

# Copepod species abundance from the Southern Ocean and other regions (1980 - 2005) – a legacy

Astrid Cornils, Rainer Sieger†, Elke Mizdalski, Stefanie Schumacher, Hannes Grobe, Sigrid B. Schnack-Schiel†

Alfred-Wegener-Institut Helmholtz-Zentrum für Polar- und Meeresforschung, Bremerhaven, Germany

†: deceased

Correspondence to: Astrid Cornils (astrid.cornils@awi.de)

## Abstract.

**This data collection originates from the efforts of Dr. Sigrid Schnack-Schiel (1946 – 2016), a zooplankton ecologist with great expertise in life cycle strategies of Antarctic calanoid copepods, but who also investigated zooplankton communities in tropical and subtropical marine environments. Here,** we present 33 data sets with abundances of planktonic copepods from 20 expeditions to the Southern Ocean (Weddell Sea, Scotia Sea, Amundsen Sea, Bellingshausen Sea, Antarctic Peninsula), one expedition to the Magellan region, one latitudinal transect in the Eastern Atlantic Ocean, one expedition to the Great Meteor Bank and one expedition to the northern Red Sea and Gulf of Aqaba **as part of her scientific legacy. A total of 349 stations from 1980 to 2005** were archived. During most expeditions depth-stratified samples were taken with a Hydrobios multinet with 5 or 9 nets, **thus allowing inter-comparability between the different expeditions. Only during four cruises a Nansen or a Bongo net** was deployed. **Maximum sampling depth varied greatly among stations due to different bottom depths. However, during eleven cruises to the Southern Ocean the maximum sampling depth was restricted to 1000 m, even at locations with greater bottom depths. In the eastern Atlantic Ocean (PS63) sampling depth was restricted to the upper 300 m. All data are now freely available at PANGAEA via the persistent identifier <https://doi.org/10.1594/PANGAEA.884619>.**

Abundance and distribution data for 284 calanoid copepod species and 28 taxa of other copepod orders are provided. **For selected species the abundance distribution at all stations was explored, revealing e.g. that species within a genus may have contrasting distribution patterns (Ctenocalanus, Stephos). In combination with the corresponding metadata (sampling data and time, latitude, longitude, bottom depth, sampling depth interval) the analysis of the data sets may add to a better understanding how the environment (currents, temperature, depths, season) interacts with copepod abundance, distribution and diversity. For each calanoid copepod species females, males and copepodites were counted separately, providing a unique resource for biodiversity and modelling studies. For selected species also the five copepodite stages were counted separately, thus also allowing to use the data to study life cycle strategies of abundant or key species.**

## 1 Introduction

Copepoda (Crustacea) are probably the most successful metazoan group known, being more abundant than insects, although far less diverse (Humes, 1994; Schminke, 2007). They occur in all aquatic ecosystems, from freshwater to marine and hypersaline environments, and from polar waters to hot springs (Huys and Boxshall, 1991). Although copepods are evolutionary of benthic origin (Bradford-Grieve, 2002), they have also successfully colonised the pelagic marine environment where they can account for 80 – 90% of the total zooplankton abundance

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### [1] verschoben (Einfügung)

**Gelöscht:** This data compilation represents the scientific legacy of Dr. Sigrid B. Schnack-Schiel (1946-2016), revealing her expertise and great interest in polar zooplankton ecology, but also exploring the zooplankton communities of tropical and subtropical regions. Copepods are often the predominant taxa in marine zooplankton and play an important role in the food web as intermediators between primary producers, the microbial loop and higher trophic levels. Due to their short life cycles and their rapid response to changing environments they are good indicators for ecosystem health and status. Investigating the effects of environmental change on planktonic copepods and thus the pelagic ecosystem requires data on species abundance and distribution. Here,

**Gelöscht:** and occurrence

**Gelöscht:** In this data compilation a

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**Gelöscht:** These data sets are now freely available at PANGAEA via the persistent identifier <https://doi.org/10.1594/PANGAEA.884619>.

**Gelöscht:** On few occasions a

**Gelöscht:** Nansen

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**Gelöscht:** net

**Gelöscht:** The deepest sample reached down to 2880 meter. As metadata sampling date and date/time, latitude, longitude, bottom depth, sampling depth interval, volume of filtered water and information of the net type and mesh size were recorded.

