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Tintinnid Ciliates of the Marine Microzooplankton in Arctic Seas: a Compilation and Analysis of Species Records

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Abstract

We set out to examine a postulated latitudinal trend in species richness within the Arctic Ocean. We compiled species records of tintinnid ciliates in the Arctic from the literature and added our own unpublished Chukchi Sea data to produce a database (available as an Online Resource data file) consisting of 1427 records of 89 species from 414 locations above 65°N sampled from 1885 to 2015. While there was no trend of species richness throughout the Arctic, there was a significant positive relationship between species richness and the number of sites sampled in a 2° band, suggesting a sampling effect. Plotting cumulative numbers of species and cumulative number of sites sampled by year, we found a linear relationship in log cumulative numbers of species and log sites sampled, and a lack of a plateau in the species accumulation trend. Species records are highly dominated by 4 species, accounting for 45% of the records: *Acanthostomella norvegica*, *Parafavella denticulata*, *Ptychocylis obtusa* and *Salpingella acuminata*, all of which, except *S. acuminata*, have long been suspected to be morphologically variable, with different morphotypes given undue species status. Pooling all reported species of *Acanthostomella*, *Parafavella* and *Ptychocylis* yielded little qualitative differences but considerable quantitative differences. We found large discrepancies in geographic coverage. We conclude that many zones projected to experience large changes in sea ice coverage are under-sampled. Based on the historical trend, the list of Arctic tintinnid ciliate species will likely continue to grow with new sampling, regardless of changes in the Arctic Seas.

Keywords: biodiversity, biogeography, plankton, zooplankton, protist, latitudinal diversity gradient

Introduction

Microzooplankton are a key component of planktonic food webs as they are responsible for the consumption of most of the primary production in the plankton, including in the Arctic Seas (Nelson et al. 2014). The relative importance of microzooplankton as grazers in the Arctic seas is predicted to increase due to the foreseen shifts towards smaller phytoplankters (Li et al. 2009) and changes in the timing of phytoplankton blooms (Michel et al. 2012). Tintinnid ciliates are a usually minority component in the microzooplankton (McManus & Santoferarra 2013). However, in the Arctic tintinnid ciliates can occasionally be a significant component of the grazer community (Sherr et al. 2009) and can seasonally dominate the biomass of the ciliate microzooplankton (Seuth et al. 2011). Tintinnids are also a component of the sea ice fauna (e.g., Melnikov et al. 2002). Among the microzooplankton they are unique in possessing a cylindrical or cup-shaped species-specific lorica (shell) permitting species identification with relative ease compared to most other taxa of the microzooplankton. Furthermore, an easily measured dimension of the lorica, the size of oral opening, is correlated with size of preferred prey and maximum growth rate (Montagnes 2013). Species with similar lorica oral diameters share then key ecological characteristics and can be considered as ecological redundants.

For tintinnids, global distribution patterns are, compared to most microzooplankton taxa, relatively well known. On a large scale, they display the most common global biogeographic pattern found among marine species, the bimodal diversity gradient, wherein species richness increases from high to low latitudes with a slight dip near the equator (Chaudhary et al. 2016). Recently, we examined the latitudinal pattern in morphological diversity as well as species diversity in tintinnids along a Northern Hemisphere transect from the Sea of Japan/East Sea to the High Arctic sampled in late summer of 2012, the year of record low sea ice (Dolan et al. 2016). Both species richness and morphological diversity (as numbers of distinct oral diameters) declined dramatically to low levels in the High Arctic. We found just a few species in the High Arctic of very different morphologies suggesting a marked lack of ecological redundancy in the tintinnid assemblage. However, our sampling, as with all oceanographic campaigns and indeed any sampling, yielded only snapshots of certain locales and with generally large detection limits hence, the generality of our finding for tintinnids in the Arctic Ocean overall required confirmation. Diversity gradients within the Arctic have been found previously for terrestrial plants (e.g. Jedrzejek et al. 2012) terrestrial vertebrates (Glig et al. 2012) and aquatic insects (Scott et al. 2011).

To our knowledge there have been no studies of latitudinal gradients in diversity with regard to planktonic organisms of the Arctic. Following Longhurst (1998), the Arctic Ocean

along with its marginal seas comprises the Boreal Polar Province. The latitudinal gradients (roughly) characterising the province between 65°N and the High Arctic, potentially influencing species richness, include water temperature, temporal and areal extent of sea ice, water column depth as well as areal extent of the Arctic Sea (Longhurst 1998).

It has been noted that Arctic data is unusually scattered, often in obscure journals, and generally not compiled (Wassmann 2011). Previous reviews of tintinnid species distributions (i.e., Pierce & Turner 1993; Dolan & Pierce 2013), with regard to the Arctic, relied on the few easily available Arctic records and concerned only the genus-level. Furthermore, the analyses pre-dated several recent relatively large studies (i.e., Monti et al. 2013; Dolan et al. 2014; Feng et al. 2014; Matsuno et al. 2014; Li et al. 2016). To our knowledge, there is no inventory or checklist of tintinnid ciliates in the Arctic. It would be fair to say that tintinnid ciliates have been largely neglected. In recent large synthesis reports concerning Arctic Biodiversity (e.g., Meltotfe 2013; Jorgensen et al. 2016) and reviews of microbes in the Arctic (Dickson et al. 2016) tintinnid ciliates are not mentioned.

To address questions of the magnitude of tintinnid species and morphological diversity in the Arctic and the possible existence of geographic patterns it was necessary to construct *de novo* a database of species records for the Arctic. We combed the published literature for records of the presence of species in locations above 65°N. To published records, we added our own previously unpublished data from analysis of samples obtained in 2010, 2013, 2014 and 2015 from a large suite of stations in the Chukchi Sea. The questions we sought to address were 1) What is the known taxonomic diversity of tintinnids in the Arctic Seas, 2) What is the morphological diversity, in terms of distinct sizes of oral openings, of the tintinnids found in the Arctic Seas, and 3) Is there a latitudinal diversity gradient within the Arctic Sea? While the database was assembled to answer these questions, it may also serve as baseline data, often lacking and needed to detect possible distributional shifts (Jorgensen et al. 2016), as well as providing a checklist of known Arctic morpho-species against which emerging molecular data (e.g., Lovejoy 2014; Pédros-Alio et al 2015; Stecher et al. 2016) may eventually be cross-referenced.

Methods

Google Scholar (<https://scholar.google.com/>) was used to identify documents containing the name of a known Arctic tintinnid genus (i.e., *Acanthostomella*, *Leprotintinnus*, *Parafavella*, *Ptychocylis*). Searches were run also for documents containing the term “arctic” as well the word “tintinnid” or “tintinnids” or “tintinnida”. Articles citing documents containing a species

record and the reference list of the articles themselves were screened by title. For some publications that reported pooled data, e.g., species found in sets of stations, the authors were contacted with a request for their data on species by sampling location (see Acknowledgements for the authors who generously supplied data). Reports from the historic literature were obtained by searching the Biodiversity Heritage Library (<http://www.biodiversitylibrary.org/>) for documents containing mentions of known Arctic species. It should be noted that sampling among the studies varied widely from coarse plankton nets to whole water examination. For many records tintinnid species were noted as 'by-catch' as the investigations were focused primarily on phytoplankton or metazoan zooplankton. Consequently no attempt was made to analyze data in terms of species per investigator publication. A species record was logged only when a binomial species name was given and a location was given, or could be determined, with a sample date (at least the year). Data were entered into a spreadsheet containing 8 columns by which records may be sorted: species, latitude, longitude, locality, reference, sample date, note (station or sample number), and system or sea (i.e., Baffin Bay, Barents Sea, Beaufort Sea, Canadian Archipelago, Chukchi Sea, Greenland Sea, Kara Sea, Laptev Sea, Norwegian Sea, White Sea). We have likely missed some of the Russian language literature not repeatedly cited. However, we have no reason to believe that such additional data would materially effect our conclusions.

