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Diversity and abundance of pteropods and heteropods along a latitudinal gradient across the Atlantic Ocean

Alice K. Burridge, Erica Goetze, Deborah Wall-Palmer, Serena L. Le Double, Jef Huisman, Katja T.C.A. Peijnenburg

Naturalis Biodiversity Center, P.O. Box 9517, 2300 RA Leiden, The Netherlands
Institute for Biodiversity and Ecosystem Dynamics (IBED), University of Amsterdam, P.O. Box 94248, 1090 GE Amsterdam, The Netherlands
Department of Oceanography, University of Hawai’i at Mānoa, 1000 Pope Road, Honolulu, HI 96822, USA
School of Geography, Earth and Environmental Sciences, Plymouth University, Drake Circus, Plymouth, Devon PL4 8AA, UK

Abstract

Shelled pteropods and heteropods are two independent groups of holoplanktonic gastropods that are potentially good indicators of the effects of ocean acidification. Although insight into their ecology and biogeography is important for predicting species-specific sensitivities to ocean change, the species abundances and biogeographical distributions of pteropods and heteropods are still poorly known. Here, we examined abundance and distribution patterns of pteropods (euthecosomes, pseudothecosomes, gymnosome) and heteropods at 31 stations along a transect from 46°N to 46°S across the open waters of the Atlantic Ocean (Atlantic Meridional Transect cruise AMT24). We collected a total of 7312 pteropod specimens belonging to at least 31 species. Pteropod abundances were low north of 40°N with <15 individuals per 1000 m^3, varied between 100 and 2000 ind./1000 m^3 between 30°N and 40°S, and reached >4000 ind./1000 m^3 just south of 40°S. This accounted for an estimated biomass of 3.2 mg m^-3 south of 40°S and an average of 0.49 mg m^-3 along the entire transect. Species richness of pteropods was highest in the stratified (sub)tropical waters between 30°N and 30°S, with a maximum of 15 species per station. The biogeographical distribution of pteropod assemblages inferred by cluster analysis was largely congruent with the distribution of Longhurst’s biogeochemical provinces. Some pteropod species distributions were limited to particular oceanographic provinces, for example, subtropical gyres (e.g. Syllopa subula) or warm equatorial waters (e.g. Creseis virgula). Other species showed much broader distributions between ~35°N and ~35°S (e.g. Limacina helicina and Helicinae infusus). We collected 1812 heteropod specimens belonging to 18 species. Highest heteropod abundances and species richness were found between 30°N and 20°S, with up to ~700 ind./1000 m^3 and a maximum of 14 species per station. Heteropods were not restricted to tropical and subtropical waters, however, as some taxa were also relatively abundant in subantarctic waters. Given the variation in distribution patterns among pteropod and heteropod species, it is likely that species will differ in their response to ocean changes.

1. Introduction

Shelled pteropods and heteropods are holoplanktonic gastropods with aragonite shells that are vulnerable to the effects of ocean acidification (e.g., Bednaršek and Ohman, 2015; Gattuso et al., 2015; Wall-Palmer et al., 2016a). They are common components of the marine zooplankton and have highly diverse evolutionary histories, life strategies and morphologies. Through production of aragonitic shells, they are involved in biogenic carbon export from the surface to the deep ocean (Bednaršek et al., 2012a). However, especially in the open Atlantic Ocean, their species diversity and abundance are still poorly understood (e.g., Bé and Gilmer, 1977; Van der Spoel and Heyman, 1983; Richter and Seapy, 1999; Van der Spoel and Dodon, 1999; Bednaršek et al., 2012a). The pteropod distribution maps from Bé and Gilmer (1977) and Van der Spoel and Heyman (1983) are based on manual estimations of distributions based on synthesized presence-only data, precluding any quantitative comparisons within and between ocean basins. Gaining more insight into the current diversity and distribution of planktonic gastropods is important for measuring future changes in diversity and distribution, and for predicting...
species-specific sensitivities to ocean changes. New data on abundance will help estimate the biomass of holoplanktonic gastropods, about which little is known for many regions, including the open Atlantic Ocean. This is important for their incorporation as a plankton functional type in ecosystem models and to estimate their contribution to ocean carbon export in biogeochemical models (Bednáršek et al., 2012a).

The oceans appear to have more biodiversity and less homogeneous species communities than has long been assumed or expected (Angel, 1993; Norris, 2000). Biogeographical provinces of the Atlantic Ocean provide a diverse range of ecological niches as well as barriers to dispersal for zooplankton taxa (e.g., Peijnenburg and Goetze, 2013; Andrews et al., 2014; Goetze et al., 2016). For example, in Cuvierina pteropods, C. cancapse is endemic to the equatorial Atlantic and C. atlantica is restricted to the northern and southern subtropical gyres (Janssen, 2005; Burridge et al., 2015). Endemism of evolutionary lineages in the equatorial Atlantic is a pattern also observed in other plankton groups (e.g., Hirai et al., 2015; Goetze et al., 2016). The species composition of planktonic ostracod assemblages also is more similar within ocean provinces (e.g., northern temperate, northern subtropical gyre and equatorial regions) than between provinces (Angel et al., 2007). Ocean warming and acidification may cause shifting species distributions in the plankton (e.g., Beaugrand et al., 2009; Provan et al., 2009), as well as (local) extinctions or adaptation to new conditions (e.g., Collins, 2012; Lobbeck et al., 2012, 2014).

