

The sipunculan fauna of Svalbard

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Abstract

This study presents the species of Sipuncula collected in the Svalbard area (74–81°N and 10–35°E) in the summer seasons from 1996 until 2005 at depths ranging from 40 to 2553 m. The faunistic analysis of the material (1056 specimens from 251 stations) resulted in a total of six species and one subspecies, belonging to two families (Golfingiidae and Phascolionidae). One species, *Golfingia vulgaris*, has not been reported previously from Svalbard waters, and increases the total number of Sipuncula taxa known from Svalbard to nine. Three species dominated the collected material: *G. vulgaris* (53.5% of all specimens found), *G. margaritacea* (19.3%) and *Nephasoma diaphanes diaphanes* (15.5%). The study shows that compared with other northern regions, Svalbard hosts a relatively rich sipunculan fauna, which is most similar in species composition to the sipunculan fauna found in Asian Arctic waters. An easy-to-use identification key to Svalbard Sipuncula species is given to aid field researchers in the identification of this often overlooked taxon.

Sipuncula is a phylum of exclusively marine worms. According to recent revisions, it consists of two classes, six families, 17 genera and about 150 species and subspecies (Cutler 1994). Sipunculans inhabit most marine habitats from intertidal zones to abyssal depths (Cutler 1994). Sipuncula are predominantly tropical and subtropical worms (75% of the species). Boreal species account for 19% of the species, and 3% are bipolar and cosmopolitan species. Another 3% occur in Arctic, Antarctic and temperate–cold waters (Murina 1975). Sipuncula are mostly deposit feeders living in various substrata, mainly containing mud, silt and sand. Some species may seek a protective shelter in molluscan shells, polychaete tubes and foraminiferan tests (Murina 1984).

Sipuncula are very often neglected in ecological studies, although they may play a considerable role in the bioerosion of coral reefs and soft rocks (Cutler 1968; Stearley & Ekdale 1989), as food for fish, gastropods and occasionally people (Kohn 1975), and probably as bioturbators (Murina 1984; Romero-Wetzel 1987). In Arctic fjords Sipuncula may play an important role in macrobenthic communities and are among the dominant taxa (e.g. Węśławski et al. 1988; Włodarska-Kowalczyk & Pearson 2004; Włodarska-Kowalczyk et al. 2004), but

usually they are identified only to the phylum level. Even though a number of publications on Sipuncula from Arctic areas have been published (Théel 1905; Wesenberg-Lund 1930, 1933, 1934, 1938, 1955; Murina 1977), almost all information on Sipuncula in Svalbard comes from studies conducted at the beginning of the last century (Théel 1905; Fischer 1895, 1914, 1922, 1929). Some remarks on Sipuncula are made in Gulliksen et al. (1999), although their work should be treated with caution as their information on Svalbard species, including Sipuncula, is mostly based on the early published sources, which may be of questionable quality. Further information on sipunculans (mainly concerning species found in Asian Arctic Shelf seas and the Barents Sea) has been published by Murina (1977).

The main aim of this article is to present a synopsis of the composition and distribution of the Sipuncula in the waters of the Svalbard archipelago based on our own samples and published data. Species composition of Sipuncula found in the Svalbard area is compared with their distribution in the North Atlantic, Arctic seas and the waters around Antarctica. An identification key of Svalbard Sipuncula species is given to aid field researchers in the identification of this often overlooked taxon.

Material and methods

The Svalbard archipelago is located between 74 and 81°N and between 10 and 35°E. It is influenced mainly by two water masses, the warm West Spitsbergen Current (temperature above 2°C and salinity of 35 PSU (practical salinity unit)) and the cold East Spitsbergen Current (from -1.5°C to 1°C, 34–35 PSU) (Loeng 1991; Beszczynska-Möller et al. 1997). The West Spitsbergen Current is the northernmost extension of the Norwegian Atlantic Current and keeps the area free of ice throughout most of the year. The East Spitsbergen Current carries Arctic Water between Spitsbergen and Franz Josef Land, and then southward along the eastern coast of Spitsbergen (Loeng 1991). The Svalbard coast is subjected to ice scour from drifting ice in the summer months, whereas fast ice forms in the inner fjords in winter (Węśławski et al. 1988; Svendsen et al. 2002).

Material was collected during cruises with the RV *Oceania* (1996 to 2005) and RV *Polarstern* (ARK XVI and ARK XIX expeditions, deep-sea samples). Samples were mostly taken in the fjords of Kongsfjorden and Hornsund but were also taken in Van Mijenfjorden and Adventfjorden. Additional samples were taken in Magdalenefjorden, Storfjorden, Erik Eriksenstretet and near Kong Karls Land (material provided by Akvaplan-niva, Polar Environmental Center, N-9296, Tromsø, Norway) (Fig. 1; Table 1).

