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ORIGINAL PAPER



Community composition of epibenthic megafauna on the West Greenland Shelf

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Abstract Epibenthic organisms are a critical component of the marine environment, functioning as ecosystem engineers, habitat and food for other organisms. Our knowledge of the diversity, complexity and sensitivities of these habitats is limited, particularly at higher latitudes and greater depths. The West Coast of Greenland is the site of a commercially important shrimp trawl fishery, but there are few published records describing the benthic community structure of the region. Here we report results from benthic camera surveys conducted at 119 sites, over 3 years, spanning 1400 km of the West Greenland continental shelf (61-725 m depth). A total of 29 classes of epibenthic taxa were identified from the images. There are significant differences of composition and diversity in sites with hard and soft substrate. Hard-substrate communities are relatively diverse with higher abundances and are characterised by sessile, attached groups such as Hydrozoa, Anthozoa, Bryozoa and Porifera. Soft-sediment sites are less diverse and dominated by Polychaeta and have specialist Malacostraca such as the commercially exploited shrimp, Pandalus borealis. Distribution patterns and variation in epibenthic megafauna are related to substrate and the environmental parameters depth, temperature and current speed. This study represents the first quantitative characterisation of epibenthic megafaunal assemblages on the

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C. Yesson chris.yesson@ioz.ac.uk West Greenland continental shelf. These data constitute an important baseline, albeit in a region heavily impacted by trawl fisheries, and demonstrate the utility of benthic photography for examining and monitoring seabed diversity and change.

Keywords West Greenland \cdot Epibenthic communities \cdot Continental shelf \cdot Hard substrate \cdot Soft substrate \cdot Benthic invertebrates

Introduction

The epibenthic organisms that constitute seafloor communities are critical components of the marine ecosystem. They provide three-dimensional habitat which can be protective or supportive to other organisms. They function as substrate upon which and within which other organisms settle or live (Levin and Dayton 2009; Buhl-Mortensen et al. 2010). Furthermore, they are food for other organisms and can play important roles as ecosystem engineers, functioning to redistribute and remineralise carbon (Renaud et al. 2007).

West Greenland study area

The West Greenland coastline extends from the Arctic $(85^{\circ}N)$ to subarctic $(60^{\circ}N)$ and is traversed by numerous fjords, many of them acting as direct links between the inland ice sheet and the ocean. Moreover, many islands are scattered directly off the coast resulting in an extremely long coastline and a variety of benthic habitats. The continental shelf often extends >100 km offshore. A mix of shallow banks (<50 m) and deep troughs (>300 m) results in a highly complex bathymetry in the shelf area. In

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Southwest Greenland, two water masses dominate: (1) the cold low-saline polar/coastal water from the East Greenland Current and (2) the warmer, more saline Atlantic water entering the area via the Irminger Current (Myers et al. 2007). The continental shelf in this south-western region is relatively narrow and rocky with a steep shelf edge, and the West Greenland Current is relatively strong. Further north, the current slows, the influence of the relatively warm Atlantic water is weaker, and the area is characterised by seasonal sea ice cover and a wide and less well-defined shelf (Boertmann and Mosbech 2011).

There are few published records describing benthic community structure in Greenland, and all of them focus on coastal and shelf habitats, while the deep offshore basins are largely unstudied (Boertmann and Mosbech 2011). The available studies of benthic invertebrates in West Greenland reveal a considerable amount of local scale variation in individual abundances and taxon richness on both local and regional scales. This is caused primarily by differences in sediment composition, currents, food input and disturbance level (Boertmann and Mosbech 2011). Clear differences in growth rates (of bivalves and sea urchins) related to differences in temperature and food availability have also been measured in coastal Greenlandic waters (Blicher et al. 2007; Sejr et al. 2009).

Currently, little information is available to assess and describe benthic faunal composition and diversity on the larger geographic scale of the West Greenland Shelf area. Benthic diversity studies undertaken in Greenland have been largely confined to local scale, inshore areas (Schmid and Piepenburg 1993; Piepenburg and Schmid 1996; Sejr et al. 2000). However, one survey in the Godthaabsfjord system in Southwest Greenland (Sejr et al. 2010) used Van Veen grabs to survey macrobenthic infauna community composition and diversity. This survey stretched from the inner fjord, across the shallow banks, and down the continental slope. They found high beta diversity, but this covers one transect and may not be representative of the wider region.

A recent compilation of data from across the Arctic suggested that West Greenland has a relatively high diversity of benthic species richness (Piepenburg et al. 2011). Impact assessments undertaken in Disko Bay and Eastern Baffin Bay (Boertmann and Mosbech 2011) document high richness, diversity and abundance of infaunal organisms (>100 infauna species in grab samples of 0.1 m^2) including several species believed to be new to science. However, the variation within the West Greenland region remains poorly described and only a modest part of benthic biodiversity is quantified and described in the published literature at present (Boertmann and Mosbech 2011).

