GD 7517, 2012													
ner Discussion	Megafauna at the Håkon Mosby Mud Volcano E. Rybakova (Goroslavskaya) et al. Title Page														
Danor															
-	Abstract	Introduction													
	Conclusions	References													
	Tables	Figures													
Dan	I.	►I													
	•	•													
5	Back	Close													
	Full Scre	en / Esc													
ק ק	Printer-frier	dly Version													
aner	Interactive	Discussion													



17506

Table 2. Continued.

High taxon/object	Species/morphospecies/ morphotype/category	HMMV	Background
Echinodermata			
Ophiuroidea	Ophiocten gracilis	+	+
	Ophiopleura borealis	+	+
Crinoidea	Bathycrinus cf. carpenteri	+	+
Asteroidea	cf. Pontaster	+	+
	Whitish form (cf. Bathybiaster)	+	+
	Flat pentagonal form, short rays	+	
	Short arms (<i>Hymenaster</i> type)		+
	White short arms		+
Holothuroidea	cf. <i>Elpidia glacialis</i>	+	
Echinoidea	Undetermined	+	+
Pisces			
Rays	Amblyraja hyperborea	+	
Zoarcidae	Lycodes squamiventer	+	+
Unknown, morphotype1	Small brown bushes (?hydroids)		+
Unknown, morphotype 2	Small white sedentary forms (?anemones, sponges)		+
Bacterial mat	Area cover (%):		
	0	+	+
	<10	+	
	10–50	+	
	50–80	+	
	>80	+	
Siboglinidae	Area cover (%):		
Archeolinum contortum	0	+	+
	<10	+	
	<50	+	
	>50	+	
Sediment	Colour yellow (light)	+	+
	Colour grey (dark)	+	





	Discussion Paper Discussion	BC 9, 17475–1 Megafau Håkon Ma Volc E. Ryt (Goroslavs	GD 7517, 2012 Ina at the osby Mud cano Dakova skaya) et al.
-	Pa		
-	ber	Title	Page
-		Abstract	Introduction
-	Disc	Conclusions	References
_	ussior	Tables	Figures
-	Pape	14	►I
-	Pr	•	•
		Back	Close
	iscussi	Full Scre	een / Esc
	on Pa	Printer-frier	ndly Version
	aper	Interactive	Discussion

CC () BY

Table 3. Indices of megafaunal diversity and megafaunal density (ind. m^{-2}) in areas of different coverage by bacterial mats and pogonophorans inside the HMMV caldera (OFOS transects I, II and III combined) and the background (OFOS transect IV).

Index		Bacte	erial mats (B	BM)			Pogonopl	BM and P	Background		
	0%	<10%	<50 %	<80 %	>80 %	0 %	<10%	<50 %	>50 %	0%	
Number of images	576	173	84	49	12	531	84	101	178	367	151
Total species (S)	24	19	10	9	3	23	17	18	19	23	26
Pielou's evenness (J')	0.3	0.4	0.2	0.2	0.2	0.2	0.4	0.6	0.5	0.2	0.4
Shannon diversity (H')	0.4	0.4	0.2	0.2	0.2	0.2	0.5	0.7	0.6	0.2	0.6
Simpson diversity $(1 - \lambda)$	0.2	0.2	0.1	0.1	0.1	0.1	0.3	0.4	0.3	0.1	0.3
Total megafaunal density (±SD)	43.7 ± 44.3	4.7 ± 8.0	1.9±3.2	1.4 ± 2.0	1.3±0.7	26.4 ± 25.1	16.7±15.8	14.6 ± 28.7	52.9 ± 71.5	37.7 ± 22.3	25.3±21.0

Table 4. Mean density (ind. m ²) and standard deviations (SD) of the most abundant taxa in ar-
eas of different coverage (%) by bacterial mats and pogonophorans. Data from OFOS transects
I, II and III combined and from the background transect IV.