A total of 251 stations were sampled, 450 samples were taken (Fig. 1) and 1056 specimens were identified. Most

(378 of 450) samples were taken with a van Veen grab (0.1-m² catch area), the remainder were taken with either dredges (74 samples) or box corers (14 samples). Material was washed on a 0.5-mm sieve and fixed in a 4% buffered formaldehyde and sea water solution. All organisms were later sorted and *Sipuncula* specimens were picked. The taxonomic nomenclature is adopted from Cutler (1994). The material is deposited at the Institute of Oceanology, Polish Academy of Science, in Sopot, Poland.

Cluster analysis (using group-average linking) was performed on the presence/absence data. To perform the cluster analysis, 23 species were included in total, of which nine were reported from Antarctica only. The detailed matrix with information on species occurrence in different polar areas is presented in Fig. 2. The similarities between samples were calculated using the Bray-Curtis index. Multivariate analyses were performed with the use of the PRIMER package version 6 (Clarke & Warwick 2001).

Results

Sipuncula specimens were not very numerous and were found only at 115 of the 251 (45.8%) stations. At some stations *Sipuncula* species represented a significant part of the benthos samples and were among the dominant species. Especially rich were samples collected at the deep

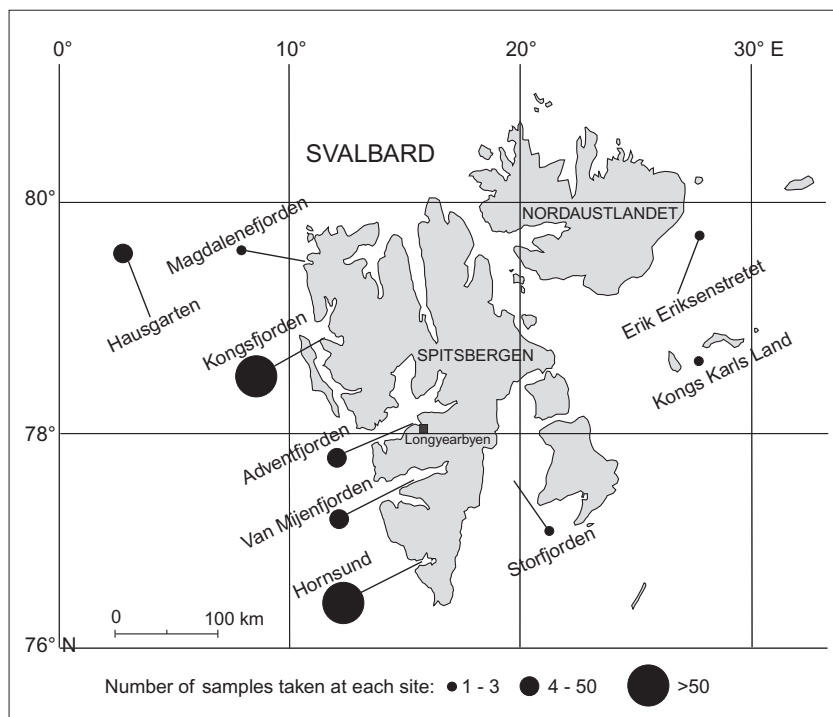


Figure 1 Location of sampling sites in the waters off Svalbard.

Table 1 Basic information about cruises and sampling details.

Cruise information (no. of cruises)	Area	Sampling year	Sampling gear	Number of stations	Number of samples	% of stations where Sipuncula were found
RV <i>Oceania</i> (10)	Kongsfjorden	1996–2005	van Veen	70	155	52.4
		1996; 1997; 1999	dredge	63	63	27.7
	Hornsund	2002; 2003; 2005	van Veen	68	102	22.7
		2003	dredge	11	11	57.1
	Van Mijenfjorden	2000; 2001	van Veen	10	44	70
Adventfjorden	2004	van Veen	6	20	66.6	
RV <i>Polarstern</i> (2) ARK XVI & ARK XIX	Hausgarten	2000; 2003	Box corer	14	14	57.1
Akvaplan-niva (1)	Storfjorden	1996	van Veen	3	15	100
	Kong Karls Land			1	5	100
	Magdalenefjorden			3	15	33.3
	Erik Eriksenstretet			2	10	66.6
Total (13)				251	450	45.8

