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Biogeography of key mesozooplankton species in the North Atlantic, by manual counting methods, and egg production of *Calanus finmarchicus*

W. Melle¹, J. A. Runge², E. Head³, S. Plourde⁴, C. Castellani⁵, P. Licandro⁵, J. Pierson⁶, S. H. Jonasdottir⁷, C. Johnson³, C. Broms¹, H. Debes⁸, T. Falkenhaus¹, E. Gaard⁸, A. Gislason⁹, M. R. Heath¹⁰, B. Niehoff¹¹, T. G. Nielsen⁷, P. Pepin¹², E. K. Stenevik¹, and G. Chust¹³

¹Institute of Marine Research, Research Group Plankton, P.O. Box 1870, 5817 Nordnes, Bergen, Norway

²School of Marine Sciences, University of Maine, Gulf of Maine Research Institute, 350 Commercial Street, Portland, ME 04101, USA

³Fisheries and Oceans Canada, Bedford Institute of Oceanography, P.O. Box 1006, Dartmouth, NS, B2Y 4A2, Canada

⁴Pêches et Océans Canada, Direction des Sciences océaniques et Environnementales, Institut Maurice-Lamontagne, 850 route de la Mer, C.P. 1000 Mont-Joli, QC, G5H 3Z4, Canada

⁵Sir Alister Hardy Foundation for Ocean Science (SAHFOS), Citadel Hill, Plymouth, PL1 2PB, UK

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⁶Horn Point Laboratory, University of Maryland Center for Environmental Science, 2020 Horns Point Road, Cambridge, MD 21613, USA

⁷National Institute for Aquatic Resources, Technical University of Denmark, Jægersborgs Allé 1, 2920 Charlottenlund, Denmark

⁸Faroe Marine Research Institute, Box 3051, FO-110 Torshavn, Faroe Islands

⁹Marine Research Institute, Skulagata 4, P.O. Box 1390, 121 Reykjavik, Iceland

¹⁰MASTS Marine Population Modeling Group, Department of Mathematics and Statistics, University of Strathclyde, Livingstone Tower, 26 Richmond Street, Glasgow, G1 1XH, Scotland

¹¹Alfred Wegener Institute for Polar and Marine Research, Polar Biological Oceanography, 27570 Bremerhaven, Germany

¹²Northwest Atlantic Fisheries Centre, Fisheries and Oceans Canada, P.O. Box 5667, St. John's, Newfoundland A1C 5X1, Canada

¹³AZTI-Tecnalia, Marine Research Division, Txatxarramendi ugartea, 48395 Sukarrieta, Spain

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Correspondence to: W. Melle (webjorn@imr.no)

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2.2 Mapping of key species with CPR

The CPR survey is an upper layer plankton monitoring program that has regularly collected samples, at monthly intervals, in the North Atlantic and adjacent seas since 1946 (Warner and Hays, 1994). Water from approximately 6 m depth (Batten et al., 2003a) enters the CPR through a small aperture at the front of the sampler and travels down a tunnel where it passes through a silk filtering mesh of 270 μm before exiting at the back of the CPR. The plankton filtered on the silk is analyzed in sections corresponding to 10 nautical miles (approx. 3 m^3 of seawater filtered) and the plankton microscopically identified (Jonas et al., 2004). In the current ESSD we present CPR data that represent basin scale distributions of *C. finmarchicus* (CV-CVI), *C. helgolandicus* (CV-CVI), *C. hyperboreus* (CV-CVI), *Pseudocalanus* spp. (CVI), *Oithona* spp. (CI-CVI), total euphausiida, total pteropoda and the presence/absence of Cnidaria (Fig. 2). Monthly data collected between 2000 and 2009 were gridded using the inverse-distance interpolation method (Isaaks and Srivastava, 1989), in which the interpolated values were the nodes of a 2° by 2° grid. The resulting twelve monthly matrices were then averaged within the year and the data log-transformed (i.e. $\log_{10}(x + 1)$). The Phytoplankton Colour Index (PCI), which is a visual assessment of the greenness of the silk, is used as an indicator of the distribution of total phytoplankton biomass across the Atlantic basin (Batten et al., 2003b; Richardson et al., 2006). After comparing distribution of *Calanus finmarchicus* by CPR and vertical net sampling Melle et al. (2014) concluded that maximum *C. finmarchicus* abundances are found in the deep basins of the Norwegian and Labrador seas to some extent north of the CPR sampling routes. For this reason, since 2008, the spatial coverage of CPR monitoring has been expanded to cover the core areas of *C. finmarchicus* distribution in the Norwegian Sea. These data are not included in the present ESSD.