**Gelöscht:** The taxonomic concept was consistent throughout the data sets. The density of

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**Gelöscht:** individual

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79 (Longhurst, 1985). In the Southern Ocean, copepods are next to Antarctic krill and salps the most important  
80 zooplankton organisms, both in abundance and biomass (e.g. Pakhomov et al., 2000; Shreeve et al., 2005;  
81 Smetacek and Nicol, 2005; Ward et al., 2014; Tarling et al., 2017). In the Southern Ocean, copepods are also the  
82 most diverse zooplankton taxon accounting for more than 300 species (Kouwenberg et al., 2015). However, only  
83 a few species dominate the Antarctic epipelagic assemblage: the large calanoids *Calanoides acutus*, *Calanus*  
84 *propinquus*, *Metridia gerlachei*, *Paraeuchaeta antarctica*, the small calanoids *Microcalanus pygmaeus*,  
85 *Ctenocalanus citer* and the cyclopoids *Oithona* spp. and species of the family Oncaeidae (e.g. Hopkins, 1985;  
86 Atkinson, 1998; Schnack-Schiel, 2001; Tarling et al., 2017). Together these taxa can comprise up to 95% of the  
87 total abundance and up to 80% of the total biomass of copepods (Schnack-Schiel et al., 1998). However, the  
88 smaller calanoid species alone can account for up to 80% of the abundance of calanoid copepods (Schnack-Schiel,  
89 2001). Stage-resolve counts for selected species will also allow future users to study life cycle strategies of  
90 abundant or key species.  
91 Numerous studies on zooplankton have been conducted in the past in the Atlantic sector of the Southern Ocean  
92 (e.g. Boysen-Ennen and Piatkowski, 1988; Hopkins and Torres, 1988; Boysen-Ennen et al., 1991; Pakhomov et  
93 al., 2000; Dubischar et al., 2002; Ward et al., 2014; Tarling et al., 2017). A major zooplankton monitoring  
94 programme in the Southern Ocean is the Continuous Plankton Recorder survey (SO-CPR), providing a large-scale  
95 coverage of surface Antarctic zooplankton species distribution abundances over the last 25 years (Hosie et al.,  
96 2003; McLeod et al., 2010). A recent review summarizes the present knowledge on abundance and distribution of  
97 Southern Ocean zooplankton (Atkinson et al., 2012). Especially in the Weddell Sea, occurrence data of copepods  
98 and other zooplankton species are scarce. One of our aims is to fill this gap with the here presented data sets from  
99 the Southern Ocean, collected by Dr. Sigrid Schnack-Schiel (1946 - 2016) over a period of 1982 to 2005.  
100 In recent years there is ample evidence that marine ecosystems are greatly affected by climate change and ocean  
101 acidification (e.g. Beaugrand et al., 2002; Edwards and Richardson, 2004; Rivero-Calle et al., 2015; Smith et al.,  
102 2016). In the Southern Ocean, the pelagic ecosystem is likely to be severely affected by increasing water  
103 temperatures and the resulting reduction of sea ice coverage in the Southern Ocean (Zwally, 1994; Smetacek and  
104 Nicol, 2005). It has already been observed over decades that the biomass of Antarctic krill decreases (Atkinson et  
105 al., 2004), but little is known about the environmental effects on copepods. Within the pelagic ecosystem  
106 zooplankton communities and thus copepods are good indicators for ecosystem health and status due to their short  
107 life cycles und their rapid response to changing environments (Reid and Edwards, 2001; Chust et al., 2017).  
108 Furthermore, they are generally not commercially exploited and thus are likely to reflect impacts of environmental  
109 changes more objectively. To better understand the effects of environmental change on planktonic copepods e.g.  
110 via biodiversity analyses and ecological niche modelling, data on species occurrence, abundance and distribution  
111 are essential. Often modelling studies however are limited by the scarcity of available plankton data (Chust et al.,  
112 2017). Thus, freely available data sets on abundance and presence/absence of copepod species are of great  
113 importance for future studies on environmental changes in the pelagic realm. The data sets presented here on  
114 copepod species and life stages (female, male, copepodites) occurrences and abundance from the Southern Ocean,  
115 the eastern Atlantic Ocean, the Magellan region and the Red Sea provide a unique resource for biodiversity and  
116 modelling studies. They may also help to enhance our understanding how the environment (currents, temperature,  
117 depths, season) interacts with copepod abundance, distribution and diversity.  
118 \_\_\_\_\_

119 **2 Methods**

Gelöscht: copepod community

Gelöscht: et al.

Gelöscht: The CPR is a plankton sampler that can be towed in approximately 10 m depth by ships of opportunity, thus allowing to rapidly sample vast regions of the oceans (Reid et al., 2003). In the device zooplankton organisms are retained by a mesh and preserved in formalin.

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139 **2.1 Sampling locations**

140 The presented data sets were collected during 24 research cruises with several research vessels from 1980 to 2005  
141 (Table 1). Most of the data sets (28 datasets from 20 cruises) are based on samples from the Southern Ocean (Fig.  
142 1), collected onboard R/V Polarstern (25 data sets from 16 cruises), R/V Meteor (1 data set), R/V John Biscoe (1  
143 data set) and R/V Polarsirkel (1 data set). Southern Ocean sampling locations were restricted to the Weddell Sea,  
144 the Scotia Sea, the Antarctic Peninsula, the Bellingshausen Sea and the Amundsen Sea (Fig. 1).