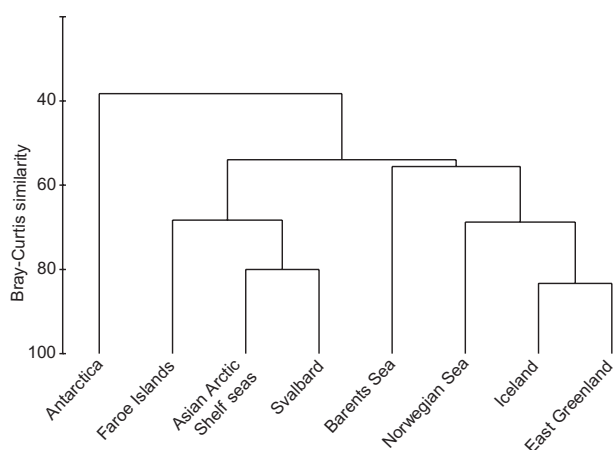


Figure 2 Dendrogram showing the faunistic resemblances in the composition of the sipunculan fauna of various northern polar and subpolar regions: Faroe Isles (Murina & Sørensen 2004); Iceland (Wesenberg-Lund 1937b); East Greenland (Wesenberg-Lund 1937a); Norwegian Sea (Murina 1977); Barents Sea (Sirenko 2001); Asian Arctic Shelf seas (Sirenko 2001); Antarctica (Saiz-Salinas 1995; Saiz-Salinas & Pagola-Cardé 1999); and Svalbard (this study).

sea sites, at Hausgarten, (a maximum of 404 specimens per 1 m² were found) and in Kongsfjorden (a maximum of 840 and 520 specimens per 1 m² were found at the two sites sampled). Nevertheless, at most of the stations we found only a few specimens or a single specimen. Table 2 shows the distribution, type of substrata on which particular species were found and abundances of Sipuncula species in Svalbard waters.

Seven species have been reported from the Svalbard archipelago in the past (Table 3). We have recognized seven taxa: six species and one subspecies, belonging to two families (Golfingiidae and Phascolionidae), of which two have not been reported previously from the Svalbard

area (*Golfingia vulgaris* and *Nephasoma diaphanes corrugatum*). In the collected material three species dominated among the Sipuncula fauna found in this study: *G. vulgaris* (53.5% of all specimens found), *G. margaritacea* (19.3%) and *N. diaphanes diaphanes* (15.5%). *G. elongata* was the rarest species, occurring at only seven stations (nine specimens—less than 1% of all specimens found).

For each species found in our material some remarks on their distribution and biotope are provided. Table 4 contains the information on the depth distribution in the World Ocean, the North Atlantic, the Arctic and Svalbard waters (this study).

Golfingia margaritacea margaritacea (Sars, 1851)

This cold water and cosmopolitan species is widely distributed. Many reports are from cold and temperate (boreal) regions, especially in the shelf and bathyal zone. It has rarely been found in the tropical and subtropical zone. It occurs in the Atlantic, Arctic and Antarctic oceans (from 80°N to 78°S) as well as in the Pacific ocean (above 30°N and 30°S), but it is unknown in the Indian Ocean. It inhabits mud and sand/gravel bottoms, and red clay in the abyssal (Gibbs 2001).

Golfingia vulgaris vulgaris (De Blainville, 1827)

This cold water and cosmopolitan species is encountered extensively in the World Ocean, particularly in the north-eastern Atlantic, including the waters off Greenland, Scandinavia and the British Isles, as well as in the Mediterranean and Red seas. Its absence from the western Atlantic Ocean is noteworthy. This species occurs rarely in the Arctic zone: there are some reports from the Barents, East Siberian and Chukotsk seas. It usually inhabits silt, mud, different types of sand and clay in deep waters (Saiz-Salinas & Villafranca Urchegui 1990).

Table 2 Distribution of Sipuncula in Svalbard waters (number of specimens found in each area) and the substrate types in which specimens were found.

	K	H	VM	A	EE	S	M	KK	HG	Total
<i>Golfingia margaritacea</i> mud	11	182	10			1				204
<i>G. vulgaris</i> mud, sandy mud, bioturbated mud, clay/mud (30–40% clay)	382	161	1	5	2	2	2	10		565
<i>G. elongata</i> mud and sandy mud	5	5								10
<i>Nephasoma lilljeborgi</i> silty/sandy clay									55	55
<i>N. diaphanes diaphanes</i> sandy mud, clay, mud	43	3	1	2	2				113	164
<i>N. diaphanes corrugatum</i> sandy mud, mud	3	2								5
<i>Phascolion strombus</i> sand, muds, sandy silt	41	3			2	7				53

K, Kongsfjorden; H, Hornsund; VM, Van Mijenfjorden; A, Adventfjorden; EE, Erik Eriksenstretet; S, Storfjorden; M, Magdalenefjorden; KK, Kong Karls Land; HG, Hausgarten.