We included previously unpublished data from cruises in the Chukchi Sea area in 2010, 2013, 2014, and 2015. Sampling and sampling analysis followed methods described in detail in Dolan et al. 2013, 2016. Briefly, for most stations a 20 μ m mesh plankton net was towed from approximately 100 m to the surface. Net material was fixed with either Acid Lugol's solution or Bouin's fixative and in the lab sample aliquots were examined with an inverted microscope. Our previously unpublished species records were entered under "reference" as ARA 01 Cruise for 2010, ARA 03 Cruise for 2013, ARA 05 Cruise for 2014, and ARA 06 Cruise for 2015 (see the Online Resource data file containing all species records).

To examine morphological characteristics of the Arctic species pool, each species was assigned lorica dimensions. We used the average of the range of lorica dimensions for the species given in the text descriptions by Kofoed & Campbell 1929 as many investigators relied on the monograph for species identification. For species depicted in Kofoed & Campbell 1929 but unaccompanied by a text description, we assigned dimensions of forms with text descriptions depicted in the plates as having very similar dimensions. For species described after 1929 the dimensions given in the original descriptions were assigned. We realise that dimensions and scales of depictions in Kofoed & Campbell 1929 contain errors compared to the

preferred authority of the original description or re-descriptions (e.g., Dolan 2016b; Santoferrara et al. 2016). However, the dimensions given in the text, or implied in the illustrations of Kofoed and Campbell, are likely to correspond to those of the organism observed by individual investigators compared to a more authoritative description. The final species list was cross-checked against GenBank to identify taxa for which sequence data has been deposited as of July 25th 2016.

We also distinguished some possibly doubtful species identities. Some of the tintinnid species commonly found in sub-Arctic and Arctic waters have long been suspected to form variable loricas with the morphological variants given undue status as distinct species. These suspected polymorphs are species of *Acanthostomella* (our own observations), *Parafavella* (Schulz & Wulff 1927; Burkovsky 1973; Cardinal et al. 1997; Davis 1978) and *Ptychocylis* (Davis 1981).

Results

Qualitative and Quantitative Characteristics of the Species Records

We extracted data from 39 publications published between 1900 and 2015 containing records of species from locations from 65° to 82°N. Including our own new data for the Chukchi Sea, the database contains 1427 records of 89 species from 414 locations sampled from 1885 (Gran 1900) to 2015 (this report). The vast majority of records are from single time point sampling of unique localities from oceanographic cruises in summer months. Few reports gave time series data and only 2 had data from winter sampling, which interestingly yielded species similar to those found in summer sampling at the same site (Gaarder 1938; Tibbs 1967).

Table 1 summarises the contents of the database giving the list of species, the number of records by “Sea”, for each species, as well as the first year the species was reported from the Arctic. Taxa for which one or more nucleotide sequences have been deposited in GenBank are shown in bold. Of the 19 tintinnid genera, 13 are represented in GenBank and of the 89 tintinnid species GenBank contains sequence data for 23 species. The complete database of species records is supplied as a Online Resource data file. Figure 1 shows the locations of all sampling points revealing large discrepancies in the geographic coverage of the data. Notably, there are no data for the Eastern Siberian Sea and there is very little data available for the Laptev Sea as well as the Canadian Archipelago.

The 4 most often reported species, in order of the number of records, were *Ptychocylis obtusa*, *Parafavella denticulata*, *Acanthostomella norvegica* and *Salpingella acuminata*, species of quite distinct morphologies (Fig. 2). The four species accounted for 45% of species

records (Fig. 3). These species were also the most widely distributed in terms of the number of seas from which they were reported (Table 1) and their apparent latitudinal range (Fig. 3). Remarkably, no species has been reported from all of the Arctic Seas. The number of potentially questionable records, the 'oncers' that is species recorded but once, represent 20 out of the 89 species. The suspected "polymorphs", possibly morphological variants of the most common species of *Acanthostomella*, *Parafavella* and *Pytchocylis*, represent 21 of the 89 species. Thus, even after subtracting potentially dubious records, the Arctic tintinnid fauna is substantial, consisting of 48 species in 16 genera. The morphological diversity, in terms of numbers of species in different size-classes of lorica oral diameter, is large as well, considering the entire species pool recorded, or the species pool remaining after removing 'oncers' and 'polymorphs' (Fig. 4). Most size-classes contained several species. Notably, the size-classes containing, or neighbouring, the 4 most commonly recorded species were the size classes containing the largest numbers of species (Fig. 4).

Latitudinal and Historical Patterns

We pooled species records in bands of 2° latitude and plotted species richness against latitude. The resulting graph (Fig. 5) showed large differences in species richness among bands of latitude with perhaps a linear decline from 74° to 82°, but no overall gradient of species diversity with latitude. The heterogeneity in species richness among bands led us to examine the relationship between the number of sites sampled within a 2° band of latitude and the number of species recorded as occurring in the band. We found a significant positive relationship between species richness of a 2° latitudinal band and the number of sites sampled in the band (Fig. 5). The lack of a latitudinal gradient and the apparent positive relationship of species richness with sampling effort held when excluding "polymorphs", the species recorded which are suspected of being morphological variants of another species recorded from the same latitudinal band. Examination of data concerning the Chukchi Sea, arguably the most sampled and with a large latitudinal gradient revealed the same patterns (data not shown). The Arctic Seas vary considerably in basic characteristics of latitudinal range, area, average depth, importance of freshwater input, etc. (see the online resource figure Arctic Ocean bathymetric map). However, plotting the apparent species richness of the Arctic Seas as a function of sampling effort revealed the same positive relationship (Fig. 6). The apparent strong effect of sampling effort on apparent species richness led us to examine the large-scale (over the past 120 years) historical trends.

We found strikingly similar patterns plotting accumulated sites sampled with time from 1895 to 2015 and accumulated apparent species richness from 1985 to 2015 (Fig 6.). A simple linear correlation characterised the historical data cumulative sampling effort (log cumulative sites) and (log) cumulative number of species recorded (inset graph in Fig. 6). The relationship resembles a typical species-area curve of a census of contiguous habitats (Preston 1962). It is important to note that temporal increases in the Arctic species inventory do not represent increases in the number of species new to science but rather new to the Arctic. There was no indication of a plateau in the relationship suggesting that further increasing sampling effort in the Arctic, that is simply by continuing to sample, will likely yield increases in the number of tintinnid species known from Arctic waters.

Discussion

Qualitative and Quantitative Characteristics of the Species Records

To put the magnitude of tintinnid species diversity in the Arctic, 89 species, in perspective, one can compare the number with total known diversity of living forms. The standard taxonomic monograph of Kofoed & Campbell (1929) lists and illustrates about 750 species. In a more recent compendium, the list extends to over 900 species (Zhang et al. 2012). Although many catalogued “species” are likely to be synonyms (e.g. Dolan 2016b), the forms reported from the Arctic Seas appear to represent a considerable fraction of described tintinnid species not restricted to warm waters, which number about 500, based on genus characterisation (Pierce & Turner 1993; Dolan & Pierce 2013). A majority of the Arctic genera are represented in GenBank with at least one nucleotide sequence deposited (Table 1). Of those genera without any GenBank record, most have been recorded only a few times. However, of 2 the most abundant and long-known taxa, *Acanthostomella* and *Parafavella*, are completely absent from GenBank.