Substrate types (rock, mud and gravel, mixed substrate) in areas that are exploited by the shrimp fishery in West Greenland have been characterised and charted in the fishery management plan (Lassen et al. 2013). The distribution and abundance of corals and sponges has not been systematically studied. A review of coral and sponge by catch collected during experimental trawls between 2010 and 2012 included notably few corals collected from depths <500 m, and knowledge of shallower distributions remains sparse. Only one high diversity coral area was identified between 63° N and 64° N and at 1000–1500 m (Jørgensen et al. 2013).

The aims of this study are to identify taxa from benthic images of West Greenland and quantify taxon richness and diversity of epibenthic megafauna (>1 cm) at the taxonomic rank of class. Community composition will be examined in relation to environmental variables along a large range of the continental shelf.

Materials and methods

Image collection

Three benthic invertebrate surveys were carried out in June and July 2011, 2012 and 2013 as part of the annual stock assessment of the West Greenland shrimp fishery. Fieldwork was conducted aboard the R/V Paamiut, a 1085 tonne shrimp trawling vessel operated by the Greenland Institute of Natural Resources (GINR). Sampling took place between the hours of 6 p.m. and 6 a.m. Images of the seabed were taken with a drop camera, deployed from a winch. The camera system consisted of a Nikon digital SLR camera, DSC-10000 Digital Ocean Imaging Systems (DOIS) deep-sea camera housing and 200 W-S Remote Head Strobe flash unit (DOIS, Model3831), all mounted on a weighted steel frame. A weight suspended below the frame triggered the camera on contact with the seabed. Each image sampled an area of approximately 0.3 m².

Ten images were taken at each sampling station. The location, time, depth and length of winch wire extension were recorded for each image. In between pictures, the camera was raised 10–20 m off the seabed for 1 min to ensure subsequent pictures did not sample the same area. Typically, the ship and camera would drift 20–50 m (based on ship GPS) during the 1-min interval between pictures. Stations were selected to represent a spread of geography, depth, seabed and fishing impact within the geographic limitations imposed by daytime operations (sampling had to be relatively nearby the start/end locations of daytime operations).

Taxon identification

We identified, counted and recorded taxa from images with the aid of guides and collaborators. Colonial organisms (such as encrusting Bryozoa) were counted as 1 'individual' per continuous 'patch' or 'unit'. Images were compared with physical samples collected from grabs and bycatch. Twenty-nine taxonomic classes were observed for which identification could be regarded as reliable (Online Resource 1). The majority of fauna identified were epifauna, but some clams could be identified due to visible siphons, and others (polychaetes, holothuria) were also often identifiable. Images were processed with the aid of the software 'Poseidon', developed by computer scientists at University College London, in collaboration with the authors of this study, specifically to aid identification of benthic taxa.

Two phyla were not examined at class level due to difficulties with identifications. Porifera were subdivided into three categories based on their morphology. These were (i) encrusting sponges, those forming a continuous mat over another object such as a stone, for example, *Myxilla* spp. and *Halichondria* spp.; ii) arborescent sponges, those with a branching structure, such as *Haliclona* spp.; iii) massive sponges, large (unbranched) sponges such as *Geodia* or *Polymastia*. Bryozoa were also subdivided into three categories (i) encrusting Bryozoa (as encrusting sponges—example *Escharina* spp.); (ii) soft Bryozoa, such as *Securiflustra* spp. and *Alcyonidium* spp.; (iii) stony Bryozoa, with rigid branching (for example *Hornera* spp.) or lattice structures (such as *Reptorella* spp.).

Analysis

Although 10 images were taken at each station, some stations did not produce 10 images suitable for analysis, due to sediment disturbance in soft-sediment areas or a tilted camera in rocky areas. Five images per station were selected for analysis (representing the best balance between image quality and maximising the number of stations to analyse). Data from these images were aggregated into station-level data to be used for analysis.

Taxon richness, abundance, evenness (Pielou's measure of evenness) and Shannon–Wiener's diversity index were determined for each camera station. Abundance was defined as the total number of individuals (or distinct individual colonies). Taxa were identified as generalist, or specialist to hard or soft seabed using the 'clam' multinomial classification method (Chazdon et al. 2011). Samplebased taxon accumulation curves were used to test the degree to which all taxa were successfully observed using seabed images. The expected total number of taxa in hardand soft-substrate communities was calculated using three different extrapolation methods: bootstrap, first-order jackknife and Chao (Magurran and McGill 2010).