Table 3 Sipunculan species reported from the Svalbard archipelago by various sources.

Species	Fischer 1895	Théel 1905	Fischer 1914	Fischer 1922	Fischer 1929	Murina 1977	Gulliksen et al. 1999	This study
<i>Golfingia margaritacea</i>		X	X	X	X	X	X	X
<i>G. vulgaris</i>								X
<i>G. elongata</i>						X		X
<i>Nephasoma diaphanes diaphanes</i>				X	X		X	X
<i>N. diaphanes corrugatum</i>								X
<i>N. lilljeborgi</i>	X	X			X			X
<i>N. eremita</i>		X		X	X	X	X	
<i>N. abyssorum</i>	X			X	X	X		
<i>Phascolion strombus</i>	X	X		X	X	X	X	X

Table 4 Depth range (m) of Sipuncula taxa found in the World Ocean (Murina 1977; Cutler 1994; Murina, unpubl. data), North Atlantic from 60°N to 40°N (Cutler 1994; Murina 1977; Murina, unpubl. data), North Atlantic >60°N (Murina 1977; Murina, unpubl. data) and in Svalbard waters (this study).

Taxon	World Ocean		North Atlantic 60°–40°N		North Atlantic >60°N		Svalbard	
	min.	Max.	min.	max.	min.	max.	min.	max.
<i>Golfingia margaritacea</i>	0	5300	72	4600	8	3230	67	290
<i>G. vulgaris</i>	5	2000	37	248	15	1480	64	374
<i>G. elongata</i>	10	590	0	73	?	?	79	162
<i>Nephasoma diaphanes diaphanes</i>	0	5300	10	4800 ^a	446	2701	40	2553
<i>N. diaphanes corrugatum</i>	80	5900						
<i>N. lilljeborgi</i>	65	2734			40	2200	1912	2553
<i>N. eremita</i>	0	3867			73	515	—	—
<i>N. abyssorum</i>	95	5300			95	973	—	—
<i>Phascolion strombus</i>	1	4030	9	2090	15	1890	79	390

^aIn the Iberian trench.

***Golfingia elongata* (Keferstein, 1862)**

This tropical and boreal species is cosmopolitan in the Northern Hemisphere, in the north-western Atlantic (from Newfoundland to Bermuda and Cuba) and north-eastern Atlantic (from Spitsbergen to the Iberian Peninsula and in the Mediterranean Sea) and in the Pacific Ocean (eastern and southern China and Vietnam). It is rare in the Arctic zone: Greenland, Spitsbergen and Iceland (Cutler 1994). It prefers the upper sublittoral (Saiz-Salinas & Villafranca Urchegui 1990). It often inhabits hard ground, gravel, sand, shelf rock; less often it inhabits silt, clay and mud (Cutler 1994; Saiz-Salinas & Villafranca Urchegui 1990).

***Nephasoma diaphanes diaphanes* (Gerould, 1913)**

This is a cosmopolitan species with widely ranging vertical distribution. It is the most common sipunculan in deep-sea communities (Cutler & Cutler 1987a). In Arctic waters it was reported from Spitsbergen, Novaya Zemlya and the Barents Sea. This species inhabits silt, sand and mud, and is often found in foraminiferal tests, small polychaete tubes and mollusc shells (Cutler 1994). In this study it was usually found in the shelter of foraminiferal tests.

Cutler & Cutler (1986) regarded *N. minutum* (Keferstein 1862) as a hermaphrodite species confined to the north-eastern Atlantic. In contrast, *N. diaphanes*, which is morphologically and anatomically similar to *N. minutum*, is dioecious and cosmopolitan (Cutler & Cutler 1986). Murina & Sørensen (2004) considered *N. diaphanes* as a junior synonym of *N. minutum* and agreed with Gibbs' (1975) opinion: "The morphologically similar specimens now regarded as a single species have two distinct forms, a dioecious form found chiefly in deep water, and a hermaphroditic form found on the shore and in shallow water around Europe and Scandinavia". Accordingly, the name *N. diaphanes* is used in this paper.

***Nephasoma diaphanes corrugatum* Cutler and Cutler, 1986**

This subspecies is widely distributed in the Atlantic and Pacific oceans, as well as in the Mediterranean and Red seas.