Arctic species diversity appears to be lower than that reported for the Southern Ocean, reviewed recently in Dolan et al. 2013. The records from the Southern Ocean are roughly comparable in quantity (2000 records from 402 locations) but contain 192 species. However, it is important to recall that the Southern Ocean is a much larger system as its northern border is usually taken as 40° S. Even using a northern border of 60°S, the surface area of the Southern Ocean is 21,000 km² compared to the Arctic Ocean's 15,000 km² (Eakins & Sharman 2010). Among the 89 species recorded from Arctic Seas, 20 have only been found once from one locality and possibly represent stray species rather than resident forms. In comparison, among the Southern Ocean species, over a third appear to be strays, that is with only one recorded presence (Dolan et al. 2013). While many species have been found in the Arctic, the records

mirror early reports of tintinnid assemblages being overwhelming dominated by just a few species (Gran 1900) as nearly half the records concern only 4 species (Fig. 3). In the size-class of each these frequently recorded species, are several other species of similar mouth size or lorica oral diameter (Fig. 4).

If we consider species of similar mouth size to be ecologically similar, then the pattern of several species in each size-class suggests that there is considerable ecological redundancy overall in the pool of Arctic tintinnid species based on the inventory pooling all historical records. This is in contrast to findings of little ecological redundancy from our recent study which was based on samples from a transect of stations across the western Arctic to 82°N (Dolan et al. 2016). The discrepancy likely reflects the fact the overall species pool described here sums a large number of samplings conducted over a long period of time. However, both the single transect data and the overall pooled data suggest that the High Arctic is apparently species-poor. The High Arctic, characterised by a smaller areal extent and extensive sea ice coverage relative to lower latitude Arctic Seas, may indeed be species-poor. However it is also poorly sampled and actually may harbor more species than known at present.

Interestingly, among the 4 species which dominate species records of the Arctic, 2 are also found in the Antarctic. *Acanthostomella norvegica* appears to be a bi-polar species found in high latitude waters of both hemispheres but absent from tropical, subtropical and temperate systems (Dolan et al. 2013). However, it is rarely abundant in Antarctic waters. The other Arctic dominant found also in Antarctic assemblages is *Salpingella acuminata*. It is an apparently cosmopolitan species found as a dominant form in assemblages from tropical waters (Dolan et al. 2007) to the Chuckchi Sea (Dolan et al. 2014). However, like *A. norvegica* it is not known to be dominant form in Antarctic waters.

Latitudinal and Historical Patterns

Our study began with a desire to confirm or refute the existence of a latitudinal diversity gradient in the Arctic Ocean. Geographic distribution of species records showed no gradient throughout the Arctic Ocean, only a decline from 74° to 82°. While the highest latitudes do harbour a lower number of species, the highest latitudes are also the least sampled. We found a positive relationship between apparent species richness of a latitudinal zone and the number of sites sampled within the zone. This led us to examine the historical trends of species records and sampling effort. Additions to the species inventories closely corresponded with cumulative sampling effort in the crude form of the number of sites sampled. Based on our findings we predict that the species inventory will continue to grow simply as a result of increased

cumulative sampling effort. There appears also to be a lack of baseline data on the distribution and composition of phytoplankton taxa (Daniëls et al. 2013). This considerably complicates diagnosis of the effects of changes in the Arctic for both microzooplankton and phytoplankton because baseline data is necessary to detect shifts (Jorgensen et al. 2016). In contrast, a similar situation does not appear to be the case for metazoan taxa of the zooplankton. Zooplankton diversity has been described as well characterised except for taxa restricted to the bathypelagic layer where increased sampling is expected to reveal species new to science (Kosobokova et al 2010). Changes in circulation in the Arctic are predicted to yield major changes in the composition of metazoan zooplankton (Wassmann et al. 2015) that in contrast to phytoplankton and microzooplankton, may be detectable. In reality, the lack of long time-series data such as that available for Mediterranean zooplankton (e.g. Berline et al. 2012) prevents rigorous examination of temporal changes in the species composition at any one particular site, much less the Arctic Sea overall.

The historical species accumulation curve for tintinnid species recorded from the Arctic is difficult to assess as comparative data for other taxa is unavailable, as far we know. There are data for another locality. A recent study examined increases with time in the species inventory of tintinnid ciliates in the Bay of Villefranche in the NW Mediterranean Sea (Dolan 2016a) and data do exist for the discovery of species new to science as species described with time. Figure 8 shows data retrieved from the WoRMS database for all marine biota catalogued showing accumulated number of species with time from 1880 to 2015 along with the species accumulation curve for Arctic tintinnids from this study and the Bay of Villefranche from Dolan (2016a). Increases in inventory of described marine species worldwide with time can be attributed to sampling effort directed toward novel or previously under-sampled environments (rather than re-examining archived samples) and thus increases in cumulative sampling effort. There is a striking similarity in the slopes reflecting close correspondence in the relative rates of increases over time in the number of described species in marine biota overall and the number of tintinnid species known from sampling in the Arctic as well as a bay in the Mediterranean Sea. We suggest that this indicates that sampling effort in the Arctic has remained roughly proportional to sampling for new marine species world-wide since the beginning of the 20th century. Unfortunately, testing such a hypothesis would be quite challenging and well beyond the scope of the present study.

Conclusion

We found large discrepancies in geographic coverage of tintinnid species records. Many areas likely to experience large changes in sea ice coverage are under-sampled. While we found no clear trend of species richness with latitude throughout the Arctic, there was a significant positive relationship between species richness reported and the number of sites sampled in a 2° band, suggesting a sampling effect. Examination of temporal trends in sampling effort and changes in the species inventory revealed a near-linear relationship in cumulative numbers of species recorded and sites sampled with time, and a lack of a plateau in the species accumulation trend. Species records are highly dominated by 4 species, accounting for 45% of the records: *Acanthostomella norvegica*, *Parafavella denticulata*, *Ptychocylis obtusa* and *Salpingella acuminata*, all of which, except *S. acuminata*, have long been suspected to be morphologically variable, with different morphotypes given undue species status. Pooling all reported species of *Acanthostomella*, *Parafavella* and *Ptychocylis* yielded little qualitative differences but considerable quantitative differences. The list of Arctic tintinnid ciliate species will likely continue to grow with new sampling, regardless of changes in the Arctic Seas. A perhaps more useful harbinger of change in the microzooplankton community of the Arctic would be shifts in the occurrences and ranges of the 4 historically dominant species or replacement of one the dominants by another species.

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Table 1. List of species recorded from Arctic Seas. Number of records for the Greenland Sea (GS), Baffin Bay (BB), Canadian Archipelago (CA), Beaufort Sea (Be S) Chukchi Sea (CS), Laptev Sea (LS), Kara Sea (KS), Barents Sea (BS), White Sea (WS) and Norwegian Sea (NS). References given in Table 2. Taxa in bold denote sequence data present in GenBank as of July 25 2016.