Pteropods are a group of heterobranch gastropods (Jörger et al., 2010), a superorder comprised of the orders Thecosomata, also referred to as “sea butterflies”, and Gymnosomata, or “sea angels” that play an important role in marine food webs (Lalli and Gilmer, 1989; Pierrot-Bults and Peijnenburg, 2015). According to the World Register of Marine Species (WoRMS, www.marinespecies.org), a total of 83 extant thecosome species and 43 gymnosome species occur worldwide, and many fossil species also have been described (Janssen and Peijnenburg, 2013). Most taxa are between 0.05 and 2 cm in size and occur in the upper 300 m of the global ocean, although some species occur at bathypelagic depths (Bé and Gilmer, 1977; Van der Spoel and Dodan, 1999). Many species are known to be vertical migrants that move to shallower depths at night (Bé and Gilmer, 1977). Most thecosomes have aragonite shells, which they maintain as adults. They include the cavolinid endoecosmes with uncoiled, bilaterally symmetrical shells (52 species), limacinid endoecosmes with coiled shells (8 species), and pseudothecosomes, with coiled shells or a semi-soft pseudocochn without an aragonitic shell (23 species; Van der Spoel and Dodan, 1999). All thecosomes produce and deploy mucous webs to feed on microplancton (Gilmer and Harbison, 1986). Gymnosomes have larval shells, but shed them as they grow into adults. They are predators that feed exclusively or primarily on thecosomes (Lalli and Gilmer, 1989).

Heteropods or “sea elephants”, formally known as Pterotraucoidea, are a group of Caenogastropoda. According to WoRMS, this group consists of 35 extant species that occur in moderately low abundance in the global ocean, primarily at tropical and subtropical latitudes (Richter and Seapy, 1999). However, there is evidence that some taxa thrive in cold waters, e.g., south of Australia and during glacial periods in the geological past (Howard et al., 2011; Wall-Palmer et al., 2014). The vertical distribution of heteropods is not well understood, but they have been found to live at epipelagic and upper mesopelagic depths (e.g., Lalli and Gilmer, 1989; Richter and Seapy, 1999; Ossenbrügger, 2010). Heteropods are visual predators with well-developed eyes. To the extent that it is known, adults feed on other zooplankton and small fishes, and juveniles feed on phytoplankton and small zooplankton (Lalli and Gilmer, 1989; Seapy et al., 2003). The three heteropod families, Atlantidae, Carinariidae and Pterotracheidae, are highly diverse in size and body form. The Atlantidae have the highest species diversity (21 species; one additional species described by Wall-Palmer et al., 2016b), are generally less than 1 cm in size, are the least efficient swimmers of all heteropods, and can retract their bodies entirely into their keeled, dextrally coiled aragonite shells (Seapy et al., 2003). They are the most thoroughly sampled heteropod group in the southern subtropical Atlantic, with species densities of 410–1710 individuals per 1000 m² accounting for 80–99% of the total sampled heteropod community (Richter and Seapy, 1999). The Carinariidae (9 species) have a cylindrical body that is very large in relation to their shells, and can be as long as 50 cm (Lalli and Gilmer, 1989). The Pterotracheidae (5 species) only have larval shells, and have an elongated body that can reach a length of 33 cm in the Atlantic Ocean (Richter and Seapy, 1999). The Carinariidae and Pterotracheidae are thought to be more efficient swimmers than Atlantidae because of the relatively small body size and large shell of the latter group (Lalli and Gilmer, 1989).

In this study, we quantitatively sampled pteropods and heteropods from 46°N to 46°S along a transect of >12,000 km across the Atlantic Ocean, during Atlantic Meridional Transect (AMT) cruise 24. The AMT is a multidisciplinary programme aimed at understanding biological, chemical, and physical oceanographic processes, with annual transect sampling across the Atlantic Ocean (Rees et al., 2015). Here we (1) assess species distributions, abundances and biomass of pteropods and heteropods at 31 sampling stations along the transect, (2) estimate biogeographical trends in species diversity, and (3) compare inter-station similarities in community structure with Longhurst’s (1998) biogeochemical provinces. To our knowledge, this is the first paper to report quantitative abundance data of pteropods and heteropods across a large-scale latitudinal gradient in the Atlantic.