***Nephasoma lilljeborgi* (Danielssen and Koren, 1880)**

This species is of Arctic/boreal origin. According to Cutler & Cutler (1987a) it is only known to inhabit the northern deep-sea basins in the far north-eastern Atlantic Ocean

between 70° and 80°N, and might therefore be endemic to these regions (Cutler 1994). It was reported from Spitsbergen, Greenland, Baffin Bay, the Norwegian coast, the western part of Sweden, the eastern part of Iceland and the Russian seas (the Kara, Laptev, East Siberian and Chukchi seas). It inhabits muds, clay and sandy mud.

Nephasoma diaphanes often occurred with *N. lilljeborgi* in the same samples, although *N. diaphanes* is a more eurybathic species that was found at a wider range of depths (40–2553 m).

***Phascolion strombus strombus* (Montagu, 1804)**

This is a cosmopolitan but mainly cold water species that is widely encountered in all oceans (rare findings in the Indian Sea). It is common in Russian arctic seas. It prefers the littoral and sublittoral, penetrates in bathyal and is rare in abyssal zones. Usually it inhabits sand, mud and silt, and is often found in mollusc shells and polychaete tubes (Hylleberg 1970). In this study it was found sometimes in Gastropoda shells and Polychaeta tubes.

Discussion

The results of this study show that the Svalbard area hosts a relatively rich (for the High Arctic region) sipunculan fauna. We increased the total number of sipunculan taxa now known from the Svalbard area from seven to nine. Cluster analysis of Sipuncula assemblages from different regions showed that the Sipuncula species composition of the Svalbard region is most similar to the one from the Asian Arctic seas. The sipunculan fauna includes members of the same taxa in both areas except for *G. elongata*, which has not been reported from the Asian Arctic seas (Sirenko 2001). The Barents Sea (Sirenko 2001) clusters (although not very well) with the other North Atlantic areas except for the Faroe Islands. A relatively low number of sipunculan taxa (six) was reported from East Greenland (Wesenberg-Lund 1937a) and Iceland (Wesenberg-Lund 1937b). The Arctic regions are well separated from the Antarctic, where twice as many species of Sipuncula are found (Saiz-Salinas 1995; Saiz-Salinas & Pagola-Carte 1999; Cutler et al. 2001) (Fig. 2).

In general, there is an increase in species richness from high to lower latitudes. This trend has been shown for many groups of marine organisms, both benthic and planktonic (Stevens 1989; Angel 1996; Ormond et al. 1997). Even though the paradigm of the latitudinal distribution of species richness is still subject to discussion (Gray 2001), Sipuncula seem to follow this rule. The phylum is undoubtedly centred in warmer waters, where the great majority (75%) of species is found. The highest

species diversity is found in the Indo-West Pacific and particularly in the Indo-Malayan archipelago. These regions are thus regarded as the main centre of origin and development of *Sipuncula* (Murina 1975). In the Atlantic Ocean the number of *Sipuncula* species decreases towards higher latitudes, although the decrease towards the south is not as strong as towards the north. In the North Atlantic the diversity of *Sipuncula* is highest off Ireland (21 species) and in the North Sea (17 species). It decreases with latitude to Svalbard (eight species) and the Asian Arctic seas (seven species). Even though more species are reported from the Antarctic area (16) (Saiz-Salinas 1995; Saiz-Salinas & Pagola-Cardé 1999; Cutler et al. 2001), there is a progressive loss of genera and species along a polar gradient from locations above the Antarctic Convergence to the high latitudinal Antarctic regions (Saiz-Salinas 1995). The effects of extreme cold temperatures appear to be the main structuring factor, although other, as yet unknown, factors may be of importance in determining this biodiversity gradient (Saiz-Salinas & Pagola-Cardé 1999).

The higher number of *Sipuncula* species in the Southern Ocean than in the Arctic may be caused by the greater age of the former region. Antarctica has been isolated for about 40 million years (Dunton 1992), whereas the Arctic Ocean has existed for only 2–3 million years (Dayton et al. 1994). The fact that the Southern Ocean covers a larger area may contribute to its greater species richness. Bouchet & Waren (1979) supposed that in addition to the effects of age and isolation, the homogeneity and absence of geographic isolating barriers within the Arctic abyssal area could result in low species richness. The Arctic Ocean fauna, located between two major oceans, is young and of mixed origins. Repeated Pleistocene glaciations were lethal to inhabitants of the Arctic Ocean and are the reason for the youth of the fauna and its low diversity (Dunton 1992). The harshness and relative homogeneity of habitats in the Arctic are also common explanations (Clarke & Crame 1997).