Species	G S	BB	CA	Be S	C S	LS	KS	Ba S	WS	NS	references	First Year
<i>Acanthostomella conicoides</i>					5						27,35	2010
<i>Acanthostomella gracilis</i>		1	3								20,34	1988
<i>Acanthostomella norvegica</i>	35	2			63		12	16		25	1,2,3,4,5,11,12,13,15,16,17,18,19,23,24,26,27,28,29,35,36,37,3	1899
<i>Bursaopsis vitrea</i>					1						10	2012
<i>Canthariella pyramidata</i>					2						35	2010
<i>Codonellopsis frigida</i>					10						14,22,35	2010
<i>Codonellopsis lagunela</i>										2	28,30	1945
<i>Codonellopsis morchella</i>					1						35	2010
<i>Codonellopsis ovata</i>									1		3	1970
<i>Codonellopsis pusilla</i>									1		3	1970
<i>Codonellopsis schabi</i>					1						35	2010
<i>Eutintinnus apertus</i>					1						9	2011
<i>Eutintinnus tenuis</i>	2										13	1994
<i>Favella azorica</i>					10						35	2010
<i>Helicostomella fusiformis</i>										1	12	1930
<i>Helicostomella subulata</i>					2				2	6	12,22,3,24,30,14	1898
<i>Leprotintinnus bottnicus</i>						1	1		1		3,33	1927
<i>Leprotintinnus pellucidus</i>	2	4		6	24		12	14		8	1,2,3,4,9,11,12,14,17,19,22,30,32,36,39	1907
<i>Metacylis vitreoides</i>	1				3		3				1,3,22,32	1907
<i>Ormosella trachelium</i>					1						35	2010
<i>Parafavella acuta</i>							7				1,3	1925
<i>Parafavella cylindrica</i>								17		1	1,8,36	1921
<i>Parafavella denticulata</i>	76	9	1		1	1	9	10	2	20	1,2,3,24,8,30,29,12,13,15,23,16,17,18,19,27,25,32,33,34,36,37,	1895
<i>Parafavella dilatata</i>										1	8	1975
<i>Parafavella edentata</i>	3									2	5,11,12,30	1898
<i>Parafavella elegans</i>	1	1	9	2	2					1	7,14,15,26,34	1975
<i>Parafavella faceta</i>					3						14	2014
<i>Parafavella gigantea</i>	24	2		8	1		18	5	13		22,6,8,11,29,26,15,16,17,18,31,21	1907
<i>Parafavella hemifusus</i>								3			1	1925
<i>Parafavella jorgensi</i>					15						14,35	2010
<i>Parafavella media</i>	3										5	1898
<i>Parafavella obtusa</i>									1		3	1970
<i>Parafavella obtusangula</i>	1									1	8,26	1975
<i>Parafavella parumdentata</i>					9		2		9	1	8,10,22,39	1975
<i>Parafavella promissa</i>					1						14	2014
<i>Parafavella robusta</i>	14						9			1	1,9,18	1906
<i>Parafavella rotundata</i>							6			1	1,30	1925
<i>Parafavella subrotundata</i>					1		10				1,6,22	1921
<i>Parafavella tenuis</i>							2				1	1925
<i>Parafavella ventricosa</i>					2						14	2014
<i>Ptychocyliis acuta</i>	1	1						3		1	7,29,26,15	1975
<i>Ptychocyliis arctica</i>					1				1	4	3,4,7,12,30	1926
<i>Ptychocyliis cylindrica</i>										3	12,30	1930
<i>Ptychocyliis drygalski</i>	9			10							16,21	1935
<i>Ptychocyliis minor</i>										1	7	1975
<i>Ptychocyliis obtusa</i>	71	5	9	1	116		20	9	13	13	22,9,10,1,2,3,4,5,7,11,12,30,29,13,14,15,16,17,18,25,32,35,36,3	1898
<i>Ptychocyliis urnula</i>	31				1		7			8	4,12,16,30,36,38	1921
<i>Salpingacantha perca</i>					1						35	2010
<i>Salpingacantha simplex</i>					14						22	2010
<i>Salpingella acuminata</i>	18			23	66		2	2	1	11	2,3,4,9,10,12,16,21,22,28,30,36,37,38	1921
<i>Salpingella curta</i>										2	17	2008
<i>Salpingella faurei</i>					65						9,10,22	2010
<i>Salpingella secata</i>							3	3			29	1925
<i>Salpingacantha unguiculata</i>								1			29	1993
<i>Schmidingerella serrata</i>										5	12,30,38	1930
<i>Schmidingerella</i>												
<i>taraikaensis</i>									1		3	1970
<i>Steenstrupiella robusta</i>									1		3	1970

Stenosemella nivalis					8		1			1	14,17,35,36	1921
<i>Stenosemella olive</i>								1	1	2	3,29,12	1930
Stenosemella steinii	2										15	1988
Stenosemella ventricosa					4		9			1	1,22,23,35,36	1921
Tintinnidium mucicola					17						35	2010
<i>Tintinnopsis acuminata</i>					17						14,22	2010
<i>Tintinnopsis amphora</i>								1			40	2002
<i>Tintinnopsis angusta</i>								5			40	2002
Tintinnopsis baltica					5						14,22	2014
Tintinnopsis beroidea	2	1			6		9	5	1	9	1,2,3,4,9,12,15,30,36,37,38	1899
<i>Tintinnopsis campanula</i>									1	5	3,12,30,38	1898
Tintinnopsis elongata								7			40	2002
Tintinnopsis fimbriata					1	1	1		1		3,22,33	1927
<i>Tintinnopsis japonica</i>					2						3,14	2014
<i>Tintinnopsis karajacensis</i>	1				5		10				4,22,32,36	1907
<i>Tintinnopsis kofoidi</i>					1						14	2014
Tintinnopsis lata					9		3				1,22	1925
Tintinnopsis major										1	12	1930
<i>Tintinnopsis mayeri</i>					4						14	2014
<i>Tintinnopsis meureri</i>							3				1	1925
Tintinnopsis nana					1				1		3,22	1970
<i>Tintinnopsis nitida</i>							4				1,36	1921
Tintinnopsis parvula	1				1				1	3	3,4,11,12,30	1930
<i>Tintinnopsis plagiotoma</i>								2			40	2002
Tintinnopsis rapa					16						3,9,14,22	2011
Tintinnopsis subacuta					2						9	2012
Tintinnopsis tubulosa							4		1		3,36	1921
<i>Tintinnopsis turbo</i>					2			1		2	22,2	1988
<i>Tintinnopsis undella</i>						1	1				3,33	1927
<i>Tintinnopsis urnula</i>					2					2	9,12	1930
<i>Tintinnopsis vasculum</i>								14			40	2002
Species	G	BB	CA	Be S	C	LS	KS	Ba S	WS	NS	references	First Year

Table 2. Sources of Arctic records of tintinnids, References numbers as noted in Table 1.

Reference #	Citation (for full reference see “References”)
1	Bernstein 1931
2	Boltovskoy et al. 1991
3	Burkovskii 1976
4	Bursa 1963
5	Cleve 1899
7	Davis 1981
8	Davis 1978
9	Dolan et al. 2014
10	Dolan et al. 2016
11	Feng 2014
12	Gaarder 1938
13	Levinsen et al. 1999
14	Li et al. 2016
15	Lovejoy et al. 2002
16	Meschkat 1939
17	Monti et al. 2013
18	Ostenfeld & Paulsen 1911
19	Paranjape 1987
20	Smayda, 1958
21	Tibbs 1967
22	Present Study: Chukchi Sea data 2010, 2013, 2014, 2015
23	Lutter et al 1989
24	Chernova & Primakov 2011
25	Yokoi et al 2016
26	Kubiszyn et al 2014
27	Piontek et al 2014
28	Braarud et al 1958
29	Jensen & Hansen 2000
30	Heimdal 1974
31	Ratkova et al. 1994
32	Ostenfeld 1910
33	Burkovskii et al. 1974.
34	McLaughlin et al 2009
35	Matsuno et al 2014
36	Rossolimo 1927
37	Boltovskoy et al 1995
38	Gran 1900
39	Yang et al. 2004
40	Lee et al. 2003

Figure 1. Map of the locations sampled from 1885 to 2015 from which one or more tintinnid species was reported. Note that there does not appear to any historical bias with regard to a dominance of nearshore compared to open water sampling.