2. Methods

2.1. Sampling and sorting

Holoplanktonic gastropods were collected during the AMT24 expedition (United Kingdom to Falkland Islands) between 46°23′N and 46°05′S from September 28th to October 30th, 2014 (Table 1). We conducted oblique tows at night from the upper 311 (216–401) m at 31 stations, representing the epipelagic and upper mesopelagic zones. The stations were assigned to biogeochemical provinces according to Longhurst (1998) and Reygondeau et al. (2013) (Table 1). We used a bongo net of 0.71 m diameter and 200 μm mesh size with a General Oceanics flowmeter (2030RC) mounted in the mouth of the net to measure the volume of seawater filtered during the tow. The flowmeter was calibrated both pre- and post-cruise. Bulk zooplankton samples were quantitatively split using a Folsom plankton splitter and one or more quantitative fractions were immediately preserved in 96% ethanol. The alcohol was replaced within 24 h of collection, and samples were stored at ~20°C. Pteropods and heteropods were sorted from 25% or 50% of the quantitative zooplankton samples, depending on the bulk sample volume (Table 1).

Pteropod and heteropod material was sorted by species and counted after the expedition. The majority of taxa could be identified to species morphologically. However, we counted individuals in larval or juvenile growth stages separately if they could not be assigned to species because they lacked the morphological characters to distinguish between closely related species. This ambiguity occurred for juvenile Cuvierina endoecosmes (Bé et al., 1972); larval stages of Diacria and Cavolinia endoecosmes, Peracela pseudothecosomes, as well as larval stages of some atlantid heteropods. Some other taxa were registered only at the genus
level because of their complex and unresolved taxonomies, including *Diacavolinia* euthecosomes, *Corolla* and *Glea* pseudothecosomes, and *Pterotrachea* heteropods (Van der Spoel et al., 1993; Richter and Seapy, 1999; Van der Spoel and Dadon, 1999). Two for-}mae of the euthecosome *Clio pyramidata*, forma *lanceolata* and forma *pyramidata*, were recorded together as *C. pyramidata* because these varieties were not found to be genetically distinct (Jennings et al., 2010), and likely represent a single species. We identified gymnosomes to order because we could not identify them to species following fixation in ethanol, and excluded them from many subsequent analyses. The ratio between the abun-}dances of gymnosomes, if present, and thecosomes was calculated to estimate the potential predation intensity by gymnosomes on thecosomes across the AMT24 transect. This ratio was calculated only for stations with more than 10 pteropod specimens.

### 2.2. Species diversity, abundance and biomass

To summarize the diversity of thecosomes and heteropods along AMT24, we calculated species richness $R$, Shannon-Wiener’s diversity index $H'$ and Pielou’s evenness index $J'$ at each station. Shannon-Wiener's $H'$ is commonly used as a measure for species diversity in a community and accounts for abundance and evenness of the species present (Shannon and Weaver, 1949; Spellberg and Fedor, 2003):

$$H' = -\sum_{i=1}^{R} (p_i \ln p_i)$$

In this formula, $R$ is the species richness or total number of species, and $p_i$ is the fraction of individuals belonging to species $i$ relative to the total number of specimens. Pielou’s evenness $J'$ is derived from $H'$ and quantifies how close in numbers or abundance each species is in an environment, given that there are at least two species present per location (Pielou, 1967):

$$J' = \frac{H'}{\ln R}$$

To assess trends in species diversity and abundance of theco-}somes and heteropods, we analyzed our gastropod diversity and abundance data in relation to oceanographic data obtained during AMT24. Abundances of pteropods and heteropods were quantified in terms of individuals per 1000 m$^3$ of seawater filtered. Ocean temperature and chlorophyll $a$ concentrations along AMT24 were obtained using a Sea-Bird Electronics 3P Temperature Sensor and a CTG Aquatracka MKIII Fluorometer. Oceanographic data were calibrated and archived by the British Oceanographic Data Centre (BODC, www.bodc.ac.uk). Sea surface temperature was measured at 10 m depth. Contour plots of temperature and chlorophyll $a$ concentration in the upper 300 m were prepared in Ocean Data View 4 (Schlitzer, 2015). This depth range represents the zooplankton tow depth as well as the depth range of most pteropod and heteropod species. We examined whether species richness $R$ showed a relation-ship with sea surface temperature and maximum chlorophyll $a$ in the upper 300 m using regression analysis in the software package PAST 2.17 (Hammer et al., 2001).

