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# A century of change in North Sea plankton communities explored through integrating historical datasets

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## Abstract

Plankton communities make useful ecosystem indicators, and taking a historical perspective on plankton community composition provides insights into large-scale environmental change. Much of our understanding of long-temporal scale change in plankton communities in the North Sea has been provided by the Continuous Plankton Recorder (CPR) survey, operating since 1931, with consistent time-series data available since 1958. This paper further increases the temporal scale of our understanding of community change in the North Sea by combining the CPR dataset with a digitised collection of plankton surveys undertaken by ICES from 1902 to 1912. After steps taken to integrate the two disparate datasets, differences in overall community composition between time-periods suggest that the multidecadal changes observed through the CPR survey time-period may have occurred from a non-stable baseline that was already on a trajectory of change. Therefore, a stable historical time period in which plankton communities are assessed against for any impact of human pressures may be hard to define for the North Sea and instead underlying variation needs to be encompassed within any baseline chosen. Further evidence for the influence of large scale changes in SST driving change in plankton community composition was found using the extended dataset.

## 1 Introduction

Climate change is causing widespread changes in marine ecosystems, superimposed on a background of climate variability that acts at different temporal scales (Hoegh-Guldberg and Bruno, 2010). Plankton communities are sensitive to changes in the physical marine environment, and have been shown to be responsive to interannual and multi-decadal climate variability as well as anthropogenic climate change (Hays et al., 2005). As the base of the pelagic food web, phytoplankton are primary producers (Boyce and Worm, 2015), transferring energy through zooplankton to higher trophic levels (Richardson, 2008). This sensitivity to environmental conditions and their role in the pelagic foodweb makes tracking plankton community change useful as an indicator of change in the wider ecosystem. Much of our understanding of multi-decadal change in plankton communities in the North Sea comes from the Continuous Plankton Recorder (CPR) survey (McQuatters-Gollop et al., 2015). Consistent monitoring data available from 1958 through the present has documented widespread shifts in both phytoplankton and zooplankton communities, specifically the occurrence of basin-scale regime shifts in the North Atlantic (Beaugrand et al., 2014; Reid et al., 2015).

The value of plankton time series as evidence for policy and management increases with time. Through using long temporal scale data, the influence of multi-decadal changes in environmental conditions on plankton communities can be investigated, and the most important environmental influences structuring plankton communities on this scale can be identified (Edwards et al., 2010; Giron-Nava et al., 2017). For example, the Atlantic Multidecadal Oscillation is a term for the natural low-frequency SST variability in the North Atlantic that oscillates between warm and cool phases on a ~60yr time scale (Edwards et al., 2013). It has been identified as the second largest macroecological signal in North Atlantic plankton communities, but requires long temporal-scale time-series in order to detect the influence of transitions between oscillatory phases on community change (Edwards et al., 2013). Furthermore, the long temporal scale of the CPR survey can help separate these wider oceanographic and climatic influences on plankton communities, such as the influence of SST, from direct anthropogenic pressures such as eutrophication, which is particularly useful during formal policy assessments (McQuatters-Gollop et al., 2015).

'Rescuing' historical ecological datasets, that otherwise may be lost or deemed redundant, has been identified as a useful way of increasing temporal scale in ecological studies, and can be used to address contemporary marine

54 policy challenges, including understanding effects of long-term climate change (Hawkins et al., 2013). Specifically,  
55 the use of rescued historical datasets in avoiding ‘shifting baselines syndrome’ in biodiversity state has received  
56 much attention (Pauly, 1995). This is the phenomenon where neglecting historical changes obscures the magnitude  
57 of change or variability in ecosystem components. Therefore, rescued historical plankton data can be a tool for  
58 avoiding shifting baseline syndrome in our understanding of the multi-decadal dynamics of plankton communities  
59 (Ward et al. 2008). The ICES historical plankton dataset used in this study is a dataset of plankton samples collected  
60 in the North Atlantic between 1902 and 1912, digitised from historical log books. Hällfors et al. (2013) compared  
61 phytoplankton records from this ‘rescued’ ICES historical dataset in the Baltic Sea with contemporary phytoplankton  
62 samples, and documented compositional differences between the two time periods, potentially driven by both  
63 climate change and eutrophication. By comparing the ICES historical dataset with North Sea data from the CPR  
64 survey, we can better understand changes occurring in North Sea plankton communities pre-1950s, facilitating  
65 further exploration of the effects of large scale temperature change to the Continuous Plankton Recorder temporal  
66 coverage.

67 Disparities in sampling and analysis methodologies between the ICES historical data and the CPR survey, however,  
68 present challenges in their direct comparison, which need to be addressed before using the datasets together.