Sipunculans inhabit all of the seas of the world and are found at all latitudes. The zoogeographical range of many species is difficult to define because of their tolerance of a broad range of temperatures and depths. Moreover, many have planktonic larvae that play a very important role in determining their wide geographical distribution (Scheltema & Hall 1975; Scheltema & Rice 1990). Teleplanic larvae of Sipunculans are abundant across entire ocean basins (Scheltema 1975; Scheltema & Rice 1990). Ranges appear to be determined by water temperature, bottom topography and water currents (Cutler & Cutler 1987b). The sipunculan fauna on most oceanic islands appears to include mostly species with wide geographical distributions, and very few endemic species are known

(Scheltema & Hall 1975). However, the potential larval dispersal may be quite different from the realized larval dispersal: many marine species with teleplanic larvae may have more limited ranges than are hypothesized on the basis of their potential for larval dispersal. Furthermore, either potential or realized larval dispersal does not necessarily confer genetic homogeneity across a species with a large range (Staton & Rice 1999). There is evidence for cryptic speciation in *Apionsoma misakianum* (Staton & Rice 1999), and species delimitation may not be as clear as previously thought, especially for disjunct “populations” (Maxmen et al. 2003). However, it is unusual to find two morphologically cryptic species that are not sister species (Schulze et al. 2007). All polyphyletic species examined by Schulze et al. (2007) were geographically widespread and inhabit hard substrates—mostly coral rubble—in shallow waters, and belong to genera that do not occur in polar areas.

No endemism is shown by either Antarctic or Arctic *Sipuncula* species (except for *N. lilljeborgi*, which might be endemic to cold deep waters). Many of the most abundant species in both areas (*G. margaritacea*, *N. diaphanes* and *P. strombus*) are cosmopolitan (Saiz-Salinas & Pagola-Cardé 1999). The most numerous biogeographical group of *Sipuncula* species found off Svalbard are widespread in the Atlantic Ocean, occurring at a range of depths and in a wide variety of substrata. *G. margaritacea* and *P. strombus* have a very wide distribution in the Atlantic Ocean, but live at greater depths at lower latitudes (Cutler 1994). In Svalbard waters they were found only below 400 m. One species (*G. elongata*) is a tropical/boreal one, but in Svalbard waters it seems to occur rarely.

The genus *Nephasoma* is especially richly represented in cold bathyal and abyssal waters of the northern Atlantic and Pacific (Cutler 1994). *N. abyssorum* has repeatedly been reported from Svalbard (Fischer 1895, 1922, 1929) but was not found in this study. It is a common bathyal and abyssal species in the north-eastern Atlantic and the Arctic Ocean. It is very likely that it is still present in the Svalbard area but has been missed in our survey because of the scarcity of deep-sea samples. The same conclusion applies to *N. eremita*. This bipolar species has previously been reported from Svalbard (Théel 1905; Fischer 1922, 1929; Gulliksen et al. 1999) but did not occur in our samples (Table 3).

It is very likely that the poor knowledge regarding the *Sipuncula* may lead to erroneous statements regarding the occurrence of sipunculans in Arctic regions. That was probably the reason that *G. vulgaris* has not been reported from Svalbard, although it is a widely distributed species in Greenland, Scandinavia (Théel 1905; Wesenberg-Lund 1925, 1937a; Cutler 1994) and in many Arctic seas (Sirenko 2001).

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Appendix. Simplified identification key to families, genera and species of Sipuncula from the Arctic region (Svalbard species are indicated in boldface).

Key to families of class Sipunculidea:

1. Longitudinal muscles of body wall gathered into either separate or anastomosing bands (20–24)
 - Longitudinal muscles of body wall in uniform continuous layer
2. A single nephridium present
 - Two nephridia present

Sipunculus norvegicus
2
Phascolionidae
Golfingiidae

Key to Golfingiidae:

1. Four introvert retractor muscles present
 - Two introvert retractor muscles present

Golfingia
Nephasoma

Key to *Golfingia* species

1. Introvert hooks in rings
 - Introvert hooks scattered, if present
2. Hooks present and scattered; central part of the trunk smooth, often transparent, both trunk ends dark and heavy papillated
 - Hooks on introvert absent; small papillae present on the trunk

Golfingia elongata
2
Golfingia vulgaris (Fig. 3)

Golfingia margaritacea (Fig. 4)

Key to *Nephasoma* species

1. Hooks absent
 - Hooks scattered
 - Hooks arranged in rings or otherwise
2. Scattered small hooks; body transparent to translucent, usually <10 mm long
 - Scattered small triangular hooks; body opaque, up to 40 mm long. From bathyal depths
3. Body skin smoothly glossy and transparent; introvert with a few short tentacular lobes
 - Body skin translucent to opaque, with irregular longitudinal epidermal ridges on the introvert base and the anterior part of the trunk
4. Large hooks (>50 µm) in a spiral arrangement, like a barber's pole