We calculated the pteropod biomass in terms of total dry weight from the abundances and shell sizes using mostly genus-specific formulae from Bednarské et al. (2012a). The formulae are presented in Supplementary Table 1. These were only available for euthecosomes and gymnosomes, so we did not include shelled pseudothecosomes (*Peraclea* species) and heteropods in our calculations. These formulae use estimated averages of shell length and width. Because some suggested size averages clearly overesti-
mated the observed sizes along AMT24, we adjusted them to re-
semble the collected specimens along AMT24 (Supplementary
umber Table 1). We did this for Clio cuspidata, C. pyramidalis, C. pyramidalis
antarctica, C. pyramidalis sulcata, Cuverina sp., Styliola subula,
Hyaloclys striata and gymnosomes. Average sizes of Clio recurva,
Cavolinia gibbosa and Diacavolinia were not indicated by
Bednaršek et al. (2012a), so we used our own size estimates as well
as their genus-level formule to estimate their dry weight (using
the Cavolinia formula for Diacavolinia). We used the formula for
Limacina helicina (Bednaršek et al., 2012b) for all coiled eucthe-
somes. Bednaršek et al. (2012b) estimated that ~27% of the total
carbon of H. helicina antarctica consisted of inorganic carbon.

2.3. Species composition

For subsequent analyses of inter-station similarities of species
composition, sampling completeness and relative species domi-
nance, we reduced our abundance dataset to exclude small sample
sizes as follows. We excluded all stations with fewer than 10 the-
cosmes or heteropods, all larval and juvenile specimens that
could be assigned to two or more species, and all identified species
that only occurred at one selected station. The stations that were
excluded from analyses of thecosome species composition were
1–4 and 31 because there were fewer than 10 specimens that could
be identified to species or genus level. The thecosome species or
genera that were excluded because they were observed at only
one station were Cavolinia gibbosa, Clio recurva, Diacria major, Gleba
and Corolla. Cuverina juveniles and adults were binned into a sin-
gle genus-level group because upon exclusion of the juveniles
there would not be enough specimens (one adult: Cuverina atlant-
tica). For heteropods, we excluded stations 1–5, 12, 22, 24–26 and
30–31; also Atlanta oligogyra was excluded.

To examine whether the distributions of distinct species assem-
blages correspond with the distribution of Longhurst’s (1998) bio-
geochoreo provinces as defined by biogeochemistry and eco-
system dynamics, we quantified and visualized inter-station similarities of species composition and identified key species. For
inter-station comparisons we performed a hierarchical cluster
analysis, similarity profile analysis (SIMPROF), and non-metric mul-
dimensional scaling analysis (nMDS) in PRIMER 6, without a priori assumptions (Clarke, 1993; Clarke and Warwick, 2001;
Clarke and Gorley, 2006). To perform cluster analysis, SIMPROF
and nMDS, we standardized and transformed (log(x + 1)) the abun-
dance data and then calculated a Bray-Curtis similarity matrix. For
the cluster analysis we used the group average setting. The signifi-
cance of the clusters was tested with SIMPROF analyses using
1000 permutations and a significance level of p < 0.05. The nMDS
ordinations were performed with 25 restarts. Furthermore, we cre-
ated rank abundance curves based on the thecosome species abun-
dances at each station in order to assess patterns of dominance
across ocean biomes.

To assess sampling completeness, we created sample-based as
well as individual-based rarefaction curves for thecosomes and
heteropods in Primer 6 (Clarke and Gorley, 2006) and PAST 2.17
(Hammer et al., 2001). The sample-based curves were based on the
Jackknife 2 index (Gotelli and Colwell, 2010) and only included
non-transitional stations that belong to distinct biogeochemical
provinces.

3. Results

3.1. Species abundance, biomass and diversity

We counted and identified a total of 7312 pteropods across all
stations on the transect, traversing the northern temperate zone,
the eastern side of the northern subtropical gyre, the equatorial
upwelling zone, the southern subtropical gyre, the southern sub-
tropical convergence and the northernmost part of the subantarctic
(Fig. 1A; Table 1). Among the pteropod specimens, there were 1028
uncoiled and 5980 coiled eucthecosomes, 230 pseudothecosmes and
74 gymnosomes. Pteropod abundances were low (<15 ind./1000 m³) north of 40°N and they were absent from station
3. Their abundance varied between 100 and 2000 ind./1000 m³
between 30°N and 40°S. Abundances were highest at stations 27–
29, just south of 40°S, with a maximum abundance of
>4000 ind./1000 m³ at station 28 (Figs. 1B, C and 2A; Supplemen-
tary Table 2). Coiled eucthecosmes were particularly abundant
in this high productivity area, while pseudothecosmes were more
abundant in the subtropical gyres.

Gymnosomes feed almost exclusively on thecosomes. The ratio
between gymnosomes, if present, and thecosomes (eucthecosmes
and pseudothecosmes) ranged from 0.005 to 0.019 in (sub)tropi-
cal waters between ~28°N and ~28°S (stations 7–23) (Supplemen-
tary Table 3). Gymnosomes were always present between ~32 and
~45°S (stations 24–30), with ratios between 0.012 and 0.091, sug-
gesting higher potential predation intensity by gymnosomes on
thecosomes at these southern stations than in (sub)tropical waters.