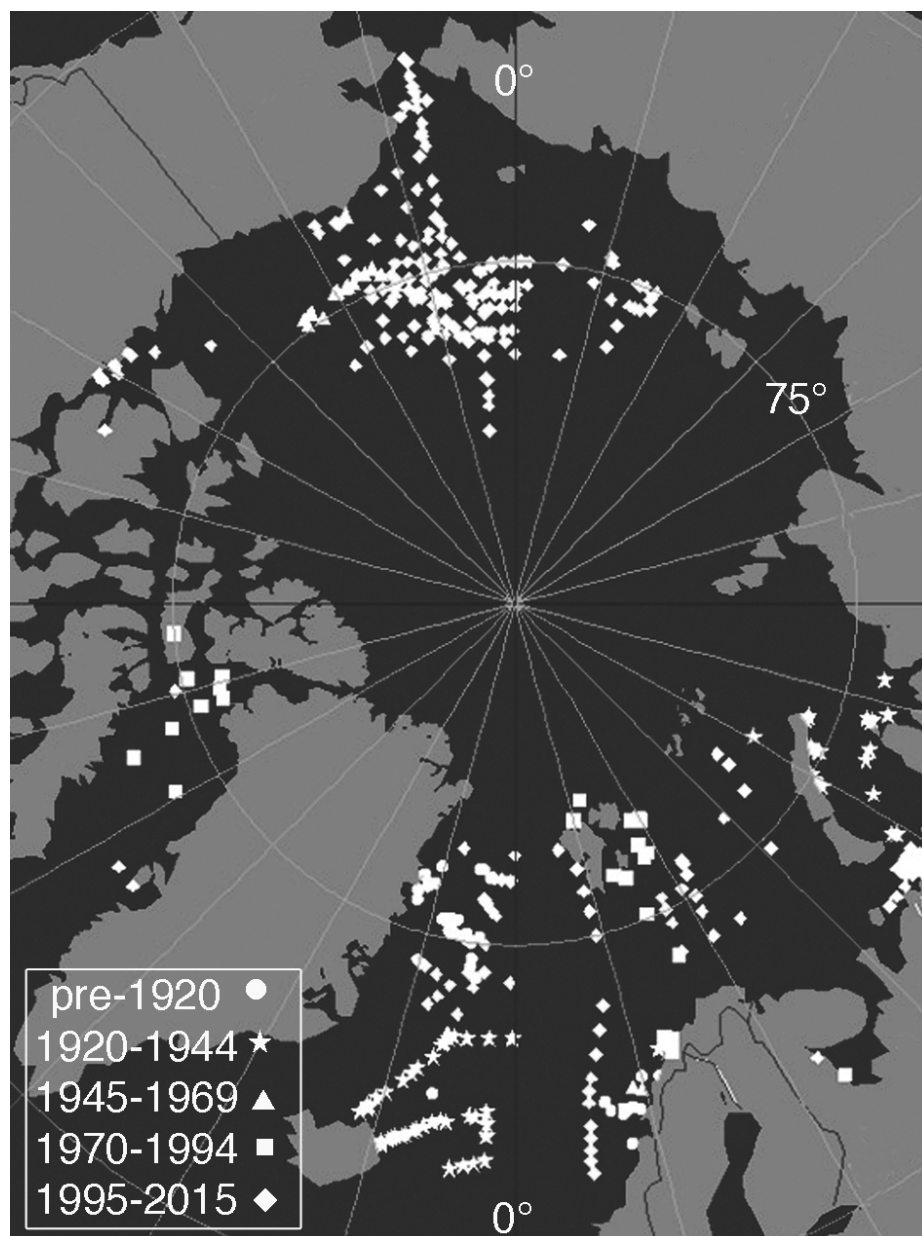


Figure 2. Micrographs of the 4 most commonly reported tintinnids from Arctic waters: A. *Salpingella acuminata*, B. *Acanthostomella norvegica*, C., *Ptychocylis obtusa*, and. D. *Parafavella denticulata*. All Lugol's-fixed specimens except for D, a Bouin's-fixed cell.

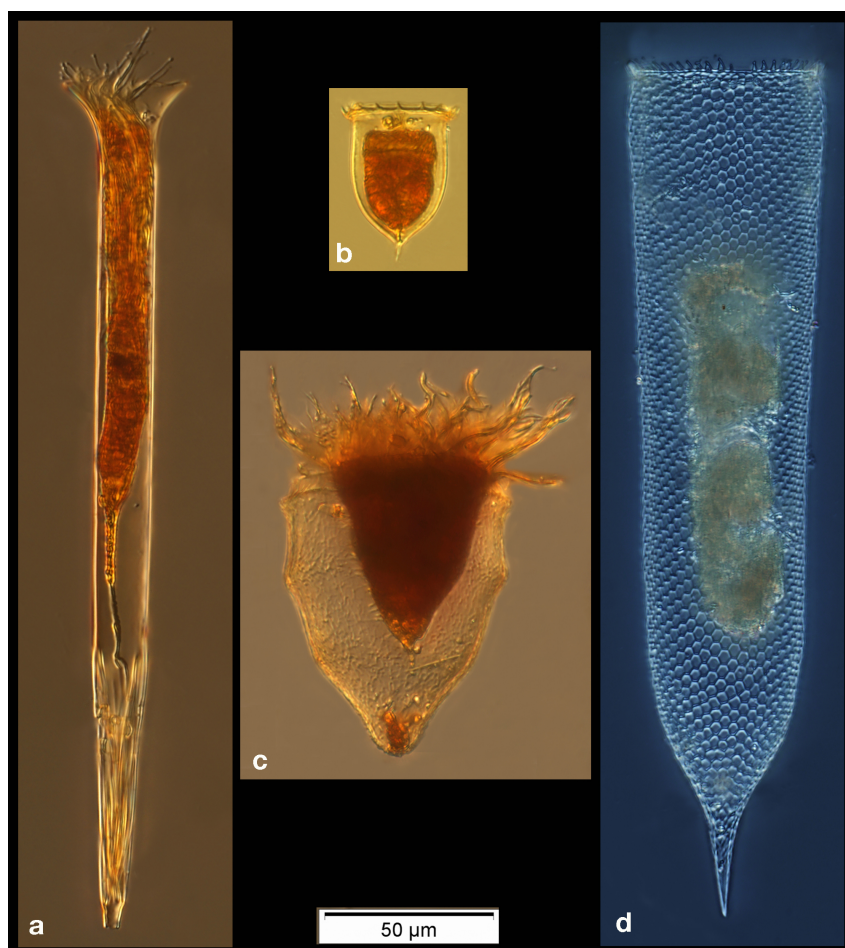


Figure 3. Plot of occurrence rank (left panel) and apparent latitudinal range vs. the number of records for the species (right panel). Remarkably, 4 species account for a large fraction of the species records and these most frequently recorded forms are also those with the largest apparent latitudinal range (species with but a single record were excluded). Note that while a species recorded but once will by definition have no latitudinal range, species with multiple records may or may be recorded across a range of latitudes.