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2.3 Seasonal dynamics and demography of *Calanus finmarchicus* by net sampling

Seasonal abundances and demography of *Calanus finmarchicus* were derived from samples taken at sites across the North Atlantic (Table 1, Fig. 1b). The sampling sites include both coastal and oceanic stations and vary from relatively cold to warm water locations (Fig. 1a). Sampling frequency also differs among sites; the more easily accessed coastal sites were generally visited more frequently than the offshore sites. An overview of sampling sites characteristics, sampling gear and methods is provided in Table 1. At all sites abundances of developmental stages were averaged over 14 days periods within the year and then for the same periods over all years.

2.4 *Calanus finmarchicus* egg production and female size

Observations of egg production rates (EPR) for female *Calanus finmarchicus* were compared for different regions of the North Atlantic (Fig. 1c). The regions were diverse in size and sampling frequency, ranging from a fixed time series station in the Lower St Lawrence Estuary, off Rimouski (RIM), where nearly 200 experiments were carried out between May and December from 1994 to 2006 (RIM), to a large-scale survey in the Northern Norwegian Sea (NNWS), where about 50 experiments were carried out between April and June from 2002 to 2004 (NNWS). For this compilation the stations were grouped mostly along geographic lines, with only limited attention being paid to oceanographic features. There is some overlap between regions, however, where stations were sometimes kept together when they were sampled on the same cruise. As well, although not shown in Fig. 1c, some stations other than RIM were occupied more than once during different years and/or in different seasons. Some of the data included here have appeared in published papers and the citations are included. Previously unpublished data were also provided by C. Broms, E. Gaard, A. Gislason, E. Head and S. Jónasdóttir. Data have been submitted to PANGEA as averages by area.

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Egg production in *C. finmarchicus* occurs in spawning bouts, which are of relatively short duration and may occur once or more per day (Marshall and Orr, 1972; Hirche, 1996). While there is evidence for diel spawning periodicity in the sea (Runge, 1987; Runge and de Lafontaine, 1996), females incubated in dishes for the first 24 h after capture do not always show a consistent night time release of eggs, as they did for *Calanus pacificus* (Runge and Plourde, 1996; Head et al., 2014). Because of the potential for diel egg-laying behaviour, the vast majority of egg production experiments have been carried out by incubating freshly caught females for 24 h. It has been shown that female *Calanus* that are kept and fed in vitro and then transferred to an incubation chamber lay the same number of eggs over the next 24 h whether or not they are fed (Plourde and Runge, 1993; Laabir et al., 1995). Thus, it has been assumed that average egg production rates of freshly caught females are the same during the 24 h following capture as they would have been in situ (Runge and Roff, 2000). In this study we include only results from such 24 h incubation experiments, and we term the eggs laid during these 24 h periods “clutches”, even though they may originate from more than one spawning bout, and the number of eggs laid by one female during a 24 h period as the clutch size (CS). In most experiments 20–30 females were incubated individually in separate chambers, and the proportion of females that laid eggs over 24 h is referred to as the “spawning frequency” (SF), which is here expressed as a percentage per day. Egg production rates (EPRs) reported here were calculated by individual contributing investigators either simply as the sum of all of the eggs produced in an experiment divided by the number of females incubated and the average incubation time (generally 1 day), or as the average of the EPRs calculated for each experimental female individually, which takes account of differences in incubation times for individual females. For the WGBB most experiments were carried out using prolonged incubation periods (e.g. 36–48 h), often with relatively few females (~ 10). For several of the analyses carried out here it was necessary to include the results of these prolonged incubations.