We observed taxa with strong preferences for particular oceanographic provinces, in particular among eucthecosmes and
pseudothecosmes (Fig. 1C–E). In the equatorial province and in
the subantarctic, the thecosome community consisted of one or
two dominant species, and especially in the equatorial region a
number of additional species occurred in low abundance. The re-
late species abundances were more equal in the subtropical gyres
(Supplementary Fig. 1; abundances per species per station are
listed in Supplementary Table 2). Uncoiled eucthecosmes were
found between 46°N and 43°S (Fig. 1D). Two Clio taxa, C. pyrami-
data sulcata and C. pyramidalis antarctica, were restricted to the
subantarctic, but were never dominant. Clio pyramidalis pyrami-
data/lancaelata was found in all other regions along the transect.
Key species that were most abundant in the subtropical gyres
included Creseis clava, Diacria danae and Styliola subula, the latter
of which was entirely absent in the equatorial upwelling region.
Although Cavolinia inflexa was present across warm water environ-
ments, this species only occurred in high abundances in the south-
ern gyre. Creseis virgula was found in the equatorial region and
occurred nowhere else along the transect (Supplementary Table 2).
Coiled eucthecosmes were found between 35°N and 46°S (Fig. 1E).
Helicocinoides inflatus was abundant along the entire transect and
did not show a clear preference for particular provinces. However,
H. inflatus specimens in the subantarctic region were morphologi-
cally distinct from warm water H. inflatus specimens. Shells of the
cold water form (listed as H. inflatus S herein) appeared to be coar-
ser and thicker and, in contrast to the warm water form, they had a
reddish hue along the whorls and aperture. Limacina helicina
antarctica was dominant in subantarctic waters. Limacina buli-
moides had a strong preference for the subtropical gyres, although
it was also present in low numbers in the equatorial region. Limac-
ina lessuerti was present throughout the warm water regions and
L. trochiformis was only found in the southern gyre, though never in
high numbers (maximum N = 9 at St. 20; Fig. 1E). Most Peracel
pseudothecosme species demonstrated a strong preference for the
subtropical gyres (Fig. 1C), but the large, soft-bodied genera
Corolla and Gleba were only found in cooler waters north of 40°N
and south of 40°S in very low numbers (never more than N = 2
per station).

Areas with high chlorophyll a concentrations had the highest
estimated pteropod biomass, except the northern temperate zone,
where hardly any pteropods were captured (Fig. 2B; Supplemen-
tary Table 4). A total biomass of more than 3.1 mg m⁻³ in the upper
~300 m was reached in the southern subtantarctic region, mainly

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caused by high abundances of *Limacina helicina antarctica* and *Heli-conoides inflatus* S. Just south of the equator there was an estimated biomass of more than 1.9 mg m\(^{-3}\) because of high abundances of *Cavolinia inflexa* and *Clio pyramidata*. The median and average pteropod biomass across all stations along AMT24 was 0.28 mg m\(^{-3}\) and 0.49 mg m\(^{-3}\), respectively.

The species richness of thecosomes was highest in the stratified (sub)tropical waters between 24°N and 24°S and was consistently high (11–13 species) in the southern gyre (Fig. 2C and D), a pattern that was also evident from the Shannon-Wiener’s diversity indices \(H'\) per station (Supplementary Table 3). Uncoiled euthecosomes generally had a higher species richness than coiled euthecosomes and pseudothecosomes (Fig. 2C). Although most equatorial stations had slightly lower species richness than the gyre stations, the highest species richness (15 species) was observed close to the equator at station 14. Species richness showed a positive relationship with sea surface temperature \(R^2 = 0.55; N = 30; p < 0.001\); Fig. 3A) and a negative relationship with chlorophyll \(a\) concentration at the deep chlorophyll maximum \(R^2 = 0.57; N = 30; p < 0.001\); Fig. 3B). We note that sea surface temperature and chlorophyll \(a\) were only weakly correlated \(R^2 = 0.28; N = 31; p < 0.01\), and hence the degree of collinearity between these two explanatory variables was low. The species evenness \(J'\) was highest in the subtropical gyres, especially in the southern gyre \(J'\) up to 0.84), when disregarding the high evenness values observed at the northern temperate stations 1, 2 and 5 because of their relatively low sample sizes. At equatorial stations 13–16 the evenness was lower \(J'\) of 0.25–0.44) than in the gyres. From the 1812 heteropods, a total of 1312 were atlantids, 325 were pterotracheids and 175 were carinariids. Heteropods were not found at stations 1–4 and 30–31. Highest heteropod species richness (up to 14 species) as well as abundances (maximum 704 ind./1000 m\(^3\)) were found in the (sub)tropical waters between 30°N and 20°S (Fig. 4). A high abundance of heteropods (>100 ind./1000 m\(^3\)) was also found in the subantarctic region (Supplementary Table 2). There were no clear patterns in species evenness along the transect except a lower evenness at stations 28–29 \(J'\) of 0.10–0.37; Supplementary Table 3).