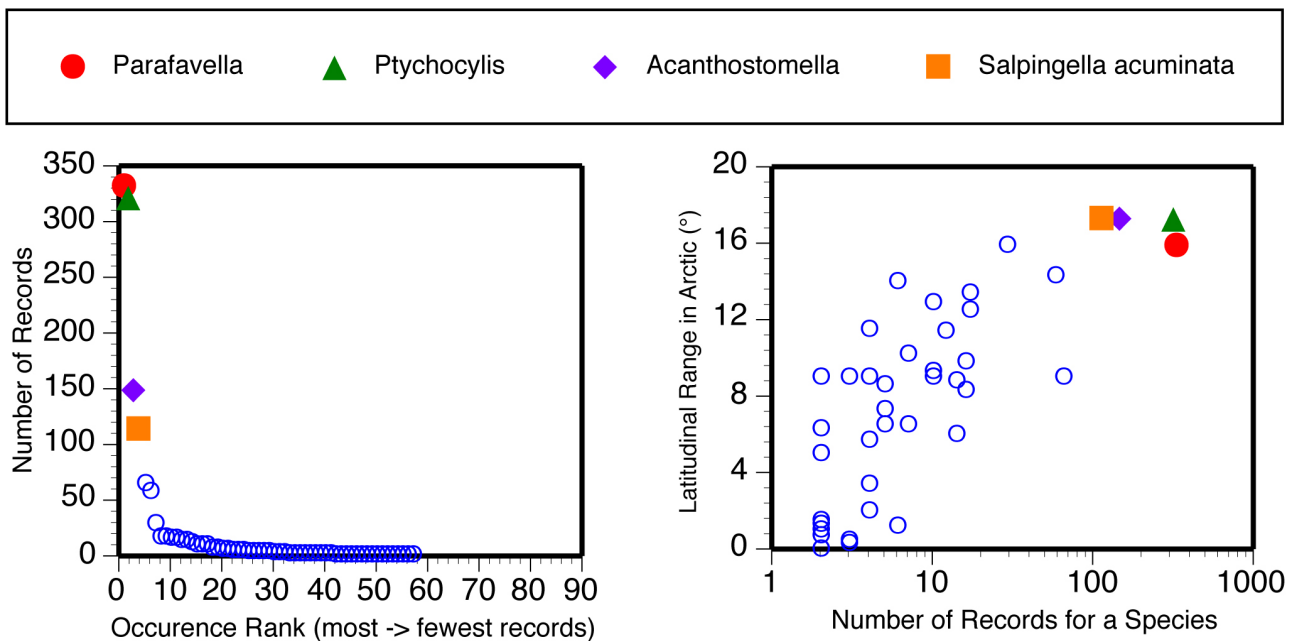


Figure 4. Morphological characteristics of tintinnid species found in the Arctic, frequency distribution of lorica oral diameters, or mouth sizes. Species were binned in size classes of 4 μm . Data shown for all reported species (all Arctic Species) and the reduced pool of species remaining after removing species recorded only once and suspected morphological variants of a main species (Arctic Species w/o Oncers & Polymorphs). The size classes containing the most frequently recorded species are shown with an asterisk. Note that x-axis breaks used to include the large size classes omitting empty intermediate size-categories.

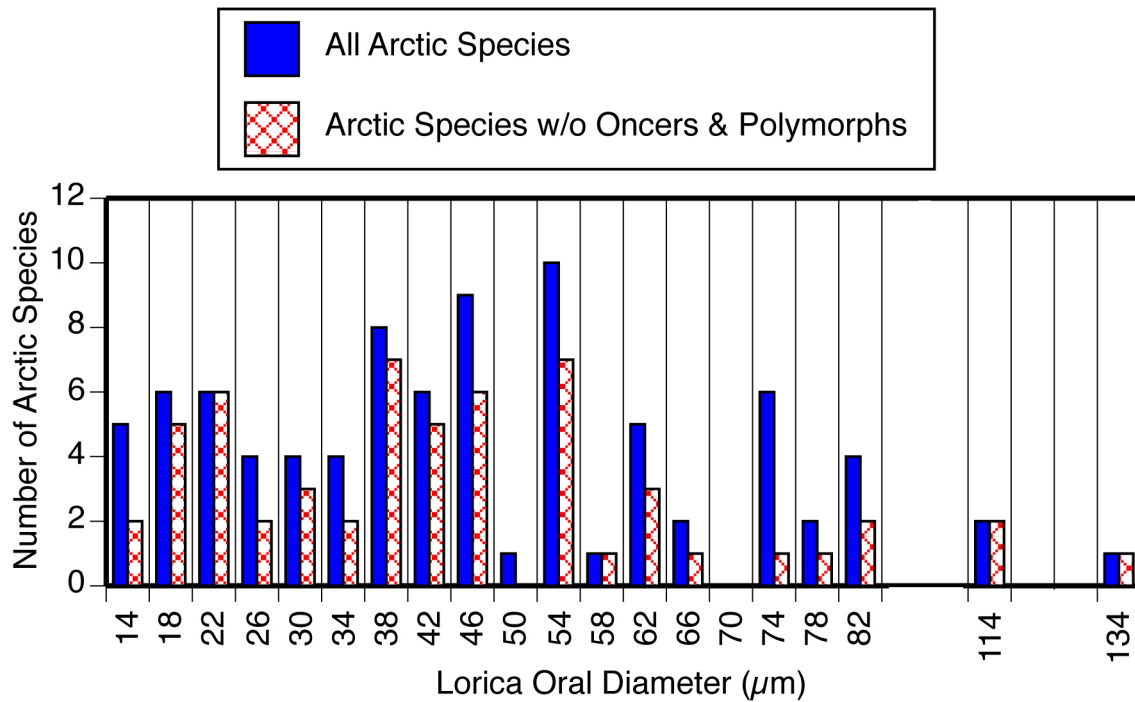


Figure 5. Species richness as a function of latitude, pooling data in bands of 2° latitude (left panel) and species richness as a function of the number of sites sampled within the 2° band of latitude (right panel). For both plots total numbers of species as recorded in publications are shown as well numbers of species without suspected polymorphs, morphological variants of the other species also recorded in the band of latitude. Note the lack of consistent latitudinal gradient in species richness throughout the Arctic and the correlation of species richness within a band of latitude with the number of sites sampled within the band. For total species (# spp) $r = 0.70$; for species excluding records of suspected morphological variants (w/o polymorphs) $r = 0.63$.

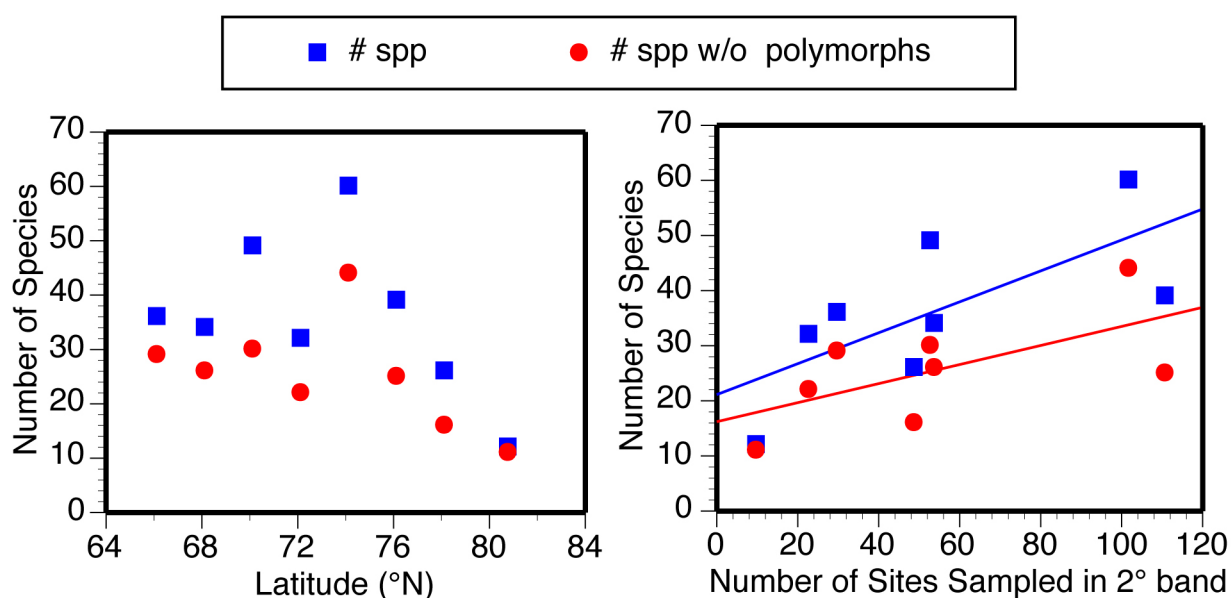


Figure 6. Species richness of the Arctic Seas as a function of the number sites sampled within the Sea. The Seas differ considerably in basic characteristics of area, latitudinal range, average depth, freshwater input, ice cover, etc., (see Online Fig. 1). Note the positive relationship of species richness with sampling effort, $n = 10$, $r^2 = 0.66$, $p < 0.05$.