145 Additionally, four data sets were collected in other regions (Table 1). In 1994 net samples were collected onboard  
146 R/V Victor Hensen in the Magellan region. Two data sets are based on research cruises with R/V Meteor, to the  
147 Great Meteor Bank in the North Atlantic (1998) and to the northern Red Sea and the Gulf of Aqaba (1999). In  
148 2002, plankton net samples were taken during a research cruise with R/V Polarstern along a transect in the eastern  
149 tropical Atlantic Ocean (Table 1).

150 Maximum sampling depth varied greatly among stations due to different bottom depths (Table 1). However, during  
151 eleven cruises to the Southern Ocean the maximum depth was restricted to 1000 m, even at locations with greater  
152 bottom depths. In the eastern Atlantic Ocean (PS63) sampling depth was restricted to the upper 300 m.

153

154 **2.2 Sampling gear**

155 Three types of plankton nets were deployed: Bongo nets, single opening-closing Nansen nets and multiple  
156 opening-closing nets. During all expeditions vertical hauls were taken, thus allowing no movement of the vessel.

157

158 **2.2.1 Nansen net**

159 During the expeditions PS04, DAE1979/80, and JB03 net sampling was carried out with a Nansen net (Table 1).  
160 The Nansen net is an opening-closing plankton net for vertical tows (Nansen, 1915; Currie and Foxton, 1956).  
161 Thus, it is possible to sample discrete depth intervals to study the vertical distribution of zooplankton. The Nansen  
162 net has an opening of 70 cm diameter and is usually 3 m long. Two different mesh sizes were used: 200 µm for  
163 the cruises PS04 and JB03, and 250 µm for DAE1979/80. To conduct discrete depth intervals the net is lowered  
164 to maximum depth and then hauled to a certain depth and closed via a drop weight. Then the net is hauled to the  
165 surface and the sample is removed. This process of sampling depth intervals can be repeated until the surface layer  
166 is reached. The volume of filtered water was calculated using the mouth area and depth interval due to the lack of  
167 a flowmeter.

168

169 **2.2.2 Multinet systems**

170 Most presented data sets are based on plankton samples taken with a multinet system (MN) from Hydrobios (Table  
171 1) a revised version (Weikert and John, 1981) of the net described by Be et al. (1959). The multinet is equipped  
172 with five (midi) or nine (maxi) plankton nets, with a mouth area of 0.25 and 0.5 m<sup>2</sup>, respectively. These nets can  
173 be opened and closed at depth on demand from the ship via a conductor cable. Thus, they allow sampling of  
174 discrete water layers. The net system was hauled with a general speed of 0.5 m/s. Mesh sizes varied between the  
175 data sets from 55 to 300 µm (Table 1). In the Southern Ocean the mesh sizes were consistent within regions: In  
176 the Weddell Sea 100 µm mesh size was used with a few exceptions during PS06. In the Scotia Sea and near the  
177 Antarctic Peninsula a mesh size of 200 µm was employed. In the Bellingshausen Sea and the Amundsen Sea  
178 multinet hauls with 55 µm mesh sizes were carried out. In other regions mesh sizes of 100 µm (PS63, M42/3), 150

**Gelöscht:** Plankton nets are designed to capture zooplankton organisms.

181  $\mu\text{m}$  (M44/2) and  $300 \mu\text{m}$  (VH1094) were used. The MN maxi was only deployed during the research cruise M44/2  
182 in the northern Red Sea.

183 Generally, the volume of filtered water was calculated from the surface area of the net opening (midi:  $0.25 \text{ m}^2$ ,  
184 maxi:  $0.5 \text{ m}^2$ ) and the sampling depth interval. For the data sets from PS63, PS65, PS67 and M44/2 a mechanical  
185 digital flowmeter was used to record the filtering efficiency and to calculate the abundances (see Skjoldal et al.,  
186 2013, p. 4). The flowmeter is situated in the mouth area of the net and measures the water flow, providing more  
187 accurate volume values of the filtering efficiency.

188

### 189 **2.2.3 Bongo net**

190 During one research cruise (PS06) 61 additional samples were taken with the Bongo net (McGowan and Brown,  
191 1966) to study selected calanoid copepod species. The Bongo net contains two nets that are lowered simultaneously  
192 for vertical plankton tows. The opening diameter is 60 cm, and the length of the nets is 2.5 m with a mesh size of  
193  $300 \mu\text{m}$ . The volume of filtering water was recorded with a flowmeter and used for the calculation of abundance.

194

### 195 **2.2.4 Effects of variable net types and mesh sizes**

196 Quantitative sampling of copepods and zooplankton is challenging. Major sources of error are patchiness,  
197 avoidance of nets and escape through the mesh (Wiebe, 1971; Skjoldal et al., 2013). These errors are defined by  
198 mesh sizes and net types, in particular the mouth area. The effect of patchiness cannot be investigated here due to  
199 the lack of replicates.