Prior to subsequent analysis, 'singleton' taxa that appeared as a single observation in just one site were removed from the data set. Taxon observation data were transformed using both the Wisconsin and square root transformation to reduce the influence of very abundant taxa (Legendre and Gallagher 2001).

A multidimensional scaling analysis (MDS) was performed on the community composition data (Faith et al. 1987). Environmental data were fit to the MDS ordination to test for significant association of composition and environment. Environmental data for the seabed was gathered at the location of each station using the TOPAZ4 Arctic Ocean Reanalysis oceanographic model (http://cata logue.myocean.eu.org/static/resources/myocean/pum/MYO2-ARC-PUM-002-ALL_V4.1.pdf). Mean values for salinity, temperature and current speed were calculated over the full time period of the oceanographic model (1991–2010).

Sites were clustered into groups using a Bray–Curtis distance matrix with a ward linkage method (Jørgensen et al. 2015). The top two hierarchical classifications were considered in subsequent analyses. The multi-response permutation procedure was used to test for differences between clusters. The simper method (Clarke 1993) was used to find taxa discriminating between clusters based on Bray–Curtis distances. An analysis of similarities (ANO-SIM) was used to determine significance of taxon dissimilarity by cluster (Clarke 1993). All analysis was performed using the vegan library of the R statistical software program (Oksanen et al. 2013).

Results

Fauna surveyed

A total of 119 stations were photographically sampled between 2011 and 2013. A map showing the location of sites along the West Greenland continental shelf is presented in Fig. 1. The stations span depths of 61–725 m but are concentrated on the 100–500 m depth zone around the areas of the shrimp fishery. The environmental conditions covered by these stations are broadly similar to the region covered by the 100–500 m depth zone (Fig. 2), although are over-represented in the warmer, more southerly areas. Fifty-five stations were classified as 'hard' substrata (i.e. rocky seabed), while 64 were classified as 'soft' (either sandy or muddy). A total of 29 taxonomic classes were identified from at least one station (Table 1).

A full list of taxa identified is provided in Online Resource 1. Polychaeta, Ophiuroidea and Ascidiacea are the most commonly encountered taxa, seen in the majority



Fig. 1 Location of sampling stations. Seabed images were taken on three cruises over three years between 2011 and 2013. (Map projection EPSG: 3411)

of sites and regarded as generalists for the area. There are many specialist taxa in hard-bottomed areas including sessile, attached fauna such as Porifera, Bryozoa, Hydrozoa and Anthozoa. Distribution maps for all taxa are presented in Online Resource 2. There was a large variation in abundance estimates between stations, and abundances were consistently higher in hard-bottomed communities (Table 1). Soft-bottom communities were more likely to be dominated by Polychaeta, but soft-specialist taxa include Malacostraca, observations of which are predominantly composed of the commercially caught shrimp *Pandalus borealis*. Hard- and soft-substrate communities are significantly different (ANOSIM, p < 0.001).

Community clusters

Sites were clustered based on taxonomic similarity. Example images from each cluster are presented in Online Resource 3. The primary partition of the cluster analysis (Fig. 3—Cluster 1 + 2/Cluster 3 + 4) agrees broadly with substrata (Fig. 3). The ANOSIM indicates these two clusters are significantly different (p < 0.001). Furthermore, the second tier of clustering (clusters 1, 2, 3 and 4) is also significantly different (ANOSIM test, p < 0.001). The hard-substrate clusters have the highest within-group similarities (Table 2).

Cluster similarities and their most discriminating taxa are presented in Table 2. Cluster 1 is most readily distinguished by the presence of Ascidiacea; cluster 2 is characterised by Anthozoa and encrusting Bryozoa; and cluster 3 is discriminated by the relative abundance of Gastropoda and Malacostraca; cluster 4 has the greatest abundance of Polychaeta.

Community clusters differ by environment, with the predominantly soft-bottom clusters 3 and 4 found in colder and deeper water with slower currents. Conversely, the (mostly) hard-bottom clusters 1 and 2 are found predominantly in warmer, shallower water with faster current speeds (Fig. 4). There are geographic patterns evident (Fig. 5) although there is strong geographic overlap between some clusters. Cluster 1 sites are focussed in the south (with 2 outliers), while cluster 3 sites are found predominantly in the north; clusters 2 and 4 exhibit a wider geographic range.

The MDS plot (Fig. 6) summarises taxonomic similarities between sites. The tight grouping of clusters 1 and 2 indicates that stations in these groupings have more similar community composition than those in clusters 3 and 4. Latitude, depth, current speed and temperature all have significant directional associations with community composition, although these are all strongly correlated (as they are with substrate).