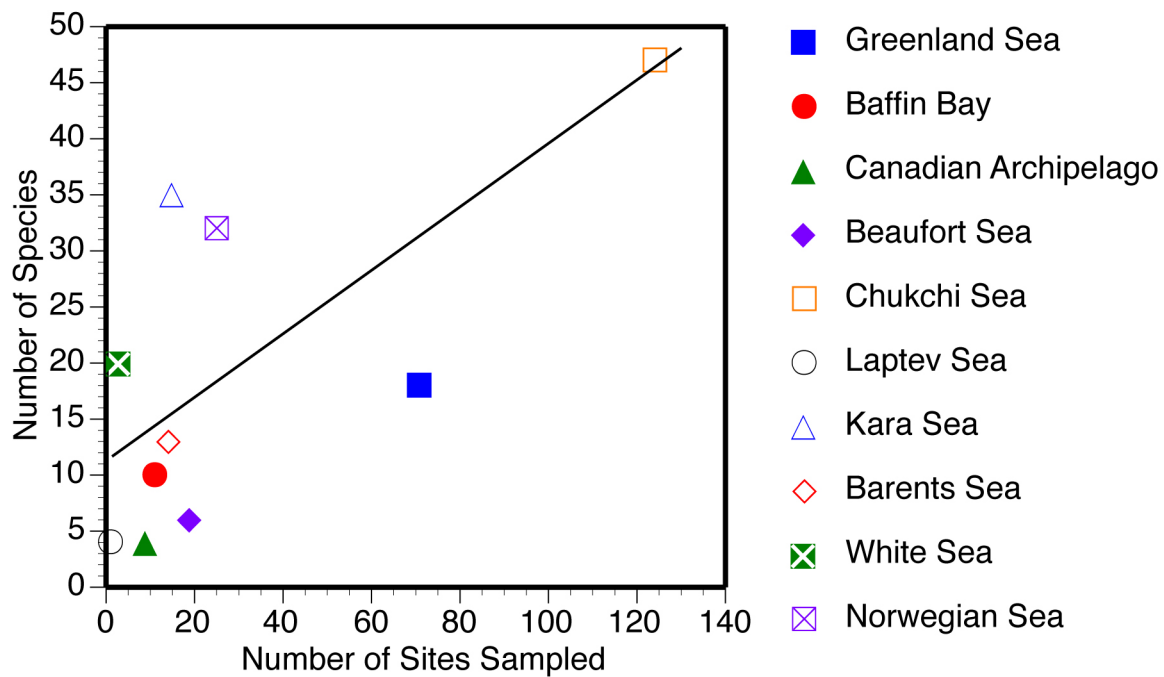


Figure 7. Temporal changes in the cumulative number of sites sampled and species recorded (all reported and excluding suspected morphological variants of other species) in the Arctic from 1895 to 2015. Note the close correspondence in temporal trends. A simple linear correlation of log cumulative number of sites and log cumulative number of species is shown in the inset graph resembling the Preston species-area curve for contiguous habit sampling (Preston 1962).