69 Handling disparate data-types is a key challenge facing regional scale monitoring and assessment where data from  
70 multiple different sampling programmes often needs to be incorporated (Olli et al., 2013; Zingone et al., 2015). For  
71 example, the OSPAR IA2017 regional-scale assessment of plankton communities incorporated multiple time-series  
72 from across Europe, where taxa were sampled using different methods, and analysed to differing taxonomic  
73 resolutions (OSPAR, 2017). In this study, by integrating and combining the CPR historical time series with the rescued  
74 ICES historical dataset, we aim to provide additional contextual information to the changes in North Sea plankton  
75 communities between 1958 and 2015 detected by the CPR survey, specifically to address the following questions:

- 76 • **Is there a difference in plankton community structure (both phytoplankton and zooplankton) between the**  
77 **early 20<sup>th</sup> century and the beginning of the consistently-sampled CPR time period (1960s)?**
- 78 • **Which plankton communities and individual taxa are most responsive to SST when examining the two**  
79 **datasets combined (1902-12, 1958-2015)?**

80

## 81 **2 Data and Methods**

### 82 **2.1 Data sources**

83

#### 84 **2.1.1 Plankton samples**

85 Data from the period 1902-1912 have become available through the ICES historic plankton digitisation project where  
86 13,379 plankton samples have been digitised from seven historical ICES volumes (McQuatters-Gollop et al. 2011).  
87 The data are collated from different sampling programmes, across the North-East Atlantic, North Sea, Irish Sea,  
88 Baltic Sea and Arctic Sea. After digitisation, data tables from the historical volumes were quality checked. The  
89 samples are all spatially referenced and consist of records of taxa at the presence/ absence level or with semi-  
90 quantitative abundance information. In this study, we used all data at the presence/absence level, as to be able to  
91 compare with the Continuous Plankton Recorder survey data. We extracted data from the months February, May,  
92 August and November, as these had the greater numbers of samples. This historical plankton dataset is now freely  
93 available via the ICES data portal ([ecosystemdata.ices.dk/HistoricalPlankton/Download.aspx](https://ecosystemdata.ices.dk/HistoricalPlankton/Download.aspx)).

94 The Continuous Plankton Recorder survey has been collecting samples in the North Sea on a routine, consistent basis  
95 since 1958 (Kirby and Beaugrand, 2009). CPRs consist of a filtering mechanism housed in an external body that is  
96 towed behind ships of opportunity at a depth of approximately 6-7m. The speed at which the silk is drawn from a  
97 storage spool is controlled by a propeller, with 10.16 cm of silk corresponding to 18.5 km of tow through the sea  
98 (Batten et al., 2003). CPR data for the months February, May, August and November were obtained for the North  
99 Sea area for phytoplankton (DOI 10.7487/2016.236.1.999) and zooplankton (DOI 10.7487/2016.236.1.998). Although  
100 abundance information is collected for each taxon identified on each sample, for this study data were converted to  
101 presence/absence to make comparable to the ICES historical database.

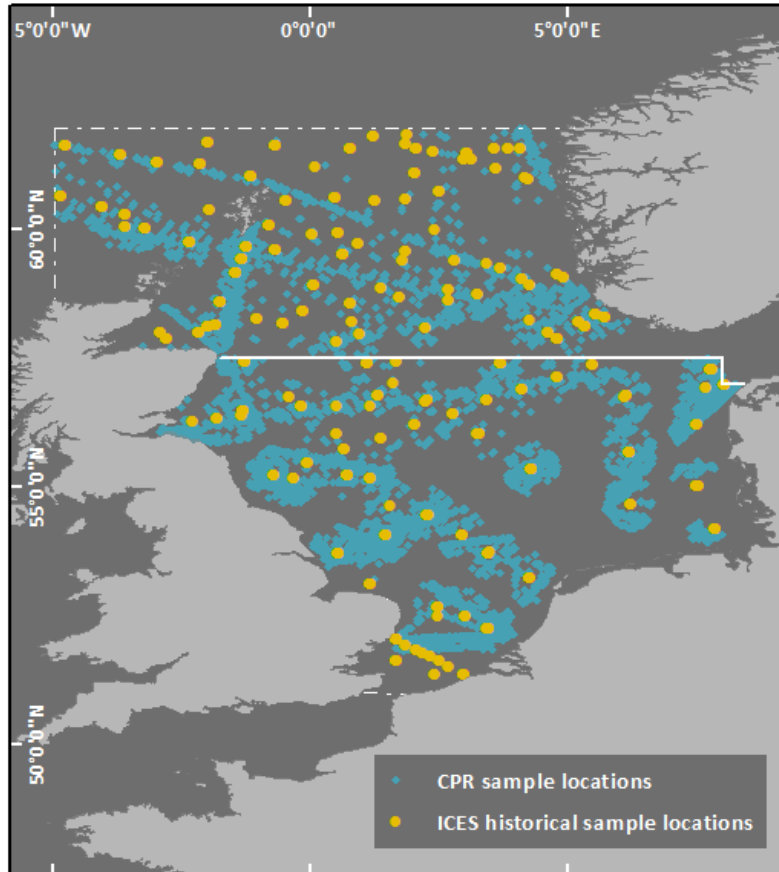
102 As well as differences in quantitative resolution between the datasets, there are major structural differences  
103 between the historical ICES surveys and the CPR survey (McQuatters-Gollop et al. 2010). Firstly, the CPR is a  
104 continuous plankton sampling method, using a 270 micron mesh size silk (Richardson et al., 2006). The ICES database,  
105 in contrast, consists of net samples, , collected at fixed point locations by a multitude of disparate sampling cruises  
106 by northern European nations. Therefore, whereas the analysis methodology has remained consistent throughout

107 the CPR series, the composite nature of the ICES dataset means that the sampling and analysis methodologies are  
108 not reliably consistent throughout the database. However, both sampling methodologies incorporated sub-sampling,  
109 where only a proportion of the sample is analysed, reducing any differences as a result of volume of water filtered  
110 (e.g. Hällfors et al. 2012).