Discussion

This study presents a baseline survey and first description of epibenthic megafauna composition and distribution along the West Greenland shelf. The outer shelf from the



Fig. 2 Environmental profile of sampling stations. Histograms represent the 119 stations sampled. The curves show the equivalent profile for the study area, based on 1000 random locations selected within the 100–500 m depth zone of Fig. 1

latitude of Disko bay to Upernavik is defined by groups such as Malacostraca, Gastropoda and Bivalvia. Moving south, we see a transition to rockier habitats with sessile, attached fauna more dominant (including Anthozoa and Porifera). Documenting the distributions of potentially habitat-forming taxa such as sponges and corals is an important first step to support their conservation.

The soft-substrate epibenthic megafaunal communities described here are notably less diverse than hard-substrate communities. This is consistent with the results of grab sample studies of infauna characterising soft-sediment habitats in other areas around the Northwest Atlantic (Kenchington et al. 2001, Sparks-McConkey and Watling 2001). Polychaeta are the most common taxa on both hard-

and soft-substrate areas and account for more than half of all observations. Ophiuroidea are found in 107 of 119 stations and are the next most widespread taxon. The predominance of this group was observed in earlier studies in NE Greenland, where Echinodermata were the most common component of epibenthic communities and Ophiuroidae the dominant taxa in each study area (Starmans et al. 1999). In NE Greenland sites, this dominance was much more pronounced (63 % of observations were Echinodermata). The substrata, depth range and latitude surveyed in these studies are not directly comparable. Echinoderms also dominated bycatch in Barents Sea surveys (Jørgensen et al. 2015) though photographic surveys and bycatch data are not directly comparable.

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	Hard	Soft	All
Number of stations	55	64	119
Total observations	40,714	23,205	63,919
Abundance range	46-2837	8-3900	8-3900
Total classes	28	29	29
Class richness (min-max)	10–21	2-18	2-21
Class evenness (min-max)	0.1-0.9	0.0-0.9	0.0-0.9
Shannon index (min-max)	0.2–2.4	0.1-2.5	0.1-2.5
Class pool estimates			
Chao	28 (±1)	29 (±0)	29 (±0)
Jackknife	29 (±1)	30 (±1)	30 (±1)
Bootstrap	29 (±1)	30 (±1)	29 (±1)
Most abundant classes			
Rank 1	Polychaeta	Polychaeta	Polychaeta
Rank 2	Ophiuroidea	Ophiuroidea	Ophiuroidea
Rank 3	Ascidiacea	Ascidiacea	Ascidiacea
Rank 4	Bryozoa (stony)	Maxillopoda	Bryozoa (stony)
Class specialisation	Hard specialists	Soft specialists	Generalists
	Bryozoa (stony)	Polychaeta	Ophiuroidea
	Bryozoa (encrusting)	Bryozoa (soft)	Ascidiacea
	Porifera (encrusting)	Malacostraca	Maxillopoda
	Anthozoa		Bivalvia
	Hydrozoa		Holothuroidea
	Porifera (massive)		Gastropoda
	Porifera (arborescent)		Asteroidea
	Rhynchonellata		Crinoidea
	Polyplacophora		Echinoidea
			Pycnogonida
			Scaphopoda
			Nemertea

Class pool estimates are based on taxon accumulation curves. Specialist/generalist taxa are defined by the clam test, which examines relative abundance across habitats

Measuring diversity

 Table 1
 Summary of station

 data for hard and soft areas

Although biodiversity indices are primarily designed for use in species-level analysis, higher taxon-based approaches are common for image-based marine studies given the difficulty of identifying organisms from in situ photographs (Freese et al. 1999; Collie et al. 2000; Compton et al. 2013). Some biodiversity dynamics will not have been described by this analysis, due to the aggregation of species at higher taxonomic levels. For example, examining taxonomic classes such as Anthozoa ignores the differences between species with very different ecological requirements. Anthozoa includes the soft-sediment specialist sea pens (such as *Pennatula* and *Umbellula* spp. which are common in northern Greenland) with organisms requiring rock attachment such as large gorgonians (i.e. *Paragorgia arborea* which is seen in some southern sites). The two phyla for which class level identification proved difficult (Porifera, Bryozoa), were subdivided into functional groups to limit the aggregation of broad ecological differences. The implications of aggregation at such a coarse taxonomic resolution should be kept in mind when interpreting the results presented herein. However, the use of higher taxonomic levels as biodiversity indices is suited to investigating broader-scale patterns (Robert et al. 2014) and the wide geographic spread of this analysis is concordant with examining this broad scale.