200 To our knowledge the sampling efficiency of the Nansen net and the MN midi have not been compared directly  
201 (Wiebe and Benfield, 2003; Skjoldal et al., 2013). However, it has been stated that the catches with Nansen net  
202 are considerably lower than with the WP-2 net (Hernroth, 1987), although the WP-2 net is considered as a modified  
203 Nansen net with a cylindrical front section of 95 cm and a smaller mouth area ( $57 \text{ cm}^2$ , Skjoldal et al., 2013). The  
204 WP-2 net with  $200 \mu\text{m}$  mesh size however, is in its sampling efficiency, measured as total zooplankton biomass,  
205 comparable to the MN midi with  $200 \mu\text{m}$  mesh size (Skjoldal et al., 2013). Thus, it has to be taken into account  
206 during future analysis that the abundance values from the Nansen net are not directly comparable to those from  
207 the MN midi.

208 The mesh size has a different effect on the zooplankton catch. It is well known that small sized copepod species  
209 ( $< 1 \text{ mm}$ ) and thus in particular non-calanoid species (e.g. Oithonidae, Oncaidae) and juvenile stages also from  
210 calanoid copepods (e.g. *Microcalanus*, *Calocalanus*, *Disco*) pass through coarse mesh sizes ( $\geq 200 \mu\text{m}$ ), while  
211 they are retained in finer mesh sizes (Hopcroft et al., 2001; Paffenhöfer and Mazzocchi, 2003). Thus, abundances  
212 of smaller specimens, and the species and life stage composition may vary considerably, when comparing samples  
213 from the Bellingshausen and Amundsen Seas ( $55 \mu\text{m}$  mesh size), around the Antarctic Peninsula ( $200 \mu\text{m}$ ) and  
214 the Weddell Sea ( $100 \mu\text{m}$ ).

215

## 216 **2.3 Sample processing and analysis**

217 All samples were preserved immediately after sampling in a 4% formaldehyde-seawater solution. Samples were  
218 stored at room temperature until they were sorted in the laboratory. The formaldehyde solution was removed, the  
219 samples were rinsed and copepods were identified and counted under a stereomicroscope, [using a modified Mini-  
220 Bogorov chamber with high transparency as described in the ICES Zooplankton Methodology Manual \(Postel et  
221 al. 2000\)](#). Abundant species were sorted from one fourth or less of the sample while the entire sample was screened

Gelöscht: from a fraction of the sample

223 for rare species. Samples were divided with a Motoda plankton splitter (Motoda, 1959; Van Guelpen et al., 1982).  
224 Abundance was calculated using the surface area of the net opening and the sampling depth interval or the  
225 recordings of the flowmeter. Samples for re-analysis are only available for the cruises M42/3 and M44/2.  
226 Except for five data sets (Cornils and Schnack-Schiel, 2017; Cornils, [et al., 2017a, b, c, d](#)) all data sets were sorted  
227 and identified by Elke Mizdalski. Thus, the taxonomic concept has been used consistently throughout the data  
228 sets. A wide variety of identification keys and species descriptions have been used to identify the copepods, which  
229 cannot be all named here. References [for the species descriptions](#) and drawings of all [identified marine planktonic](#)  
230 species can be found at Razouls et al. (2005 – 1018). Calanoid copepods were identified to the lowest taxa possible,  
231 in general genus or species. Furthermore, of each identified taxon females, males and copepodite (juvenile) stages  
232 were separated. Cyclopoid copepods were identified to species level in four data sets (Cornils et al., 2017a, b, c,  
233 d).  
234 Previously published data sets were revised to ensure consistency of species names throughout the data set  
235 collection (Michels et al., 2012; Schnack-Schiel et al., 2007; Schnack-Schiel, 2010; Schnack-Schiel et al., 2010).  
236 In the present compilation we have used the currently acknowledged copepod taxonomy as published in  
237 WoRMS (World register of Marine Species (WoRMS Editorial Board, 2018)) and at Razouls et al. (2005 –  
238 2018). Species names have been linked to the WoRMS database, so future changes in taxonomy will be tracked.  
239 In the parameter comments the “old” names are archived that were used initially when the specimens were  
240 identified. All used species names can be found as “Copepod species list” under “Further details” at  
241 <https://doi.org/10.1594/PANGAEA.884619> or at <http://hdl.handle.net/10013/epic.65463ec2-e309-4d57-8fe3-0ceb7dd7dce70>. We provided also the unique identifier (Aphia ID) from WoRMS and notes on the distribution of  
242 each species.  
243

244  
245 When specimens could not be identified due to the lack of identification material, uncertainties in the taxonomy  
246 or missing parts they were summarized under the genus name (e.g. *Disco* spp., *Diaixis* spp., *Paracalanus* spp.,  
247 *Microcalanus* spp.) or family name (e.g. Aetideidae, copepodites). In most data sets few individuals could not be  
248 assigned to any family or genus. These are summarized as Calanoida [indeterminata](#), female, Calanoida  
249 [indeterminata](#), male and Calanoida [indeterminata](#), copepodites.  
250

### 251 3 Data sets

#### 252 3.1 Metadata

253 Each data set has its own persistent identifier. The metadata are consistent among all data sets, thus ensuring the  
254 comparability of the data sets and document their quality.