### 3.2. Species composition

The cluster and SIMPROF analyses of thecosome species composition of the different stations along AMT24 resulted in six significant clusters \(p < 0.05\); Fig. 5A). These results are further confirmed by the nMDS ordination, in which the six different clusters can be clearly recognized (Fig. 5B and C). The species composition in the subantarctic region (St. 26–30) demonstrated almost no overlap with species compositions in other regions. The other clusters consisted of the equatorial stations (St. 11–16), the northern and southern gyres together (St. 6–9 and 18–23), the northern gyre-equatorial transition (St. 10), the southern subtropical convergence (St. 25), and the northern temperate zone and southern gyre.
Fig. 2. Basin-scale patterns of (A) pteropod abundance (individuals/1000 m$^3$), (B) pteropod biomass in terms of total dry weight (mg m$^{-3}$), (C) pteropod species richness $R$, (D) seawater temperature and (E) chlorophyll a concentrations in the upper 300 m of the water column, measured during Atlantic Meridional Transect cruise 24. Lines in (A) are interrupted in case of zero abundances. Because gymnosomes were not identified to the species level, they were not included in the calculation of species richness $R$ in (C).
equatorial transition (St. 5 and 17) together. Because of the overlap between species compositions in the northern temperate and in the northern and southern gyre-equatorial transitions, the ordination of St. 5 and 17 may be an artefact of the gradual nature of the transitions between oceanographic provinces (Fig. 5). According to the rarefaction curves of the equatorial, gyre and subantarctic stations, the major oceanographic provinces traversed during AMT24 were well-sampled for thecosomes (Supplementary Fig. 2A). Although transitions between oceanographic provinces were gradual, we were not able to thoroughly sample these areas at more than one location per transition (Supplementary Fig. 2B).

Heteropod species had a wider distribution than the subtropical gyres and equatorial regions: some of them also occurred in the subantarctic (Figs. 1B and 4). This was the case for an exclusively subantarctic and yet undescribed Atlanta morphotype, listed as Atlanta species A herein (Wall-Palmer et al., 2016b). In particular, the taxa that produce very large adults, the carinariids and pterotracheids, had patchy distributions, with high numbers at a few stations and no or a few specimens at the adjacent stations (Supplementary Table 2). It was therefore difficult to identify large-scale biogeographical distribution patterns among the heteropods. This patchy pattern was also reflected in rarefaction curves of heteropods, which were represented by fewer samples with at least 10 specimens compared to pteropods. Suboptimal sampling of the heteropods was demonstrated especially in the rarefaction curves representing the northern subtropical gyre and subantarctic stations (Supplementary Fig. 2C and D). We did not generate rarefaction curves for heteropods from northern and southern temperate stations because they contained too few (<6) specimens.

4. Discussion

4.1. General observations

We assessed pteropod and heteropod abundances, diversity and distributions in the Atlantic Ocean based on quantitative sampling along a basin-scale transect from 46°N to 46°S, crossing seven biogeochemical provinces. The AMT programme allowed us to combine zooplankton diversity and abundance data with ocean metadata from the same expedition, providing a rich oceanographic context to explain the observed patterns (e.g., Rees et al., 2015). Our results show that the species richness of both pteropods and heteropods varied significantly across the latitudinal gradient.
Fig. 5. (A) Hierarchical cluster analysis of the stations according to their community composition of thecosome pteropods (euthecosomes and pseudothecosomes). The analysis is based on all stations with at least 10 specimens, and resulted in the indicated six significant clusters (SIMPROF p < 0.05). (B and C) nMDS ordination based on the community composition of thecosome pteropods, (B) using all stations with at least 10 specimens, and (C) excluding the subantarctic stations 26–30. Symbols are colored according to their geographical location, with upward pyramids for the northern hemisphere, downward pyramids for the southern hemisphere and diamonds for equatorial locations. The Longhurst provinces (Longhurst, 1998; Reygondeau et al., 2013) of the sampling stations are indicated in (A): NAST E (Northeast Atlantic subtropical gyral), NAST W (Northwest Atlantic subtropical gyral), NATR (North Atlantic tropical gyral), WTRA (Western tropical Atlantic), SATL (South Atlantic gyral), SSTC (South subtropical convergence), SANT (Subantarctic water ring), FKLD (Southwest Atlantic shelves).

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and heteropods was highest in the tropical and subtropical Atlantic. Pteropod abundance and biomass were highest just south of 40°S, and also reached high values near the equator (Figs. 1 and 2; Supplementary Tables 2 and 4). Some pteropod taxa showed strong preferences for particular oceanographic provinces, for example for warm equatorial waters (e.g. Creseis virgula), subtropical gyres (e.g. Styliola subula), or in and south of the southern subtropical convergence zone (e.g. Limacina helicina antarctica). Other species occurred across a broad range of provinces (e.g. Heliconoides inflatus and Limacina lesueurii; Supplementary Table 2). Most heteropods were restricted to warm waters, but some taxa also occurred in the subantarctic region.