Nephasoma eremita (Fig. 5)
2
4
3

Nephasoma lilljeborgi (Fig. 6)

Nephasoma diaphanes diaphanes (Fig. 7)

Nephasoma diaphanes corrugatum

Nephasoma abyssorum

Key to Phascolionidae:

1. Epidermal holdfast papillae present
2. Epidermal holdfast papillae absent

Phascolion
Onchnesoma

Key to *Phascolion* species:

1. Ventral retractor much thinner than dorsal; holdfast papillae V-shaped with dark, hardened border
2. Ventral and dorsal retractors have equal width; large holdfast papillae without hardened borders

P. strombus (Fig. 8)
P. turberculosum

Key to *Onchnesoma* species:

1. Trunk covered with prominent backward directed protrusions; eight tentacles
2. Trunk flat without spherical protrusions or scales; tentacles absent

O. squamatum
O. steenstrupii

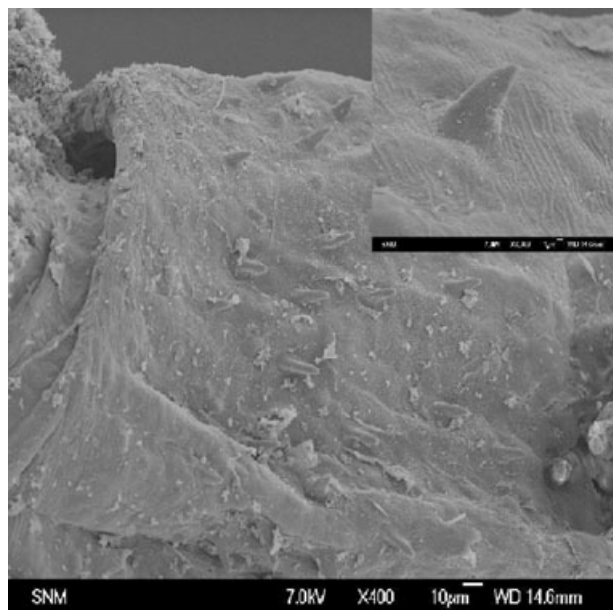


Figure 3 Scanning electron micrographs of the small hooks of *Golfingia vulgaris*. (Scanning electron microscope pictures were taken at the Zoological Museum of the University of Copenhagen.)

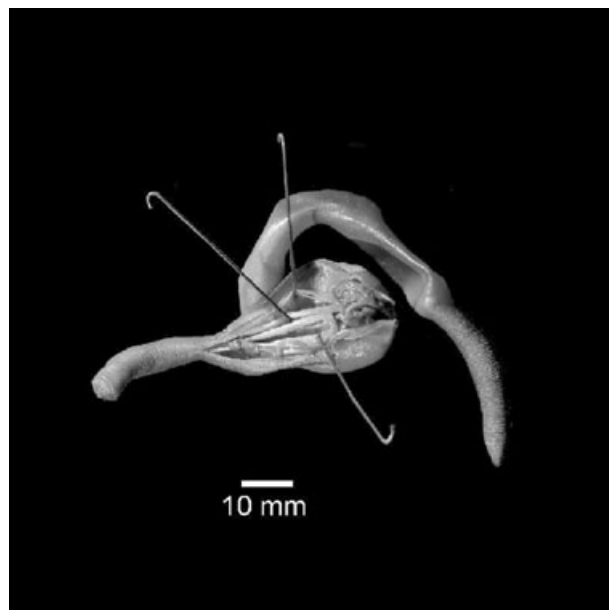


Figure 4 The interior organs of *Golfingia margaritacea* showing four introvert retractor muscles.

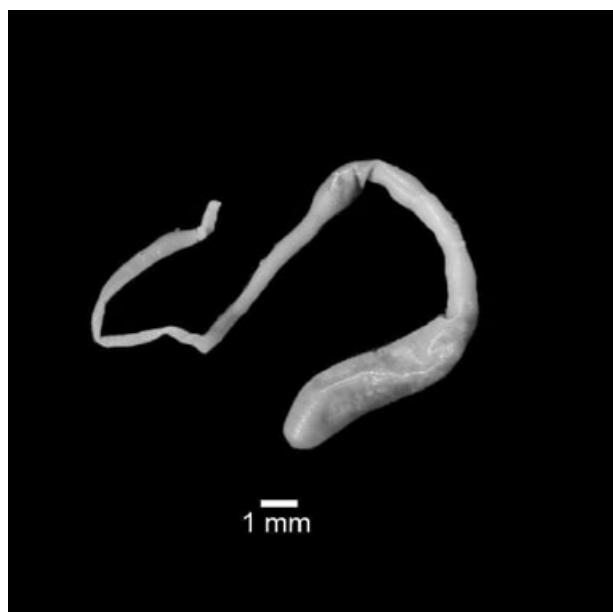


Figure 6 *Nephasoma lilljeborgi*.