255 The following metadata can be found in each data set:

- 256 - “Related to:” includes the corresponding cruise report, related data sets and scientific articles [of Sigrid](#)  
257 [Schnack-Schiel and others](#) that have used part of the data previously.
- 258 - “Other version:” In a few cases we have revised a previously published version of the data to ensure  
259 consistent species names throughout all data sets (for more information see section 2.3).
- 260 - “Projects:” shows internal projects or those with external funding. In the present case all data sets are  
261 related to internal projects of the AWI (Alfred Wegener Institut Helmholtz Centre for Polar and Marine  
262 Research) research program.
- 263 - “Coverage:” gives the min/max values of the georeferences (latitude/longitude) of all stations.

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269 - "Event(s):" comprises a list of station labels, [a combination of cruise abbreviation and station number](#),  
 270 [Latitude/longitude of the position \(Units are in decimals with six decimal places\)](#), date/time of start and  
 271 end of station, [and elevation giving the bottom depth. Latitude/longitude, date/time and elevation were](#)  
 272 [all recorded by the systems of the respective scientific vessel](#). Campaign contains the cruise label  
 273 (including optional labels), basis is the name of the research vessel. Device contains the net type, which  
 274 was deployed and the comment may show further details of the station operation.

275 - "Parameter(s):" list of parameters used in the data set with columns containing the full and short name,  
 276 the unit, the PI (which in this data compilation is always Sigrid Schnack-Schiel, except for one data set  
 277 (<https://doi.org/10.1594/PANGAEA.880239>), and the method with a comment. The parameter  
 278 "Date/Time of event" is not always identical with "Date/Time" given in the event. This is the case when  
 279 the "Device" in the event is set to "Multiple Investigations" and thus the starting time of all investigations  
 280 at this event is given. "Date/Time of event" however, is the time when the plankton net haul started.  
 281 ["Date/Time" recorded on R/V Polarstern and during the cruises M42/3 and JB03 was UTC \(Coordinated](#)  
 282 [Universal Time\) and during cruise M44/2 local time was recorded \(UTC+2\). No information on](#)  
 283 ["Date/Time" was found for the cruises DAE1979/80, M11/4 and VH1094.](#)

284 "Elevation" provides information on the bottom depth of the plankton station, if available.  
 285 Three parameters describe the sampling depths interval. "Depth, water" is the mean depth of the sampled  
 286 depth interval. "Depth top" and "Depth bot" describe the upper and lower limit of the sampling depth  
 287 interval, respectively.

288 "Volume" is the amount of water that was filtered during each net tow, either calculated using the mouth  
 289 area of the net and depth interval or with a flowmeter (section 2.2.2). "Comment" gives the detailed  
 290 information on the net type, the net number and mesh size. In the following list of parameters are the  
 291 copepod taxa for which abundance data were recorded. Calanoid taxa are separated in female, male and  
 292 copepodites. Species names are consistent throughout all data sets, which ensures the comparability of  
 293 the data sets. [Clicking the link on the species names leads to their respective WoRMS ID](#) (see section 2.3).  
 294 The "short names" of each taxon consist of the first letter of the generic name and the name of the species.  
 295 In nine cases this results in identical short names (*Pleuromamma antarctica*, *Paraeuchaeta antarctica* =  
 296 *P. antarctica*; *Temoropia minor*, *Temorites minor* = *T. minor*; *Chiridius gracilis*, *Centropages gracilis* =  
 297 *C. gracilis*; *Clausocalanus minor*, *Calanopia minor* = *C. minor*; *Heterostylites longicornis*, *Haloptilus*  
 298 *longicornis* = *H. longicornis*; *Scolecithricella abyssalis*, *Spinocalanus abyssalis* = *S. abyssalis*;  
 299 *Scaphocalanus magnus*, *Spinocalanus magnus* = *S. magnus*). Thus, we advise to use the full scientific  
 300 names of these species in further analyses.

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### 302 3.2 Temporal station distribution

303 While samples of the Magellan region (November 1994), the Gulf of Aqaba and the northern Red Sea  
 304 (February/March 1999), Great Meteor Bank (September 1998) and Eastern Atlantic Ocean (November 2002) were  
 305 restricted to one year and one season, the Southern Ocean was sampled multiple times (Table 1). Samples in the  
 306 Southern Ocean were taken from 1980 to 2005 (Table 1, Fig. 2 a, b). The highest number of zooplankton samples  
 307 was taken in the 1980s (Fig. 2 b). In the 1980s the sampling effort was concentrated to the Antarctic Peninsula,  
 308 the Scotia Sea and the Weddell Sea (Fig. 2 a). Samples were taken in multiple years. In the 1990s until 2005 most  
 309 samples were taken in the Bellingshausen and Amundsen Sea, with fewer samples in the western and eastern