We found no unexpected extremes in pteropod abundance along AMT24 (Fig. 2A) and the rarefaction curves flattened with an increasing number of samples or individuals (Supplementary Fig. 2A and B), indicating that the pteropods in this study have been representatively sampled. However, it is likely that our net aperture (0.71 m) and mesh size (200 µm) are responsible for highly variable heteropod species counts, diversities and abundances across stations, and quantitative sampling of these larger taxa requires nets with larger apertures and mesh sizes (McGowan and Fraundorf, 1966; Wells, 1973; Wall-Palmer et al., 2016a). Hence, we will focus most of our discussion on pteropods.

4.2. Distribution and abundance

The biogeographical distributions of pteropod assemblages inferred by cluster analysis were largely congruent with the distribution of Longhurst’s (1998) biogeochemical provinces for the period from September to November (Fig. 5; Reygondeau et al., 2013). Our pteropod species distributions in the Atlantic Ocean also were comparable with those from the qualitative biogeographical synthesis of Bé and Gilmer (1977), however, there were some exceptions. According to Bé and Gilmer, Limacina trochoformis was most abundant in the equatorial province (WTRA). By contrast, we observed L. trochoformis only at station 14 (N = 3) in the equatorial province, but found much higher abundances throughout the south subtropical gyre. Exact localities upon which the distribution maps of Bé and Gilmer (1977) were based were not given, so we could not deduce if and where in the South Atlantic their samples were collected. We found a cold water variant of Heliconoides inflatus (H. inflatus S herein) in the subantarctic, but the distribution map from Bé and Gilmer (1977) most likely shows the broad equatorial and subtropical distribution of the warm water H. inflatus (listed as Limacina inflata) with a southern limit at ~40°S. The distribution of Creseis virgula was limited to the equatorial Atlantic in our study, but also occurred in the subtropical gyres according to Bé and Gilmer (1977). They also listed some species that were not found in our quantitative samples: the warm water species Cavolinia tridentata and the cold water species Limacina retroversa.

Because previous quantitative abundance data in provinces traversed by AMT24 are limited, our abundance data could only be compared to a few other pteropod studies. In October, St. 30 and 31 are within a transition zone between the Subantarctic water ring (SANT) and Southwest Atlantic shelves (PKLD) provinces (Reygondeau et al., 2013). Hunt et al. (2008) synthesized pteropod abundances in the Southern Ocean from tow data (1982–2006). In October, St. 30 and 31 (45–46°S) in October. Much higher population densities of this species were found in the mid austral summer (January–February), with 10^3 to 10^5 ind./1000 m^3 at the Antarctic Peninsula and south of Australia, and even 10^5 ind./1000 m^3 near South Georgia (Hunt et al., 2008). Bednaršek et al. (2012b) report similar seasonal dynamics from the Antarctic Peninsula to South Georgia, with higher numbers of L. helicina antarctica in the austral summer and autumn than in the spring. We did find high densities of L. helicina antarctica, exceeding 10^3 to 10^5 ind./1000 m^3 in October, further up north at St. 27 and 28 (40–41°S), which may indicate that the growing season for this species started earlier at lower latitudes.

The total zooplankton abundance in the upper 200 m of the northeast Atlantic Ocean during July 1996 was characterized by Gallienne et al. (2001) by vertical hauls of 200 µm mesh size nets (aperture 0.57 m). They found low pteropod abundances of 0–75 ind./1000 m^3 between 39 and 47°N, comparable to 0–24.2 ind./1000 m^3 between 35 and 46°N in our study in late September/early October (St. 1–5). Gallienne et al. (2001) found higher summer abundances of 845–1730 ind./1000 m^3 at 37°N, whereas in our study higher abundances were found south of 31°N in early fall, which may again reflect seasonal variation.

In the Caribbean Sea, not sampled in our study, the average species abundances in the 239 µm mesh size net (aperture 1.0 m) used by Wells (1973) near Barbados resemble our findings in equatorial St. 14. For Creseis virgula, Wells (1973) found 16.9 ind./1000 m^3, which is highly comparable to the 18.4 ind./1000 m^3 in our study. For Heliconoides inflatus, Wells (1973) found 257.3 ind./1000 m^3 resembling the 239.8 ind./1000 m^3 in our study. The major difference was the high abundance of Creseis conica in the Caribbean (188.2 ind./1000 m^3) reported by Wells (1973; listed as Creseis virgula conica), whereas we found only 16.8 ind./1000 m^3 at our St. 14.