111 The mesh sizes of the net samples in the ICES historical database are missing from the sample metadata, and are  
112 likely to be varied. The mesh size of the Continuous Plankton Recorder, 270 microns, is larger than the majority of  
113 standard plankton nets, which tend to range between 5 $\mu$ m - 80  $\mu$ m for phytoplankton and 125  $\mu$ m – 200  $\mu$ m for  
114 zooplankton (John et al., 2001; Castellani and Edwards, 2017) . Importantly therefore, any biases in sampling as a  
115 result of mesh size differences between the ICES historical plankton dataset and the CPR data are likely to come  
116 from the side of the CPR survey, evidenced by a lower number of species recorded overall than the ICES historical  
117 dataset. For example, CPR methodology likely undersamples smaller phytoplankton taxa, although they often are  
118 retained on the silk strands of the mesh (taxa as small as 5-10  $\mu$ m are regularly recorded), which constitutes 30-40%  
119 of the mesh area (Batten et al., 2003) Similarly, the CPR survey likely undersamples small zooplankton taxa. A  
120 previous study however, comparing CPR data to net samples taken at the L4 sampling station in the Western English  
121 channel, that used a mesh size of 200 microns, concluded that although the abundance of zooplankton taxa were  
122 generally lower, all dominant zooplankton species recorded at L4 were also common to CPR data (John et al., 2001).  
123 In this study, occurrence frequencies of select plankton taxa, based on presence/absence resolution data, were  
124 compared between datasets.

125 Samples from both datasets located in the North Sea region were divided into a 'Northern' North Sea region and a  
126 'Central/Southern' North Sea region based on the border between ICES regions 4b and 4c (Figure 1). The two spatial  
127 areas represent a balance between the need for spatial specificity in comparing plankton communities with known  
128 differences occurring across latitudes, and the retention of a reliable sample size within each area. To ensure the  
129 depth of the ICES samples were comparable to the CPR dataset all ICES historical samples collected below 15m, or  
130 vertical hauls that started below 15m were removed from the resulting sample list, along with samples for which no  
131 depth information was given. To compare plankton communities from the same area, CPR samples within half a  
132 degree of the ICES historical sample locations were then selected.

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143 **Figure 1.** Location of historical samples (large yellow) and centre points of CPR samples (small blue), included in the  
144 study. North Sea area (dashed white line) divided into 'Northern' and 'Central/Southern' areas based on the boundary  
145 between ICES subregions 4b and 4c (solid white line).

146

### 147 2.1.2 Sea surface temperature (SST) data

148 Monthly SST data were downloaded for the North Sea region from the International Comprehensive Ocean  
149 Atmosphere Dataset (ICOADS) at a 2 degree resolution. Data points were extracted from the Northern and  
150 Central/Southern North Sea area, and averaged for each year between 1902 and 2015.

151

## 152 2.2 Data preparation

153 Taxa lists of both phytoplankton and zooplankton were extracted from the historical ICES and CPR databases and  
154 both the ICES taxa lists and the CPR taxa lists were run through the Taxon Match Tool available on the WoRMS

(World Register of Marine Species) website (<http://www.marinespecies.org>) to update all names to the most up-to-date accepted nomenclature. Due to the ICES database being a composite of multiple sampling programmes, sporadically occurring taxa were removed, as these may not have been recorded or identified inconsistently between the different sampling programmes. For both datasets, a threshold of 1% frequency of occurrence was selected as a cut-off point for taxa to include in analyses of taxonomic composition. This represented a balance between the need to remove sporadic taxa, as highlighted by Hällfors et al. (2013), but still include rare species in analyses. Because of the decade time-span of the ICES historical dataset, this list for the CPR data was constructed based on a 1% occurrence frequency threshold *in any decade*, to ensure consistency.

The taxa lists differed in the taxonomic resolutions of recorded taxa. As the CPR time-period is the longer of the two, and the taxa are generally more coarsely taxonomically resolved, the taxa within the ICES list were aggregated to their equivalent resolution within the CPR taxa list. For example, the CPR taxon name 'Radiozoa' is a phylum, whereas in the ICES taxa list there were four taxon names within the phylum Radiozoa. These taxa were therefore aggregated to the coarser CPR resolution. In some cases, new groups were constructed to aggregate multiple taxa. 'Gelatinous zooplankton' was created as Cnidarians and Ctenophores were sometimes recorded as 'Coelenterata' within the ICES dataset. This nomenclature is outdated, and is not a monophyletic group, and so it would be impossible to determine whether these records related to '*Cnidaria*' taxa or '*Ctenophora*' taxa. Some taxa had resolutions too low for aggregation, for example records of '*Crustacea*' with associated life stages 'larvae' or 'nauplius'. Samples containing these records were removed before analysis, so the low taxonomic resolution did not skew results. Lastly, taxa that are not consistently recorded throughout the CPR time series, as a result of analysis changes, were removed. Similarly, any taxa within the ICES taxa list that would not be reliably sampled by the CPR due to their small size or delicate nature were removed, thus reducing biases from differing mesh sizes.