This study is limited to epifaunal diversity. In the future, this should be supplemented with the infaunal component to complete biodiversity estimates. Grab sampling of





Fig. 3 Dendrogram of hierarchical cluster analysis based on Bray– Curtis distances using the Ward method. Numbers in parentheses represent mean within-group taxonomic distances based on a

undisturbed soft sediment has recorded high infaunal diversity, with >100 different invertebrate species per m^2 and several thousand individuals per m^2 (up to >10,000 ind. m^2) in coastal and shelf areas of West Greenland (Sejr et al. 2010; Blicher et al. 2011).

Temporal variation

The variability and accuracy of some community indices have been shown to be highly dependent on the temporal variability of species abundances (Trebitz et al. 2003). This study does not attempt to account for temporal variation. Each of our stations was sampled once at a fixed point in time from June and August between 2011 and 2013. One of the most frequently observed taxa in this study is the motile Ophiuroidea, and several stations show what appears to be large feeding aggregations. Sampling dense feeding aggregations could reveal ephemeral patterns of high local diversity. However, local aggregations of Ophiuroidea are known to persist over time (at least in dynamic, shallow environments, Dauvin et al. 2013, see also Piepenburg 2000; Blicher and Sejr 2011). In this study, all ophiuroids are grouped into a single taxon and the majority of the most populous taxa are positionally fixed and likely to endure for multiple years, which reduces the potential for large temporal variation.

Another type of temporal variation results from diurnal migration. For example, the commercially fished shrimp *Pandalus borealis* (the most commonly observed decapod seen in 41 stations) exhibits a diurnal migration pattern,

multiple-response permutation procedure. Labels preceded by *asterisk* show stations classified with hard substrata

inhabiting the seabed in the day and moving up into the water column during the night (Bergström 2000). The night-time sampling of this survey will lead to under-representation of this species. It is not known whether other diurnally migrating taxa are under-represented in these data.

Environment

We find evidence of environmental influence on community composition, with factors temperature, depth, slope and current speed showing significant association with community structure. Sejr et al. (2010) found strong infauna species turnover along sediment and depth gradients in the Godthaabsfjord system, but weak correlation of diversity and environment. Jørgensen et al. (2015) found temperature, salinity and ice cover to be significant determinants of epibenthic community structure in the Barents. It should be noted that our study area follows a north–south strip of shelf, with northern regions being notably colder, deeper, muddier, less saline and with weaker currents. It is unclear to what extent the environmental results of this study are obfuscated by co-variation along a latitudinal gradient.

Fishing impacts

A direct and current influence on the diversity and functioning of the benthic systems of West Greenland is the disturbance impact of bottom trawling, primarily for

	Cluster 1	Cluster 2	Cluster 3	Cluster 4		
N	30	27	16	46		
N (hard)	27	20	0	8		
N (soft)	3	7	16	38		
Depth range	181–323	109–517	206–411	61–725		
Latitude (min)	60°13′N	60°18′N	62°33′N	60°18′N		
Latitude (max)	63°10′N	70°34′N	72°16′N	71°44′N		
Class summary						
Total classes	24	28	16	28		
Total observations	23,413	12,712	829	26,965		
Most abundant classes						
Rank 1	Polychaeta	Polychaeta	Polychaeta	Polychaeta		
Rank 2	Ascidiacea	Ophiuroidea	Malacostraca	Ophiuroidea		
Rank 3	Ophiuroidea	Bryozoa (encr.)	Ophiuroidea	Ascidiacea		
Rank 4	Bryozoa (stony)	Bryozoa (stony)	Bivalvia	Bryozoa (encr.)		
Rank 5	Porifera (encr.)	Ascidiacea	Hydrozoa	Maxillopoda		
Rank 6	Hydrozoa	Anthozoa	Bryozoa (soft)	Bryozoa (stony)		
Discriminating classes						
Cluster 1	-	Ascidiacea, Holothuroidea	Ascidiacea, Porifera (massive)	Ascidiacea, Hydrozoa		
Cluster 2	Nemertea, Pycnogonida	-	Anthozoa, Porifera (massive)	Anthozoa, Bryozoa (encr.)		
Cluster 3	Malacostraca, Gastropoda	Malacostraca, Gastropoda	_	Malacostraca, Gastropoda		
Cluster 4	Polychaeta, Asteroidea	Polychaeta, Asteroidea	Polychaeta, Asteroidea	-		
Pairwise anosim (p)	R)					
Cluster 1	-	0.432	0.889	0.201		
Cluster 2	0.001	-	0.850	0.107		
Cluster 3	0.001	0.001	-	0.183		
Cluster 4	0.001	0.013	0.003	-		