4.3. Plankton biomass in the global ocean

Qualitatively, the pteropod biomass distribution along AMT24 showed a similar large-scale pattern as the biomass distributions of many other mesozooplankton groups, with high zooplankton biomass in arctic and subantarctic waters, elevated biomass in the equatorial regions, and lowest biomass within the gyres (Moriarty and O’Brien, 2013). Quantitatively, however, biomass data from sampling locations between 10 and 500 m deep in various parts of the Atlantic Ocean as synthesized by Bednaršek et al. (2012a) point at a higher pteropod biomass (dry weight) in the Atlantic Ocean than in the Indian and Pacific oceans. This was true for the equatorial Atlantic north of Brazil, the Mediterranean Sea, and the subantarctic near South Georgia. Our data appear to confirm this. Compared to the global median pteropod biomass of 0.058 mg m^−3 (only non-zero global data) reported by Bednaršek et al. (2012a), we found a much higher pteropod biomass per m^3 in the Atlantic Ocean (Fig. 2B; median of 0.28 mg m^−3).

4.4. Latitudinal trends in species richness

The dominant paradigm for latitudinal diversity patterns in pelagic systems is a bimodal pattern of species richness, with maxima in the gyres, slightly lower richness in the equatorial province, and a sharp decrease of diversity towards the temperate and polar zones (e.g. Hillebrand, 2004a). Our pteropod data broadly support this pattern, and with the exception of one equatorial station, diversity at most stations in the equatorial province was slightly lower than in the gyres (Fig. 2C). Many hypotheses have been formulated for advancing the ecological, biogeographic, and evolutionary understanding of the latitudinal diversity gradient, and for explaining why the gradient in the pelagic ocean differs from a steady increase of diversity towards the equator, the most widely recognized pattern in benthic and terrestrial environments (e.g. Angel, 1997; Gaston, 2000; Willig et al., 2003; Hillebrand, 2004a, b; Brayard et al., 2005). Examples of such conceptual models are
the evolutionary speed hypothesis and geometric constraints hypothesis (Willig et al., 2003).

Several pelagic groups show latitudinal trends in species richness similar to pteropods, with the highest richness in the subtropical gyres. Such a pattern was reported for hydromedusans (Macpherson, 2002), euphausiids (Angel, 1997; Tittensor et al., 2010), pelagic decapods (Angel, 1997), ostracods (Angel, 1997; Angel et al., 2007), fish (Angel, 1997), foraminifera (Rutherford et al., 1999; Tittensor et al., 2010), and tintinnid ciliates (Dolan and Pierce, 2013). Several other pelagic taxa do not show a bimodal pattern with maximum species richness in the subtropical gyres, but display a diversity peak in the equatorial region or a diversity plateau across subtropical and tropical latitudes. Examples of equatorial maxima are found for salps (Macpherson, 2002) and hyperiid amphipods (Burridge et al., 2016), and our data indicate that heteropods also show maximum species richness in equatorial waters. A wider diversity plateau across the tropics and sub tropics was found for copepods (Woodd-Walker et al., 2002; Rombouts et al., 2009), siphonophore hydromedusans and cephalopods (Macpherson, 2002). Larvacean species diversity only had a clear peak in the northern gyre (Macpherson, 2002), and chaetognaths only demonstrated a peak in the southern gyre (Macpherson, 2002) with a distinct decrease in species diversity towards cold North Atlantic waters (42–59°N; Pierröt-Bults, 2008). However, the reported species diversities in the latter two groups were lower (maxima of up to ~50 and ~40 species, respectively) than for hydromedusans or crustaceans. Species diversities registered for pteropods were also comparatively low. Accurate estimates of pteropod species diversities may require more rigorous sampling efforts and more accurate assessments of species boundaries. Our anomaly of high diversity at equatorial station 14 was largely caused by the occurrence of pteropod taxa that are generally known to appear in low abundances and that were only sporadically collected along AMT24, e.g. Diacavolinia sp. and Hyalocysis striata. Repeated transect studies may compensate for such anomalies. Overall, the latitudinal diversity gradient varies among zooplankton groups, and appears to result from a complex interplay among ecological factors, highly diverse life history strategies and roles in the marine food web, as well as different evolutionary histories.

5. Conclusions

To our knowledge, this is the most comprehensive study of the diversity and abundance of planktonic gastropods across the Atlantic Ocean to date. We found close correspondence between our pteropod assemblages and Longhurst’s (1998) biogeographical provinces. On average, our results point to a substantially higher pteropod biomass in the Atlantic Ocean than in the Indian and Pacific oceans. The dominant paradigm of a bimodal pattern of species richness in pelagic systems, with maxima in the subtropical gyres, was broadly supported by our pteropod data. Our study provides only a snapshot in time, however. Thorough repeated sampling will be essential for examining large-scale, long-term trends in the diversity and abundance of planktonic gastropods, quantifying future changes, and predicting species-specific sensitivities to ocean changes.

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Appendix A. Supplementary material

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