After integrating the taxonomic nomenclature and resolution of the two taxa lists, of taxa that occurred in over 1% of samples, 39 phytoplankton taxa and 27 zooplankton taxa were unique to the ICES list, whilst 10 phytoplankton taxa and 13 zooplankton taxa were unique to the CPR list. These differences could represent large changes in occurrence frequency over the time period, but could also still be a result of sampling biases between the two datasets, for example though different mesh sizes. We therefore only used taxa that occurred in over 1% of samples in both datasets. These lists of common phytoplankton and zooplankton taxa shared between the two datasets



182 represented taxa that were assumed to be consistently sampled by both surveys (Hällfors et al., 2013), further  
 183 minimising biases from differing mesh sizes , and consisted of 44 phytoplankton taxa and 30 zooplankton taxa  
 184 respectively (Table 1). Records of these shared common taxa were then extracted from the CPR and ICES samples,  
 185 before determining the occurrence frequency of each taxon for each sampling month. Months with fewer than 5  
 186 samples were removed before analysis.

187 **Table 1. 'Matching' taxa lists, at aggregated taxonomic resolution, used in analysis**

### Phytoplankton Matching List

#### Diatoms

<i>Asterionellopsis glacialis</i>	<i>Navicula</i> spp.
<i>Bacillaria paxillifera</i>	<i>Odontella aurita</i>
<i>Bacteriastrium</i> spp.	<i>Odontella sinensis</i>
<i>Bellerocha horoglacialis</i>	<i>Paralia sulcata</i>
<i>Ceratoneis closterium</i>	<i>Proboscia alata</i>
<i>Chaetoceros</i> spp.	<i>Pseudo-nitzschia delicatissima</i>
<i>Corethron</i> spp.	<i>Pseudo-nitzschia seriata</i>
<i>Coscinodiscus</i> spp.	<i>Rhaphoneis amphiceros</i>
<i>Coscinodiscus concinnus</i>	<i>Rhizosolenia hebetata f. semispina</i>
<i>Ditylum brightwellii</i>	<i>Rhizosolenia setigera</i>
<i>Eucampia zodiacus</i>	<i>Rhizosolenia styliformis</i>
<i>Fragilaria</i>	<i>Skeletonema costatum</i>
<i>Guinardia delicatula</i>	<i>Thalassionema</i> spp.
<i>Guinardia striata</i>	<i>Thalassiosira</i> spp.
<i>Halosphaera</i> spp.	<i>Thalassiothrix longissima</i>
<i>Lauderia danicus</i>	

#### Dinoflagellates, silicoflagellates and haptophytes

<i>Ceratium fusus</i>	<i>Tripos furca</i>
<i>Ceratium horridum</i>	<i>Tripos lineatus</i>
<i>Ceratium tripos</i>	<i>Tripos longipes</i>
<i>Dictyochophyceae</i>	<i>Tripos macroceros</i>
<i>Dinophysis</i> spp.	
<i>Gonyaulax</i>	
<i>Phaeocystis</i>	
<i>Prorocentrum</i> spp.	
<i>Protoperdinium</i> spp.	

### Zooplankton Matching List

#### Holoplankton

<i>Acartia</i> spp.	<i>Oithona</i> spp.
<i>Anomalocera patersoni</i>	<i>Para-Pseudocalanus</i> spp.
<i>Appendicularia</i> spp.	<i>Paraeuchaeta norvegica</i>
<i>Calanus</i> spp.	<i>Podon</i> spp.
<i>Centropages</i> spp.	<i>Temora longicornis</i>
<i>Centropages hamatus</i>	<i>Thecosomata</i>
<i>Centropages typicus</i>	<i>Tintinnidae</i>
<i>Chaetognatha</i> spp.	
<i>Copepoda</i> spp.	
<i>Corycaeus</i> spp.	
<i>Euphausiacea</i> spp. and <i>Mysida</i> spp.	
<i>Evadne</i> spp.	
<i>Foraminifera</i> spp.	
<i>Isias clavipes</i>	
<i>Labidocera wollastoni</i>	
<i>Metridia lucens lucens</i>	

#### Meroplankton

<i>Bivalvia</i> spp.
<i>Bryozoa</i> spp.
<i>Cirripedia</i> spp.
<i>Decapoda</i> spp.
<i>Echinodermata</i> spp.
<i>Pisces</i> spp.
<i>Polychaeta</i> spp.

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## 2.3 Multivariate analysis

To investigate whether significant change occurred in the plankton community between the ICES historical time-period and the beginning of the time-period covered by the CPR survey, we tested for an effect of time period (historical dataset, 1902-1912, to the 1960s decade of the CPR time period) on plankton community composition using multivariate generalised linear models with the *'mvabund'* package in R (Wang et al., 2012). This method fits a generalised linear model to each taxon separately, and then gives a summed likelihood ratio for the given predictors for each model, which can be used as a test statistic ('Sum-of-LR') for the effect of predictors on the community as a whole. Resampling is then done at the whole-sample level (here the sampling month) to test for significance while accounting for correlations between taxa (Wang et al., 2012). The method accounts for a mean-variance relationship in the data (Warton et al., 2012). The generalised linear models were fitted for the occurrence frequency of each taxa in each sampling month, with a complementary log-log link to accommodate the proportional, binomial data (Wang et al., 2012). For each model, the log of the sampling month occurrence frequency total was used as an offset as an approximate method of analysing relative compositional change, and weights were included so that sampling months with higher sample sizes were given stronger weighting. We extracted the univariate statistics for each taxon in the model, to examine the contribution of each taxon to any overall effect.