315 Weddell Sea. Two transects were sampled across the Weddell Sea in the 1990s in austral summer and autumn  
316 (Fig. 2 b). In general, most stations were sampled during summer (December to February), followed by autumn  
317 (March to May) and spring (September to November), while winter samples are only available from 1986 in the  
318 eastern Weddell Sea (Fig. 2 b, c). Summer and autumn samples are widely distributed from the Amundsen Sea to  
319 the eastern Weddell Sea (Fig. 2 b), while spring and autumn samples are mostly present from the Scotia Sea and  
320 Eastern Weddell Sea. Most samples were taken in January and February (Fig. 2 d). Samples are scattered  
321 throughout the entire day (Fig. 3.).

322 It should be taken into account that several copepod species in regions with pronounced seasonality of primary  
323 production, e.g. in high latitudes or upwelling regions (Conover, 1988; Schnack-Schiel, 2001) undergo seasonal  
324 vertical migration (e.g. *Rhincalanus*, *Calanoides*). They reside in deep water layers during period of food scarcity  
325 and rise to the surface layers when the phytoplankton blooms start. Furthermore, other species undergo pronounced  
326 diel vertical migrations (e.g. *Pleuromamma*) from mesopelagic layers during daytime to avoid predators to  
327 epipelagic waters at night to feed (Longhurst and Harrison 1989). Thus, to avoid biases in the comparison of the  
328 vertical distribution of copepod species season and daytime should be considered during further analysis of the  
329 data sets.  
330

### 331 3.3 Copepoda

332 In total, specimens from six copepod orders were recorded in the compiled data sets.  
333 However, in 29 data sets only calanoid copepods were identified on species level. Specimens of other copepod  
334 orders were comprised in families or orders.  
335

#### 336 3.3.1 Calanoida

337 In total 284 calanoid species could be separated in 29 data sets (see "Copepod species list" at  
338 <https://doi.pangaea.de/10.1594/PANGAEA.884619>). These species are representatives of 28 families and 91  
339 genera (Table 2). In the Southern Ocean abundance and distribution data for 96 calanoid species were archived.  
340 In the eastern Atlantic Ocean 125 and around the Great Meteor Bank 135 calanoid copepod species could be  
341 identified (Table 2). These numbers already indicate the well-known fact that species richness in the tropical and  
342 subtropical open oceans is much higher than in the polar Southern Ocean (e.g. Rutherford et al., 1999; Tittensor  
343 et al., 2010). Compared to these the number of calanoid species (60) in the subtropical northern Red Sea is low,  
344 which is expected due to the shallow sills at the entrance of the Red Sea and the high salinity (see Comils et al.  
345 2005). The lowest number of calanoid species (35) was found in the Magellan Region. Calanoid copepod families  
346 with the highest number of species were Aetideidae (33), Augaptilidae (27) and Scolecitrichidae (40; Table 2).

347  
348 For selected species from the Southern Ocean and the northern Red Sea and Gulf of Aqaba, also the five copepodite  
349 stages were counted individually (Table 3), providing valuable information on the seasonal and vertical  
350 distribution of the five copepodite stages. During four cruises, also *Rhincalanus gigas* nauplii were counted (PS09,  
351 PS21, PS23, PS29). In the 1990s Sigrid Schnack-Schiel has used these data to publish a series of papers on life  
352 cycle strategies of Antarctic calanoid copepods such as *Calanoides acutus*, *Rhincalanus gigas*, *Microcalanus* cf.  
353 *pygmaeus* or *Stephos longipes* (e.g. Schnack-Schiel and Mizdalski 1994, Schnack-Schiel et al. 1995, Ward et al.  
354 1997, Schnack-Schiel, 2001). However, the stage-resolved copepod data of most species in Table 3 have not been  
355 analyzed.

Gelöscht: All calanoid species were counted separately as females, males and copepodites.

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361 It is notable that none of the calanoid species were found in all five regions (see “Copepod species list” at  
362 <https://doi.pangaea.de/10.1594/PANGAEA.884619>). In contrast, many species were only recorded in one region:  
363 60 species were found only in the Southern Ocean, while 43 and 38 were found only in the data sets from the Great  
364 Meteor Bank and the transect in the eastern Atlantic Ocean, respectively. 24 species were found only in the Red  
365 Sea and six were identified only from samples in the Magellan region. Of the 28 calanoid families eleven were  
366 distributed in all five regions (Table 2).