As batches of eggs are released into the water column in situ, they may hatch and develop, or they may be consumed by local predators, including female *C. finmarchicus* themselves, which are sometimes the most abundant potential predators (Basedow and Tande, 2006). To avoid cannibalism, incubations are generally set up so as to minimize contact between the females and the eggs they are laying. This has been done by the investigators contributing to this work using one of five techniques. In Method A females are incubated individually in 45–50 mL of seawater in 6–10 cm diameter petri dishes. The eggs sink rapidly to the bottom surface, where they are unlikely to be caught up in the females’ feeding currents. Method B involves incubating females individually in similar but smaller “Multi-well” chambers, which have a volume capacity of 10–15 mL. In Method C females are placed individually (or in groups of 2 or 3) in cylinders, fitted with mesh screens on the bottom, which are suspended in beakers of 400–600 mL capacity (Gislason, 2005). The eggs sink through the mesh and are thus separated from the females. Method D represents a modification of Method C, in which there is flow of seawater through the chamber (White and Roman, 1992). Finally, in Method E, individual (or groups of 2 or 3) females are incubated in bottles or beakers (up to 1 L capacity), without screening (Jónasdóttir et al., 2005). For Method E the vessels are kept upright and it is assumed that the eggs will sink out and become unavailable to the females relatively rapidly.

There have been relatively few comparisons of these different experimental methods. Cabal et al. (1997) found that female *C. finmarchicus* from the Labrador Sea incubated individually in 50 mL petri dishes (Method A) or 80 mL bottles (Method E) produced similar numbers of eggs after 3 days, although only three experiments were done and over the first 24 h CSs were larger for Method A. They also found that over 24 to 72 h periods groups of females in screened cylinders within large volume chambers (Method C) gave higher egg production rates than did those in chambers without screens (Method E). Runge and Roff (2000) reported egg laying in dishes (Method A) yielded similar egg production rates to those for groups of 10–15 females incubated in 1.5 L screened beakers (Method C). However, the beaker egg production estimates declined dramati-

cally relative to dish estimates in rough weather, presumably due to increased mixing in beakers and therefore higher loss due to cannibalism. More recently, Plourde and Joly (2008) found that suspending a mesh screen within petri dishes 2 mm above the bottom made no difference to the number of eggs produced by female *C. finmarchicus* over 24 h, although it did increase the number of eggs recovered from *Metridia longa* females, which could be seen swimming actively and sweeping the bottom with their mouthparts in the unscreened dishes. In the Northeast Atlantic, at Ocean Weather Station M (included in our Southern Norwegian Sea (SNWS) region), B. Niehoff (personal communication) found that females incubated for 24 h in Multi-wells (Method B) had similar CSs to those incubated according to Method C. None of these studies compared all methods and the fact that the NW Atlantic groups have used Method A, while the central and NE Atlantic groups have used mainly Methods C, D or E introduces a question as to whether methodological differences might have contributed to the differences found among the CSs and EPRs in the different regions. Such an analysis is not possible based on the data currently available, however, and the topic will not be considered further in this work, although it merits further attention.

Another point on which investigators differed is how they dealt with small clutches. For the Georges Bank (GB), Rimouski station (RIM) and Scotian Shelf (SS) regions and for the Labrador Sea (LS) data provided by R. Campbell, clutches of < 6 eggs were routinely not included in the datasets on CSs, since they were regarded as being the result of interrupted spawning events. These small clutches were apparently very rare (J. Runge, personal communication) and indeed for the LS data reported by Head et al. (2013) clutches of < 6 eggs accounted for only 32 of the 1324 clutches observed, i.e. 2%. For regions farther east, however, the proportions of clutches of < 6 eggs were generally larger, between 13 % (SNWS) and 33 % (Northern Norwegian Sea, NNWS). Because of this difference in data reporting, CSs of < 6 eggs were excluded from the calculations of average CSs for all regions. Small clutches were, however, included by all investigators in their calculations of EPRs.