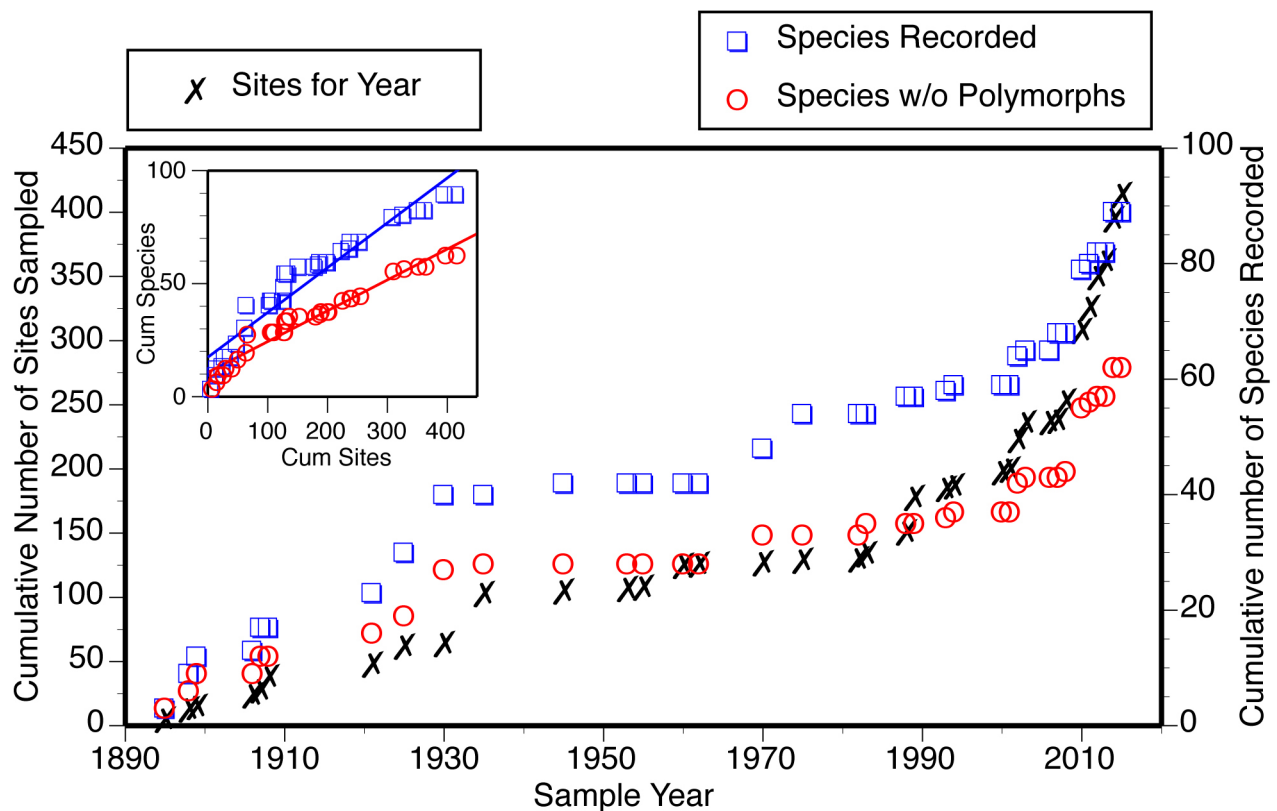
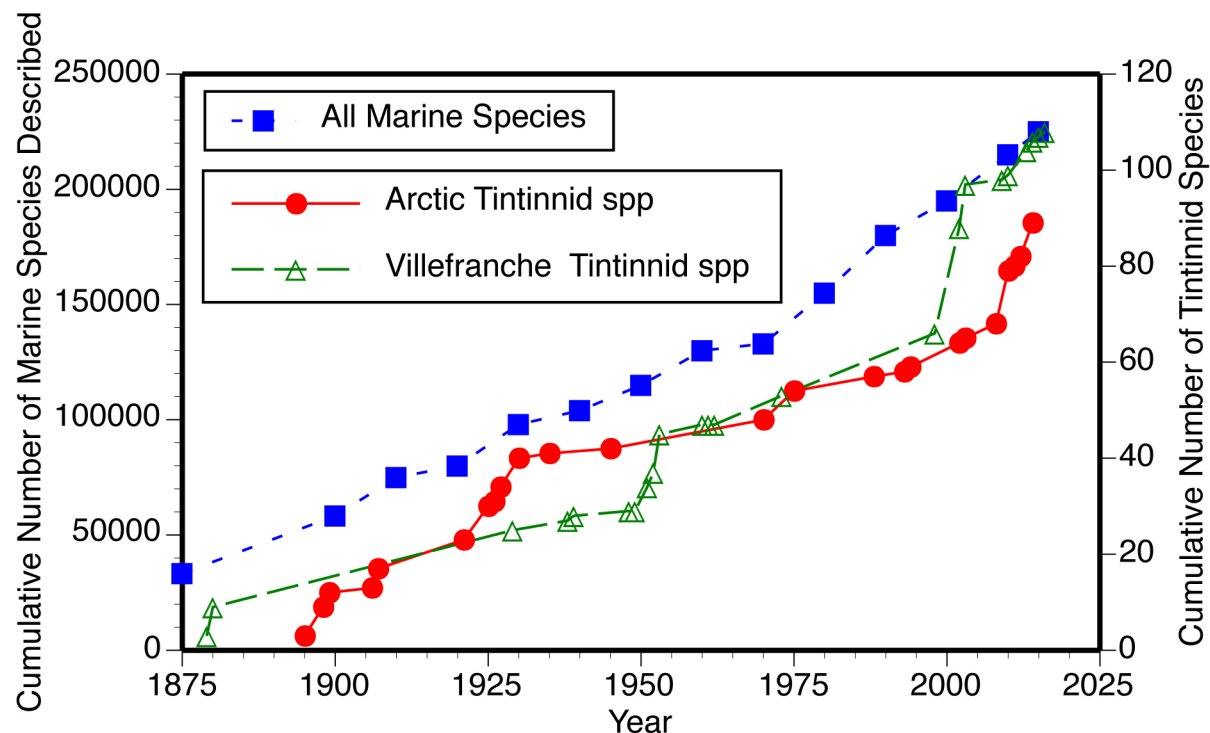
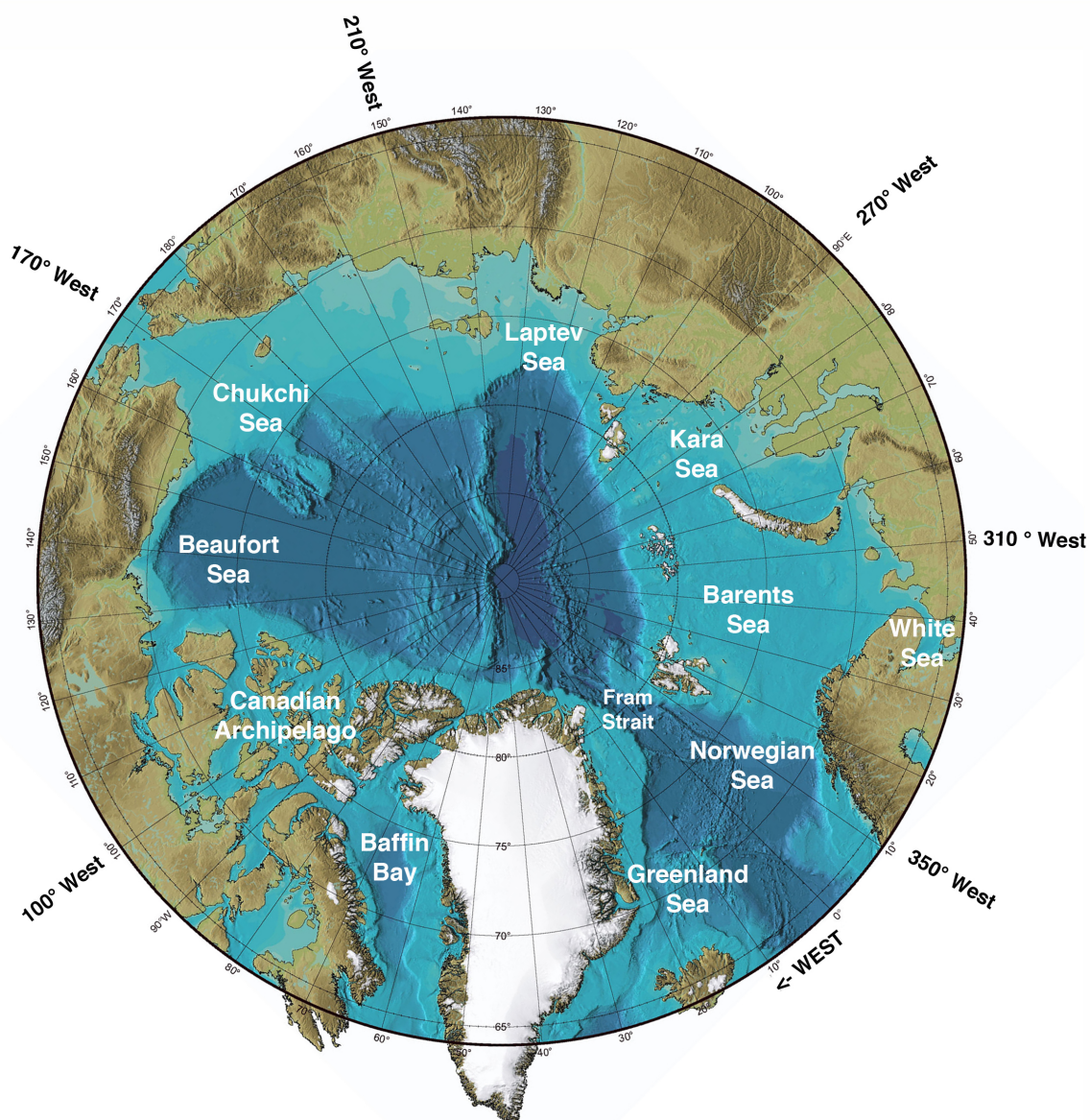


Figure 8. The cumulative number of marine species described from 1875 to 2015 and cumulative number tintinnid species reported from the Arctic from 1885 to 2015 as shown in Fig. 6. Data on the temporal changes in the species inventory of the Bay of Villefranche (NW Mediterranean Sea) are from Dolan (2016a). Data for total marine species (benthic, nekton, planktonic, parasitic, free-living, etc.) described from WoRMS Editorial Board (2016).



Online Resource Figure 1. Bathymetric Map of Arctic Seas with seas sampled labelled and 'Degrees West' indicated as given in Online Resource Data File.



Base map from: Jakobsson, M., L. A. Mayer, B. Coakley, J. A. Dowdeswell, S. Forbes, B. Fridman, H. Hodnesdal, R. Noormets, R. Pedersen, M. Rebesco, H.-W. Schenke, Y. Zarayskaya A, D. Accettella, A. Armstrong, R. M. Anderson, P. Bienhoff, A. Camerlenghi, I. Church, M. Edwards, J. V. Gardner, J. K. Hall, B. Hell, O. B. Hestvik, Y. Kristoffersen, C. Marcussen, R. Mohammad, D. Mosher, S. V. Nghiem, M. T. Pedrosa, P. G. Travaglini, and P. Weatherall. 2012. The International Bathymetric Chart of the Arctic Ocean (IBCAO) Version 3.0, *Geophysical Research Letters*, doi: 10.1029/2012GL052219.