Taxon/	Bacterial mats (BM)								Pogonophorans (P)							BM a	nd P	Backg	round			
Morphotype	0	%	<10	%	<50	%	>50	>50 % >80 %			0	0%)%	<50)%	>50)%	c)		
	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Ophiocten gracilis	37.9	38.7	1.3	5.3	0.1	0.4	0.1	0.4	0	0	24.7	24.4	11.9	14.1	6.5	19.9	41.4	62.2	35.7	21.7	22.3	21.8
Amphipoda Lysianassidae	3.7	7.9	1.6	3.0	0.5	1.7	0.2	1.2	0	0	0.3	1.3	2.5	3.5	4.9	6.8	8.9	11.2	0.5	1.6	0.3	0.7
Metacaprella horrida	0.3	1.1	0.1	0.4	0.1	0.5	0	0	0	0	0.0	0.1	0.1	0.5	0.1	0.5	0.9	1.9	0.0	0.1	0.0	0.0
Nymphon macronix	1.2	3.5	0.9	3.1	0.4	1.6	0.2	0.6	0.1	0.3	0.7	2.3	1.4	3.3	2.3	6.0	1.0	2.9	0.9	2.7	0.0	0.1
Lycodes squamiventer	0.1	0.3	0.5	0.6	0.8	0.7	0.8	0.6	1.2	0.6	0.3	0.6	0.2	0.5	0.2	0.4	0.2	0.4	0.1	0.3	0.0	0.1
Bivalvia (whitish form)	0.1	0.3	0.1	0.3	0.0	0.1	0.0	0.1	0	0	0.1	0.2	0.2	0.3	0.3	0.5	0.1	0.3	0.1	0.2	0.1	0.2
Polychaeta tubes (cf. Sabellidae)	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1.0	1.0
Porifera cf. Stylocordyla borealis	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1.0	1.2
Morphotype 2 (white seden- tary form)	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0.1	0.3







Fig. 1. Location of the Håkon Mosby Mud Volcano in the south-west Barents Sea.







Fig. 2. Position of the OFOS transects I, II and III taken across the Håkon Mosby Mud Volcano caldera area and position of the reference OFOS transect IV taken outside the volcano caldera.















Fig. 4. Most abundant species on the OFOS images. **(a)** *Nymphon macronix* (Pycnogonida), **(b)** Lysianassidae (Amphipoda), **(c)** *Lycodes squamiventer* (Pisces), **(d)** *Ophiocten gracilis* (Ophiuroidea) **(e)** Bivalvia gen. sp (whitish form).







Fig. 5. Mean densities (ind. m^{-2}) of selected taxa in areas with combinations of different coverage by bacterial mats (BM) and pogonophorans (P) along Transect III. Relative location within the volcano caldera is shown.







Fig. 6. Relative contribution of the most abundant taxa to total density in areas with different bacterial mat (BM) and pogonophoran (P) cover. Data of three OFOS transects (I, II and III) combined.









Fig. 7. Images from different habitats. (a) bacterial mat (coverage > 80 %) with fish *Lycodes* squamiventer; (b) patches of bacterial mat (<10 %) on grey sediment, fish *Lycodes* squamiventer; (c) tubeworms Archeolinum contortum (>50 %); (d) plain light-coloured sediments with numerous ophiuroids, Ophiocten gracilis.



Fig. 8. Variations in mean densities of selected taxa, area coverage of some biological objects and sediment colour on images along transects OFOS I, II and III. Density values in ind. m^{-2} on the left axis. Area coverage of bacterial mats: (0) 0%, (1) <10%, (2) 10–50%, (3) 50–80%, (4) >80%; *Archeolinum contortum*: (0) 0%, (1) <10%, (2) 10–50%, (3) >50%. Sediment colour: (1) yellow, (2) grey. Zones are shown in colour: red – bacterial mats, yellow – siboglinid settlements, pink – plain light-coloured sediment.







Fig. 9. Mean densities (ind. m⁻²) of selected taxa in different zones of the Håkon Mosby Mud Volcano. Data of three OFOS transects (I, II and III) combined.

Printer-friendly Version

Interactive Discussion