Table 2 Summary of stations by cluster group

Discriminating taxa are defined by the simper analysis, where row headings identify the groups where this taxa predominates compared with the group identified by the column heading (e.g. cluster 3 is positively associated with Gastropoda and Malacostraca in comparison with all other clusters). The pairwise anosim section shows the anosim statistic R (upper triangle) and the p value associated with this statistic (lower triangle)

shrimp. Shrimp trawling has occurred across the western Greenland shelf since at least the mid-1950s with intense fishing (>50,000 tonnes/year) occurring from 1975 onwards (Buch et al. 2004; Hammeken Arboe 2014). However, the impacts of otter trawling, and the biological implications of this disturbance, vary widely according to the environmental conditions, substrate types, natural variability and natural disturbance regime of the site in question. Major impacts of trawling on benthic biodiversity and functioning are known from other regions (Engel and Kvitek 1998; Watling and Norse 1998; Freese et al. 1999; Blanchard et al. 2004; Asch and Collie 2008; Bolam et al. 2014). An important (but not the sole) impact of trawling is the direct physical disturbance to the areas fished. Trawling reduces habitat complexity by removing habitat-forming species such as corals and sponges on hard-bottom habitats colonised by these communities. In contrast, experimental trawling on sandy bottoms of offshore fishing grounds have caused declines in some taxa, but not the associated large or long-term changes in benthic assemblages (Løkkeborg 2005), and mixed responses to trawling are reported for motile groups in these habitats (McConnaughey et al. 2000). This reduced impact observed in these soft-bottom communities may indicate some natural resistance to trawling disturbance in areas with higher natural variability (Prena et al. 1999; Kenchington et al. 2001; Kutti et al. 2005). Furthermore, the negative impact of trawling is known to be diminished in areas exposed to high natural disturbance regimes, including wave action, fluctuations in salinity and temperature (Auster and Langton 1999). The western shelf of Greenland is subject to significant seasonal ice flows, with observations of ice scouring at depths exceeding 1 km (Kuijpersa et al. 2007). This 1000-m scouring depth may be exceptional, but the target depth of

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the main fishery is 150–600 m, which may be inside the keel depth of larger icebergs. The impacts of otter trawling and the biological implications of this disturbance vary widely, and results of existing impact studies are highly specific to each site. This study does not attempt to incorporate fishing effort into the analysis, but acknowledges that trawling is likely to be a significant driver of community composition. Proper consideration of this variable must begin with a sound understanding of the faunal assemblages as they currently exist and of their spatial and temporal distributions. This study provides the first quantitative description of the epibenthic megafaunal assemblages as they currently exist on the West Greenland continental shelf.

In recent years, international fishery management organisations have recommended the initiation of monitoring of benthic communities as part of a more ecosystem oriented approach to management (FAO 2003). For countries heavily dependent upon fishing industries, such as Greenland, it is now critically important to start documenting temporal and spatial changes in the benthos, parallel to trends in trawling intensity, in order to understand, measure and manage actual disturbance impacts.

Next steps

This study could be improved by further, carefully selected sampling, particularly focussing on countering: the southern over-representation; the restrictions imposed by nighttime sampling; and spatial bias caused by fitting around existing surveys. Identifications could be improved by complementary physical sampling using methods such as grabs or beam trawl. The inclusion of fishing effort as an influencing factor will improve the analysis. These steps will increase the potential to detect future changes and improve our ability to detect environmental drivers of



Fig. 5 Map of stations by cluster groupings. Approximate seabed temperatures are shown for reference (Map projection EPSG: 3411)

composition. Such efforts are recommended by CAFF (Conservation of Arctic Flora and Fauna) and the Circumpolar Biodiversity Monitoring Programme (http:// www.caff.is/monitoring). Indeed, ongoing initiatives sampling trawl bycatch in Norway and Russia are being expanded to other Arctic territories including Greenland, which will serve to enhance the results of the present study.



Fig. 6 *Plot* of first two axes of multidimensional scaling (MDS) analysis, based on the community composition data. *Points* represent stations. Multiple taxa are excluded from the centre of the figure for clarity. *Inset* vectors show directional influence of significant environmental parameters (p < 0.05)

Conclusion

This study represents the first widespread characterisation of epibenthic megafauna on the area of the West Greenland shrimp fishery. Although the region has been heavily impacted by trawl fisheries, these data constitute an important baseline. We conclude that benthic photography can be a useful tool for examining and monitoring seabed diversity.