Previous studies of egg production have shown a significant link between clutch size and female size (Runge and Plourde, 1996; Campbell and Head, 2000; Jónasdóttir et al., 2005; Runge et al., 2006) and most of the datasets provided for this work included measurements of the prosome lengths for each individually incubated female for each egg production experiment, along with each corresponding individual clutch size (Table 3). One exception to this was in the SNWS region (data from Ocean Weather Station M), for which average female prosome lengths were determined for groups of females that had not been used in experiments, but that had been collected on the same day. In addition, there were no measurements of prosome lengths for some data from the region “Between Scotland and Iceland” (BIS) and the SNWS and NNWS regions. As well, prosome lengths were not measured for all clutch sizes enumerated at RIM.

Egg production rates for the experiments carried out within a given region were averaged seasonally. The rationale for the grouping of months into seasons within each region was based partly on observations of seasonal cycles of temperature and chlorophyll concentration, partly on what could be ascertained from the literature about the timing of the appearance of females at the surface after over-wintering, and partly on the availability of data. The spring months cover the period when water temperatures are increasing, when the spring bloom is starting or is in progress, when diatoms dominate the female diet and when the overwintered (G0) generation of females is abundant in the surface layers. Spring is the time when community egg production rates, although maybe not individual rates, are expected to be highest. In summer, temperatures are higher, the bloom may still be in progress, but the female diet may be more varied, and some females of the new year’s generation may be present. In autumn and winter relatively few females are in the near surface layers and phytoplankton levels are generally low.

Observations of in situ temperature and chlorophyll concentration were made at nearly all experimental stations. The original aim had been to use in situ temperatures from 5 m and chlorophyll concentrations integrated to 30 m in this study. Not all

data were provided in this form, however. For example, in some datasets temperature data were surface values or 0–10 m or 0–20 m averages and chlorophyll concentrations were sometimes values integrated to 50 m. The data were standardized to a comparable format by assuming that surface, 0–10 m or 0–20 m average temperatures were the same as 5 m temperatures, and that the chlorophyll concentrations were uniform throughout the 0–50 m depth range. These assumptions are likely to be most appropriate in spring and winter, when mixed layers are relatively deep.

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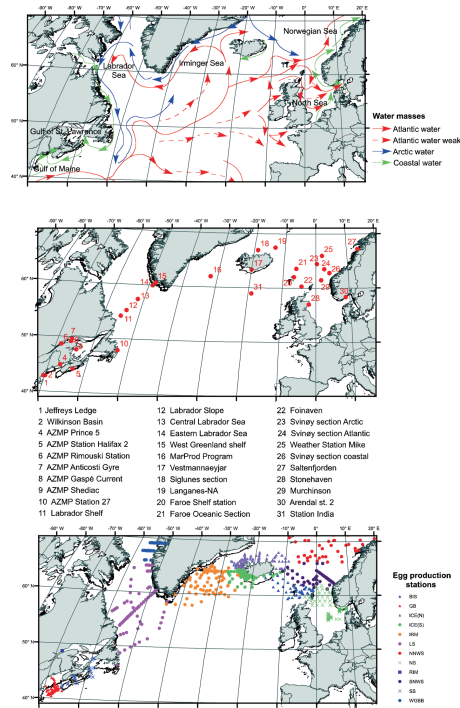


Fig. 1. (Upper panel) The northern North Atlantic Ocean, major warm and cold water currents and important seas. (Mid panel) Locations of demographic stations and transects listed in Table 1. (Lower panel) Locations of observations of *C. finmarchicus* egg production rates (and usually adult body size, chlorophyll *a* concentrations and temperature).

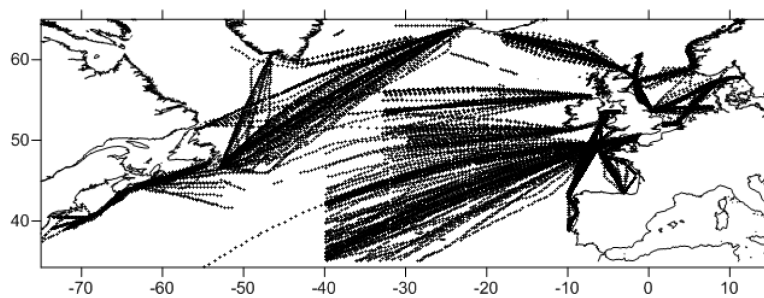


Fig. 2. CPR data sampling routes, 2000–2009.