Gelöscht: “Copepod species list”

367 As an example for the geographical and vertical distribution of the copepods three abundant genera were chosen  
368 (Fig. 4). While *Microcalanus* spp. (not separated in species due to uncertainties in the taxonomy) and *Spinocalanus*  
369 spp. (9 species; Table 2) are abundant down to 1000 m, the two species of *Ctenocalanus* (2 species, Fig. 4) and  
370 *Stephos* occur mainly in the epipelagic layer of the ocean. This is in accordance with their known vertical  
371 distribution (Schnack-Schiel and Mizdalski, 1994, Bode et al., 2018). Comparing the abundance of *Spinocalanus*  
372 and *Microcalanus* from all regions suggests that the abundance of these taxa is far greater in the Southern Ocean  
373 than in the warmer regions of the ocean. This picture however, has to be treated with caution, since the tropical  
374 Atlantic was only sampled in the upper 300 m of the water column and was thus too shallow for the meso- and  
375 bathypelagic genera (Bode et al., 2018).

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376 In the case of *Ctenocalanus* and *Stephos* our data sets reveal that closely related species within a genus may have  
377 contrasting distribution patterns. *Stephos longipes* and *Ctenocalanus citer* are restricted to colder and polar waters  
378 of the southern hemisphere, while *Ctenocalanus vanus* occurs in both the Red Sea and the warm Atlantic Ocean.  
379 *Stephos maculosus* occurs only in the Red Sea (see arrow in Fig. 4). Furthermore, the distribution patterns reveal  
380 that of the four genera only *C. citer* has a higher abundance in the samples from the Bellingshausen and Amundsen  
381 Seas, and around the Antarctic Peninsula, while *S. longipes*, *Microcalanus* spp. and *Spinocalanus* spp. all have  
382 higher abundances in the Eastern Weddell Sea. This may be due to the lower water depth at the Peninsula since  
383 *Microcalanus* and *Spinocalanus* are considered as mesopelagic to bathypelagic. Thus, they are often not found at  
384 shallow stations (< 300 m depth). In case of the sea ice-associated *S. longipes*, low sea-ice conditions and offshore  
385 stations may have caused the restricted distribution. *S. longipes* occurred mainly in the upper water layers, but it  
386 was also recorded with low abundances in deeper layers (Fig. 4). This pattern may be due to its life cycle, shifting  
387 seasonally from a sea-ice associated to a benthopelagic life cycle (Schnack-Schiel et al., 1995).

### 389 3.3.2 Other Copepoda

390 In total, 28 non-calanoid taxa were recorded. Four data sets provide only abundance and distribution data for non-  
391 calanoid copepod orders (PS06, PS10, PS29, PS35; Table 1), in particular on species of the order Cyclopoida from  
392 the families Oithonidae (2 species) and Oncaeidae (6 species; Table 2). They were separated in female, male,  
393 copepodite stages 1, 2, 3, 4, and 5. During VH1094 also *Oithona* species were identified (Table 2). In all other  
394 data sets species of these two families were not separated. In all regions representatives of the family Lubbockiidae  
395 were recorded. In the subtropical and tropical samples of PS63, M44/2 and M42/3 also abundances of species of  
396 the families Corycaeidae and Sapphirinidae, and of the genus *Pachos* were recorded. Except for PS65, species of  
397 the order Harpacticoida were not separated. In the latter five species were identified, mainly sea-ice associated  
398 harpacticoids (Table 2; Schnack-Schiel et al., 1998). Also, specimens of the orders Monstrilloida, Mormonillida  
399 and Siphonostomatoida were counted.

400 In most data sets, copepod nauplii are also recorded as one parameter. However, due to the small size of nauplii  
401 they were not sampled quantitatively and should be discarded in further analysis.

405

### 406 3.4 Further remarks on the usage of the data compilation

407 Generally, the cruise reports have been linked to each data set. The cruise reports provide valuable information on  
408 the itinerary, zooplankton sampling procedures and on other scientific activities on-board that could be useful for  
409 the data analysis (e.g. CTD data). Abundance data of selected species and data sets have been published previously  
410 in scientific articles. These articles are linked to the respective data sets (under “Related to”).

Gelöscht: We have also added scientific article that are related to individual data sets.

411 To use the data, they can be downloaded individually as tab-delimited text files or altogether as a .zip file to allow  
412 an import to other software e.g. R (R core team, 2018) or Ocean Data View (Schlitzer, 2015) for further analysis.

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413 Due to the consistent taxonomic nomenclature the individual files can be concatenated easily. It should be kept in  
414 mind however, that not all data sets are directly comparable due to difference in net type and mesh sizes (see  
415 section 2.2.4). As noted in section 3.2 several species undergo pronounced seasonal and diel vertical migrations.  
416 Therefore, nets from surface waters may not always sample the full vertical extent of the populations, particularly  
417 of the biomass dominants.