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References

- Asch RG, Collie JS (2008) Changes in a benthic megafaunal community due to disturbance from bottom fishing and the establishment of a fishery closure. Fish Bull 106:438–456
- Auster PJ, Langton RW (1999) The effects of fishing on fish habitat. Am Fish Soc Symp 22:150–187
- Bergström BI (2000) The biology of *Pandalus*. Adv Mar Biol 38:55–245
- Blanchard F, LeLoc'h F, Hily C, Boucher J (2004) Fishing effects on diversity, size and community structure of the benthic invertebrate and fish megafauna on the Bay of Biscay coast of France. Mar Ecol Prog Ser 280:249–260
- Blicher ME, Sejr MK (2011) Abundance, oxygen consumption and carbon demand of brittle stars in Young Sound and the NE Greenland shelf. Mar Ecol Prog Ser 422:139–144
- Blicher ME, Rysgaard S, Sejr MK (2007) Growth and production of sea urchin Strongylocentrotus droebachiensis in a high-Arctic

fjord, and growth along a climatic gradient (64°–77°N). Mar Ecol Prog Ser 341:89–102

- Blicher ME, Rasmussen LM, Sejr MK, Merkel F, Rysgaard S (2011) Abundance and energy requirements of eiders (*Somateria* spp.) suggest high predation pressure on macrobenthic fauna in a key wintering habitat in SW Greenland. Polar Biol 34:1105–1116
- Boertmann D, Mosbech A (eds) (2011) Eastern Baffin Bay-A strategic environmental impact assessment of hydrocarbon activities. DCE Danish Centre for Environment and Energy, Aarhus University, Aarhus
- Bolam SG, Coggan RC, Eggleton J, Diesing M, Stephens D (2014) Sensitivity of macrobenthic secondary production to trawling in the English sector of the Greater North Sea: a biological trait approach. J Sea Res 85:162–177
- Buch E, Pedersen SA, Ribergaard MH (2004) Ecosystem variability in West Greenland waters. J Northwest Atl Fish Sci 34:13–28
- Buhl-Mortensen L, Vanreusel A, Gooday AJ, Levin LA et al (2010) Biological structures as a source of habitat heterogeneity and biodiversity on the deep ocean margins. Mar Ecol 31:21–50
- Chazdon RL, Chao A, Colwell RK, Lin S-Y, Norden N, Letcher SG, Clark DB, Finegan B, Arroyo JP (2011) A novel statistical method for classifying habitat generalists and specialists. Ecology 92:1332–1343
- Clarke KR (1993) Non-parametric multivariate analyses of changes in community structure. Aust J Ecol 18:117–143
- Collie JS, Escanero GA, Valentine PC (2000) Photographic evaluation of the impacts of bottom fishing on benthic epifauna. ICES J Mar Sci: J Conseil 57:987–1001
- Compton TJ, Bowden DA, Roland Pitcher C, Hewitt JE, Ellis N (2013) Biophysical patterns in benthic assemblage composition across contrasting continental margins off New Zealand. J Biogeog 40:75–89
- Dauvin JC, Méar Y, Murat A, Poizot E, Lozach S, Beryouni K (2013) Interactions between aggregations and environmental factors explain spatio-temporal patterns of the brittle-star *Ophiothrix fragilis* in the eastern Bay of Seine. Est Coast Shelf Sci 131: 171–181
- Engel J, Kvitek R (1998) Effects of Otter Trawling on a Benthic Community in Monterey National Marine Sanctuary. Soc Conserv Biol 12:1204–1214
- Faith DP, Minchin PR, Belbin L (1987) Compositional dissimilarity as a robust measure of ecological distance. Vegetatio 69:57–68
- FAO (2003) The Ecosystem Approach to Fisheries. FAO Technical Guidelines for Responsible Fisheries no. 4, suppl. 2. FAO, Rome
- Freese L, Auster PJ, Heifetz J, Wing BL (1999) Effects of trawling on seafloor habitat and associated invertebrate taxa in the Gulf of Alaska. Mar Ecol Prog Ser 182:119–126
- Hammeken Arboe N (2014) The Fishery for Northern Shrimp (*Pandalus borealis*) off West Greenland, 1970-2014. NAFO SCR Doc. 14/061
- Jørgensen OA, Tendal OS, Hammeken Arboe N (2013) Preliminary mapping of the distribution of corals observed off West Greenland as inferred from bottom trawl surveys 2010-2012. NAFO SCR Doc. 13/07
- Jørgensen LL, Ljubin P, Skjoldal HR (2015) Distribution of benthic megafauna in the Barents Sea: baseline for an ecosystem approach to management. ICES J Mar Sci 72:595–613
- Kenchington EL, Prena J, Gilkinson KD et al (2001) Effects of experimental otter trawling on the macrofauna of a sandy bottom ecosystem on the Grand Banks of Newfoundland. Can J Fish Aquat Sci 58:1043–1057
- Kuijpersa A, Dalhoffa F, Brandtb MP, Hümbsc P, Schottd T, Zotovae A (2007) Giant iceberg plow marks at more than 1 km water depth offshore West Greenland. Mar Geol 246:60–64
- Kutti T, Høisæter T, Rapp HT, Humborstad O-B, Løkkebor S, Nøttestad L (2005) Immediate effects of experimental otter

trawling on a sub-arctic benthic assemblage inside Bear Island Fishery Protection Zone in the Barents Sea. Am Fish Soc Symp 41:519–528