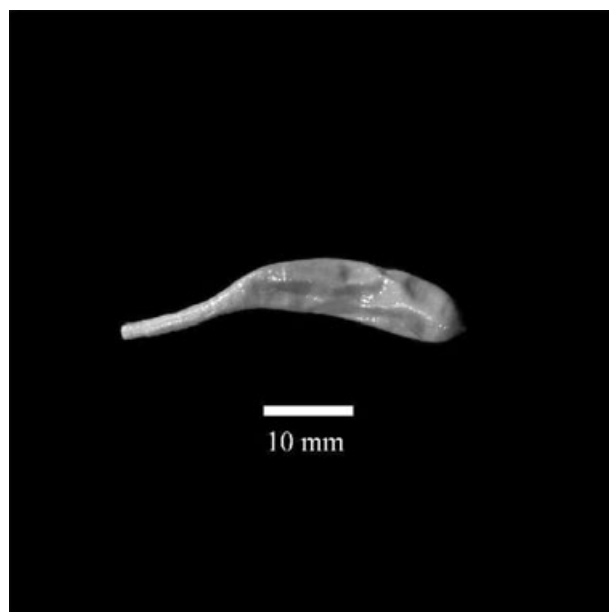


Figure 5 *Golfingia eremita*.

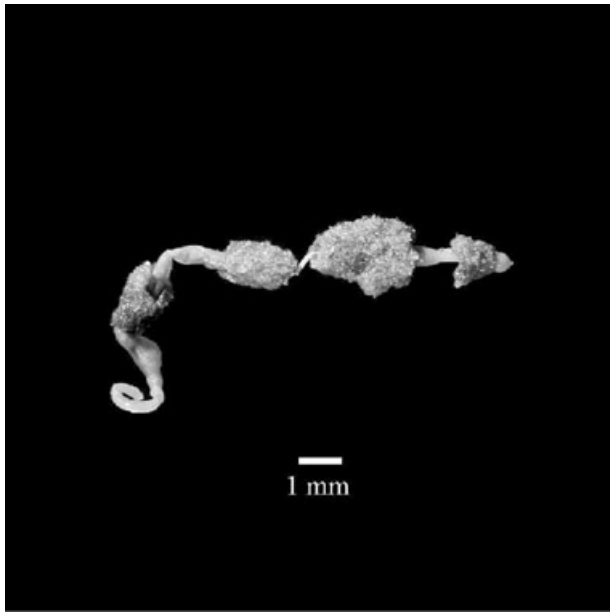


Figure 7 *Nephasoma diaphanes diaphanes* in foraminiferal tests.

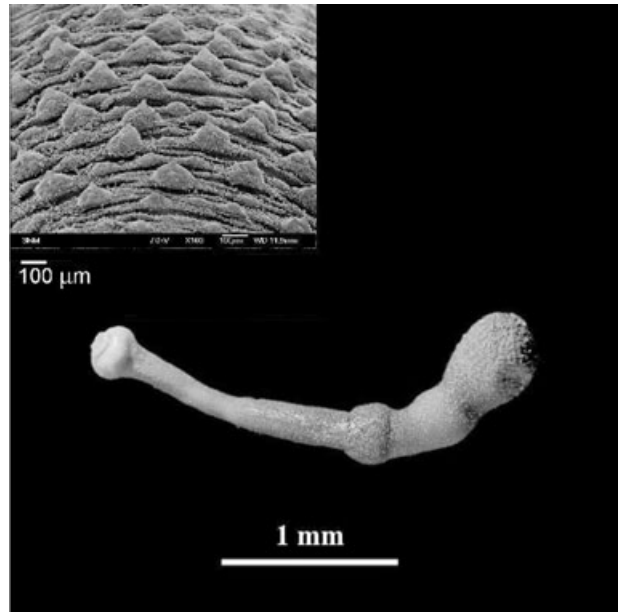


Figure 8 *Phascolion strombus*. Scanning electron micrographs of the holdfast papillae of *P. strombus*.