418 To evaluate the vertical and spatial distribution of marine plankton hydrographic information such as temperature  
419 and salinity profiles are essential. The relevant publications are available at  
420 <https://doi.org/10.1594/PANGAEA.884619>, see “Further details”. Recently, a summary of the physical  
421 oceanography of R/V Polarstern has been published (Driemel et al., 2017) with CTD data archived in PANGAEA  
422 as well (Rohardt et al., 2016), except for the cruises PS04 (ANT-II/2), PS14 (ANT-VII/2), PS21 (ANT-X/3), PS63  
423 (ANT-XX/1) and PS65 (ANT-XXII/2) (see Table 1). For these five cruises information on temperature and salinity  
424 profiles exist only for PS63 (Schnack-Schiel et al., 2010) and for PS65 the CTD profiles can be downloaded  
425 (<https://doi.org/10.1594/PANGAEA.742627>; Absy et al., 2008). For M11/4 a CTD data set is also available  
426 (<https://doi.org/10.1594/PANGAEA.742745>; Stein, 2010). To connect the CTD data with the corresponding  
427 plankton net haul the metadata “Event“ and “Date/time“ can be used. Furthermore, cruise track and station  
428 information are available in the cruise reports as well as on the station tracks for each cruise  
429 (<https://pangaea.de/expeditions/>). For the other two R/V Meteor cruises hydrographic information is available in  
430 scientific articles (M42/3: Beckmann and Mohn, 2002; Mohn and Beckmann, 2002; M44/2: Cornils et al., 2005;  
431 Plähn et al., 2002). Metadata information of the cruise JB03 can be downloaded from:  
432 [https://www.bodc.ac.uk/resources/inventories/cruise\\_inventory/report/5916/](https://www.bodc.ac.uk/resources/inventories/cruise_inventory/report/5916/). To date, no hydrographic  
433 information is publicly available for the cruises DAE79/80 and VH1094.

Gelöscht: In these cases we recommend to use only presences and absences of the species. ¶

434 Additionally, abundances of all other zooplankton organisms in the net samples used for the copepod datasets are  
435 available for the four cruises ANT-X/3, ANT-XVIII/5b, M42/3 and M44/2. These can be downloaded at  
436 <https://doi.org/10.1594/PANGAEA.883833>, <https://doi.org/10.1594/PANGAEA.884581>,  
437 <https://doi.org/10.1594/PANGAEA.883771> and <https://doi.org/10.1594/PANGAEA.883779>.

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438 All data presented here are archived in the database PANGAEA. There are however, other data archiving initiatives  
439 that also store data on copepod abundance and distribution such as COPEPOD  
440 (<https://www.st.nmfs.noaa.gov/copepod/>), BODC (<https://www.bodc.ac.uk>) or OBIS (<http://www.iobis.org>). The  
441 here presented data however, have not been published in any other cataloguing initiative before.

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### 442 4 Data availability

443 In total 33 data sets with 349 stations were archived in the PANGAEA® (Data Publisher for Earth &  
444 Environmental Science, [www.pangaea.de](http://www.pangaea.de)) database. The persistent identifier  
445

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457 <https://doi.org/10.1594/PANGAEA.884619> links to the splash page of the data compilation. We encourage the  
458 users of these data to cite both the DOI of the data collection in PANGAEA as well as the present  
459 data publication as a courtesy to Dr. Sigrid Schnack-Schiel and the people preparing the data for  
460 Open Access. Metadata include DOIs to cruise reports and related physical oceanography. Data are provided in  
461 consistent format as tab-delimited ASCII-files and are in Open Access under a CC-by license (Creative Commons  
462 Attribution 3.0 Unported).

## 463 **5 Concluding remarks**

465 Pelagic marine ecosystems are threatened by increasing water temperatures due to climate change. These  
466 environmental changes are expected to cause also shifts in the community structure of pelagic organisms. Within  
467 the pelagic food web copepods have a central role as mediator between the microbial loop and higher trophic  
468 level. Due to their short life cycles and their high diversity copepods offer a unique opportunity to study effects of  
469 environmental variables on numerous taxa with different life cycle strategies. It is also known that their species  
470 composition and abundance often reflect environmental changes such as temperature, seasonal variability or  
471 stratification (Beaugrand et al., 2002). To understand the complexity of ecological niches and ecosystem  
472 functioning, but also to investigate the effects of environmental changes a detailed knowledge of species diversity,  
473 distribution and abundance is essential. The present data compilation provides further information on spatial,  
474 vertical and temporal distribution of copepod species and may thus be used to obtain a better picture of species  
475 biogeographies. Many individual data sets can also be linked to corresponding CTD profiles (Table 1) and may  
476 thus be useful for modeling approaches such as species distribution or environmental niche modeling.

477 Furthermore, for all calanoid copepods females, males and copepodites were enumerated separately and for  
478 selected species even between copepodite stages was discriminated. This detailed resolution of abundance data  
479 will also allow future investigations on life cycle strategies and also how the different stages interact with the  
480 environment (e.g. temperature, currents, depth).

## 481 **Competing interests**

482 The authors declare that they have no conflict of interest.

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[1] nach oben verschoben: This data compilation represents the scientific legacy of Dr. Sigrid B. Schnack-Schiel (1946-2016), revealing her expertise and great interest in polar zooplankton ecology, but also exploring the zooplankton communities of tropical and subtropical regions. ¶

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