Furthermore, we visualised change in the plankton community over the extended time period using non-metric Multidimensional Scaling (nMDS) ordination plots. Plots were constructed for each area and plankton type using the *vegan* package in R (Oksanen et al., 2007). These were constructed based on the relative occurrence frequency of each of the matching list taxa in each sampling month.

After testing for the effect of time period on community composition, we tested whether SST difference between the two periods could explain any observed differences in community composition using multivariate generalised linear models. Here, models including SST were compared to models including SST and time-period, as a significant effect of time-period over and above SST suggests there is variation between the time-periods not explained by changes in SST alone. Lastly, we tested for any overall effect of SST on plankton community composition, over the whole extended time period, when examining the two datasets combined. Models with SST and season as predictors were compared against models with just season as a predictor to look for the influence over and above seasonality.

## 3 Results

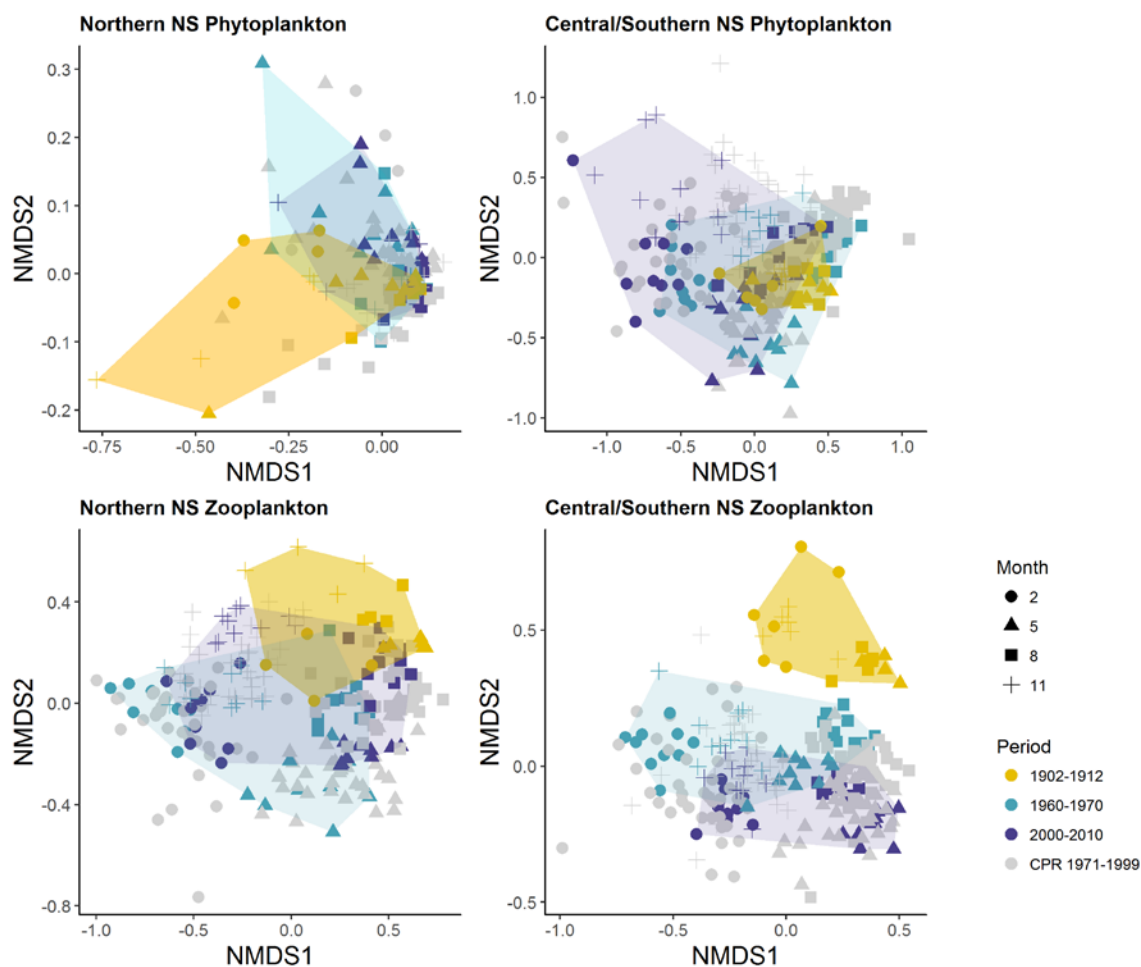
### 3.1 Changes in plankton community composition over time

Significant differences in overall community composition were found for both phytoplankton and zooplankton in both the Northern and Central/Southern North Sea areas, suggesting a change in the North Sea plankton community between the beginning of the 20<sup>th</sup> century and the 1960s. The zooplankton communities showed a stronger overall difference, with larger overall summed likelihood ratios for an effect of time period, despite a lower number of taxa within the list of shared common taxa (Northern North Sea: Sum-of-LR= 1891.3,  $p=0.004$ ; Central/Southern North Sea: Sum-of-LR= 2355.5,  $P=0.003$ ). In contrast, the overall effect of time period, although significant, was lower for phytoplankton communities, suggesting a smaller community change (Northern North Sea: Sum-of-LR= 299.44,  $p<0.001$ ; Central/Southern North Sea: Sum-of-LR= 825.65,  $p<0.001$ ).