- Lassen AH, Powles H, Bannister C, Knapman P (2013) Marine Stewardship Council (MSC) Final Report and Determination for the West Greenland Cold Water Prawn Trawl Fishery Client. (http://www.msc.org/track-a-fishery/fisheries-in-the-program/cer tified/arctic-ocean/West-Greenland-Coldwater-Prawn/assessmentdownloads-1/20130122_FR_PRA126.pdf)
- Legendre P, Gallagher ED (2001) Ecologically meaningful transformations for ordination of species data. Oecologia 129:271–280
- Levin LA, Dayton PK (2009) Ecological theory and continental margins: where shallow meets deep. Trends Ecol Evol 24: 606–617
- Løkkeborg S (2005) Impacts of trawling and scallop dredging on benthic habitats and communities. FAO Fisheries Technical Paper No 472. FAO, Rome
- Magurran AE, McGill B (2010) Biological Diversity: Frontiers in Measurement and Assessment. Oxford University Press, Oxford
- McConnaughey RA, Mier KL, Dew CB (2000) An examination of chronic trawling effects on soft-bottom benthos of the eastern Bering Sea. ICES J Mar Sci 57:1377–1388
- Myers PG, Kulan N, Ribergaard MH (2007) Irminger Water variability in the West Greenland Current. Geophys Res Lett 34: L17601
- Oksanen J, Blanchet FG, Kindt R, Legendre P, Minchin PR, O'Hara RB, Simpson GL, Solymos P, Stevens MHH, Wagner H (2013) vegan: Community Ecology Package. R package version 2.0-10. (http://CRAN.R-project.org/package=vegan)
- Piepenburg D (2000) Arctic brittlestars. Oceanogr. Mar Biol 38:189– 256
- Piepenburg D, Schmid MK (1996) Distribution, abundance, biomass, and mineralization potential of the epibenthic megafauna of the Northeast Greenland shelf. Mar Biol 125:321–332
- Piepenburg D, Archambault P, Ambrose WG et al (2011) Towards a pan-Arctic inventory of the species diversity of the macro- and megabenthic fauna of the Arctic shelf seas. Mar Biodivers 41:51–70
- Prena J, Schwinghammer P, Rowell TW, Gordon DC Jr, Gilkinson KD, Vass WP, McKeown DL (1999) Experimental otter trawling on a sandy bottom ecosystem of the Grand Banks of Newfoundland: analysis of trawl bycatch and effects on epifauna. Mar Ecol Prog Ser 181:107–124
- Renaud PE, Morata N, Ambrose WG, Bowie JJ, Chiuchiolo A (2007) Carbon cycling by seafloor communities on the eastern Beaufort Sea shelf. J Exp Mar Biol Ecol 349:248–260
- Robert K, Jones D, Huvenne V (2014) Megafaunal distribution and biodiversity in a heterogeneous landscape: the iceberg-scoured Rockall Bank, NE Atlantic. Mar Ecol Prog Ser 50:67–88
- Schmid M, Piepenburg D (1993) The benthos zonation of the Disko Fjord, West Greenland. Medd Grønl 37:1–21
- Sejr MK, Jensen KT, Rysgaard S (2000) Macrozoobenthic community structure in a high-arctic East Greenland fjord. Polar Biol 23:792–801
- Sejr MK, Blicher ME, Rysgaard S (2009) Sea ice cover affects interannual and geographic variation in growth of the Arctic cockle *Clinocardium ciliatum* (Bivalvia) in Greenland. Mar Ecol Prog Ser 389:149–158
- Sejr MK, Włodarska-Kowalczuk M, Legeżyńska J, Blicher ME (2010) Macrobenthic species composition and diversity in the Godthaabsfjord system, SW Greenland. Polar Biol 33:421–431
- Sparks-McConkey PJ, Watling L (2001) Effects on the ecological integrity of a soft-bottom habitat from a trawling disturbance. Hydrobiologia 456:73–85
- Starmans A, Gutt J, Arntz WE (1999) Mega-epibenthic communities in Arctic and Antarctic shelf areas. Mar Biol 135:269–280

- Trebitz AS, Hill BH, McCormick FH (2003) Sensitivity of indices of biotic integrity to simulated fish assemblage changes. Environ Manage 32:499–515
- Watling L, Norse EA (1998) Disturbance of the seabed by mobile fishing gear: a comparison to forest clearcutting. Conserv Biol 12:1180–1197