However, when extracting the individual contributions of each taxon to the overall community response, a low number of taxa in all communities showed significant contributions to overall community responses. Furthermore, the overall community responses were largely dominated by a low number of taxa. For example, in each community over 20% of the variation was driven by one individual taxon, which showed changes in relative occurrence frequency in all months. These were *Protopteridinium* (a heterotrophic group) and *Tintinnidae* in the Northern North Sea area for phytoplankton and zooplankton communities respectively, which showed declines. In the Central/Southern North Sea area *Guinardia striata* showed a decline, whilst '*Euphausiacea* and *Mysida*' showed an increase. Out of these taxa, only the decline in *Tintinnidae* in the Northern North Sea was a statistically significant contribution to community change. Other taxa showing large contributions to overall effect were *Dinophysis* within the Northern North Sea phytoplankton community, and *Anomalocera patersoni* within the Northern North Sea zooplankton community, both of which showed a decline, although the decline in *Dinophysis* was not a statistically significant contribution to community change. Aside from these particular taxa, the overall community change between the beginning of the 20<sup>th</sup> century and the 1960s was distributed relatively evenly between the taxa, suggesting a holistic community change between the two time periods.

As sampling biases between the datasets, such as varying mesh sizes, may have influenced the taxa that had disproportionate contributions to overall community change, we removed taxa contributing over 20% of variation

244 between time periods before visualising community composition over the extended time period using nMDS plots  
 245 (Figure 2). 'May 1912' was removed due to being highly anomalous. Here, the stronger effect of time-period on  
 246 zooplankton composition can be seen with a clearer distinction between the historical (1902-1912) decade and the  
 247 1960s. Furthermore, there is a clearer distinction between the 1960s and the 2000s within the zooplankton plots,  
 248 especially for the Central/Southern North Sea, suggesting phytoplankton to be more stable in terms of change in  
 249 community composition over multi-decadal scales.



263 **Figure 2.** nMDS plots using Bray Curtis dissimilarity, based on monthly occurrence frequency of the matching list taxa  
 264 in each North Sea region. Data points from the ICES historical dataset (1902-1912) are shown in orange and are  
 265 bound by orange polygon (convex hull).  $K=3$  for all except Northern NS zooplankton, where  $k=4$  to lower stress. Data  
 266 points from 1971-1999 from the CPR survey are shown in grey, with data from the 1960s shown in blue and bounded  
 267 by blue polygon. Data from the 2000s decade are in purple and bounded by purple polygon, for additional context.

269

### 270 3.2 Influence of SST change on plankton communities

271 Taxa contributing over 20% of between-dataset variation then remained removed when analysing the effect of SST  
 272 on plankton community composition, to ensure any effects of SST found were not being driven by a small proportion  
 273 of the taxa. SST has increased in both the Northern and Central/Southern North Sea areas and particularly sharp  
 274 increases occurred during the late 1920s and 1980s (Figure 3). The average annual SST for the ICES historical time  
 275 period (1902-1912) was 9.00 °C for the Northern North Sea area, rising to 9.53 °C in the 1960s. In the  
 276 Central/Southern North Sea area, the average SST for the ICES historical time period was 9.59 °C, rising to 9.86 °C in  
 277 the 1960s.

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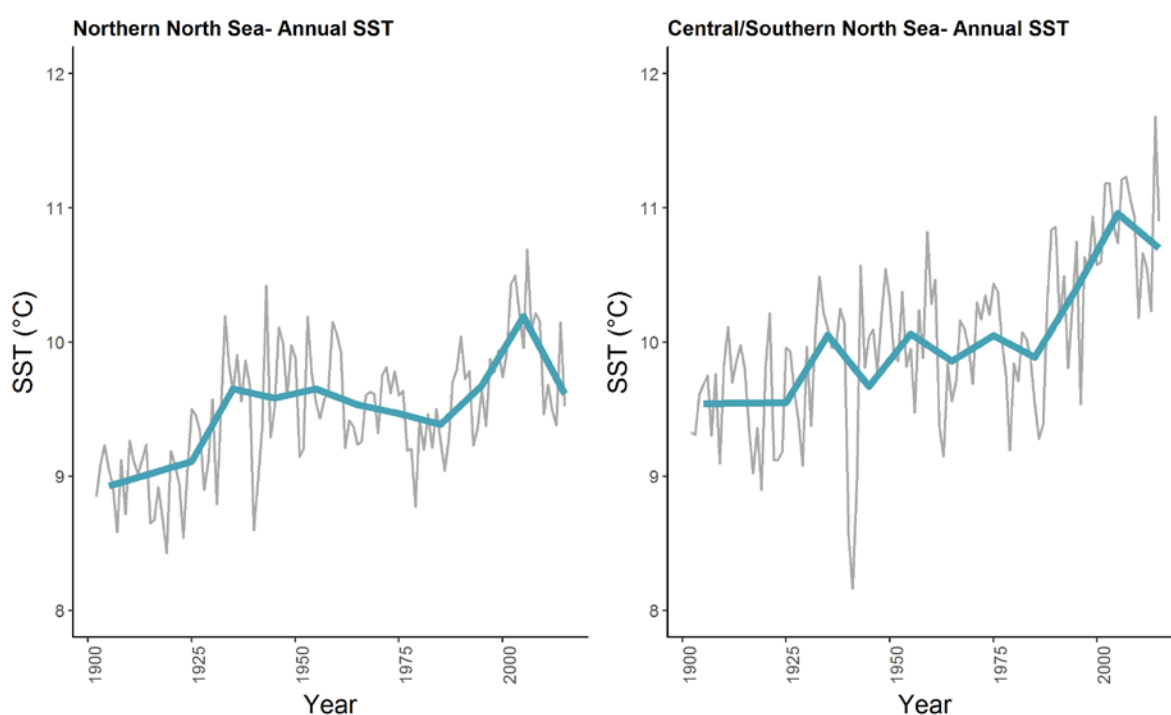
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286 **Figure 3.** North Sea SST variation between 1902 and 2015. Annual data are in grey and the 5 year mean is in blue.

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288 Differences in SST between the time periods suggest that changes observed in overall plankton community

289 composition between 1902-1912 and the 1960s coincided with changes in environmental conditions within the

290 North Sea. We tested this further using multivariate generalised linear models; a significant effect of time-period

291 over and above SST suggests there is variation between the time-periods not explained by changes in SST alone. A  
292 significant effect of time period over and above SST was found only in the Central/Southern North Sea  
293 phytoplankton community ( $p=0.023$ ), suggesting variation between time-periods could not be explained by SST  
294 change only in this community. In the Northern North Sea zooplankton and phytoplankton communities, as well as  
295 the Central/Southern zooplankton community there was no significant effect, suggesting variation could be linked to  
296 large-scale SST change.

297  
298 When then using both the ICES historical dataset and the full CPR dataset together, giving an extended temporal  
299 coverage, we found significant effects of SST on phytoplankton and zooplankton communities in both the Northern  
300 and Central/Southern North Sea areas (Table 2). SST had a greater influence in the Central/Southern North Sea than  
301 the Northern North Sea area on both phytoplankton and zooplankton composition, and a larger influence on  
302 zooplankton than phytoplankton overall. No phytoplankton taxa showed individual significant contribution to overall  
303 community response. In contrast, there were multiple individual significant contributions to the overall response  
304 within zooplankton communities, with the most number of significant individual contributions shown in the  
305 Central/Southern North Sea. These included both meroplankton and holoplankton taxa, with the largest  
306 contributions to overall community response from *Centropages typicus* and the multi-species group *Bivalvia*.  
307 *Centropages typicus* showed an increase in relative occurrence frequency over time, whilst *Bivalvia* showed a  
308 decrease in relative occurrence frequency over time, coinciding with increasing annual SST (Figure 4).

314 **Table 2.** Plankton community responses to SST when examining both datasets combined (1902-1912, 1958-2015).  
 315 Sum-of-LR= Summed likelihood ratio.

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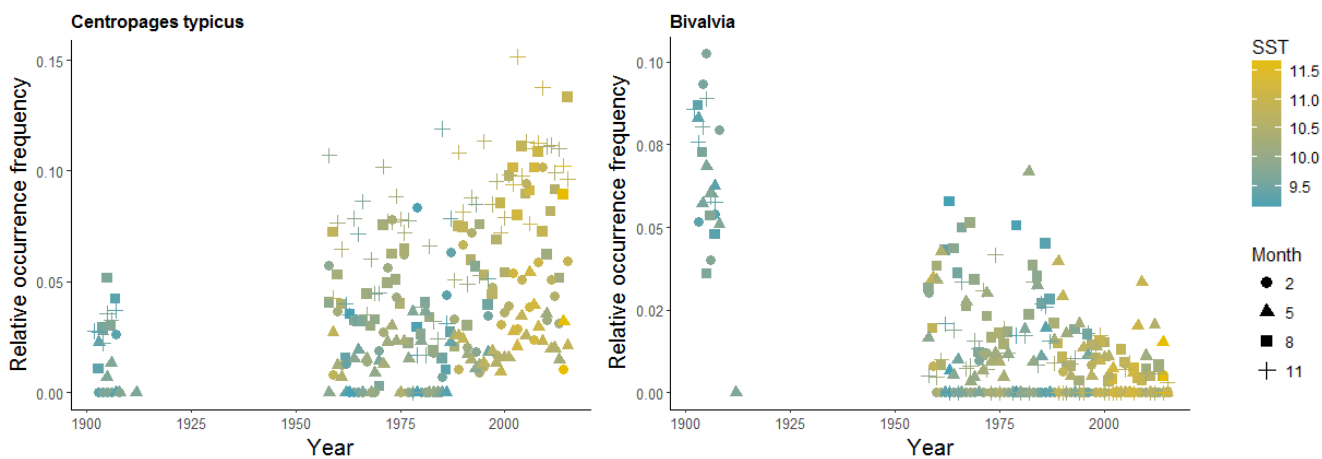
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Plankton community	Overall community response		Taxa with significant contributions to community response to SST over the extended time period
	Sum-of-LR	p	
Northern NS phytoplankton	195.7	0.044	N/A
Central/Southern NS phytoplankton	542.86	<0.001	N/A
Northern NS zooplankton	669.94	<0.001	<i>Anomalocera patersoni</i> <i>Decapoda spp.</i> <i>Echinodermata spp.</i>
Central/Southern NS zooplankton	1999.7	<0.001	<i>Bivalvia</i> <i>Calanus spp.</i> <i>Centropages typicus</i> <i>Corycaeus spp.</i> <i>Decapoda spp.</i> <i>Oithona spp.</i> <i>Para-Pseudocalanus spp.</i> <i>Polychaeta spp.</i>



325

326 **Figure 4.** A) Occurrence frequency of *Centropages typicus* by month from wider time-period. B) Occurrence frequency  
 327 of *Bivalvia* by months from wider time-period.

328 Although overall community composition change between 1902-1912 and the 1960s may be linked to changes in  
329 SST, taxa that had the largest univariate contributions to community change did not necessarily have large responses  
330 to SST across the wider time period (1902-1912, 1958-2015). This suggests that although a change in temperature  
331 conditions may have contributed to the overall community response, it does not necessarily explain individual taxon  
332 changes between the two time periods. Furthermore, any potential influences of specific environmental drivers on  
333 community composition differences between the two time-periods may be at least partially obscured by the  
334 differences in sampling and analysis methodologies between the two datasets used, and the low quantitative  
335 resolution available.

## 336 **4 Discussion**

338 Here, we have demonstrated the value of 'rescued' historical plankton data in increasing the temporal scale of  
339 understanding of community change. By harmonising the taxonomic lists from the two datasets in order to ensure  
340 comparability and then further selecting a subset of shared, common taxa based on a 1% occurrence frequency  
341 threshold, and using presence/absence semi-quantitative resolution, we have reduced the influence of disparate  
342 sampling and analysis methodologies. Results suggest that the 1960s had a significantly different plankton  
343 community composition compared to the early 1900s, indicated by variation in the relative occurrence frequency of  
344 shared common taxa. Differences in community composition between time periods were largely driven by a small  
345 number of taxa. The remaining effect was shared relatively evenly between the remaining taxa, suggesting the  
346 overall significant changes in community composition are a result of subtle change across the taxa list, with  
347 individual taxa having mainly non-significant contributions to overall community response.

348 Zooplankton communities showed a greater difference between the ICES historical time period and the 1960s  
349 decade of the CPR time period than phytoplankton communities. The nMDS plots also revealed clearer visual  
350 distinctions between the 1960s decade and the 2000s decade within the zooplankton communities than within the  
351 phytoplankton communities. This suggests that although differences between the time periods were found within  
352 phytoplankton communities, over the whole time period the phytoplankton community showed less directional  
353 change in community composition at the multi-decadal scale than zooplankton communities. A similar result was



354 found during the OSPAR (Oslo-Paris Convention for the Protection of the North-East Atlantic) Intermediate  
355 Assessment 2017, where larger changes in indicators of zooplankton community structure were found compared to  
356 phytoplankton communities (OSPAR, 2017). This assessment result could therefore be representative of multi-  
357 decadal patterns of variation occurring at the century-scale.

358 Furthermore, we found that the plankton community change identified between 1902-1912 and the 1960s could be  
359 explained through changes in SST in Central/Southern North Sea zooplankton and Northern North Sea  
360 phytoplankton and zooplankton. These community changes in response to SST could therefore be attributed to a  
361 regime shift that has been shown to have occurred in the North Atlantic during the 1920s and 1930s, which is argued  
362 to be the largest and most significant climate-induced regime shift of the 20<sup>th</sup> century (Drinkwater, 2006), associated  
363 with increases in SST. Furthermore, change in the Central/Southern North Sea phytoplankton community could not  
364 be explained by SST change. It is likely, therefore, that finer scale changes, in variables other than SST, drove the  
365 change in the Central/Southern North Sea phytoplankton community. Hällfors et al. (2013) similarly described an  
366 unknown 'period effect' between the ICES historical time period and contemporary phytoplankton samples in the  
367 Baltic Sea, where variation could not be explained by environmental change alone, and instead they hypothesise a  
368 potential signal of eutrophication in the change observed. At the regional scale in the North Sea however, previous  
369 research has suggested that eutrophication occurs mainly in coastal regions, rather than open sea (McQuatters-  
370 Gollop et al., 2009). Furthermore, although we are confident that differences in taxonomic nomenclature and  
371 resolution are not driving any patterns observed, we cannot rule out an influence of the low quantitative resolution  
372 resulting from sampling and analysis biases, especially for the taxa showing disproportionate contributions to the  
373 overall community response.

374 By integrating the CPR survey with the ICES historical data, we facilitated exploration of the influence of warming  
375 SSTs on multidecadal plankton community change at the century-scale, although focusing on occurrence frequency,  
376 rather than abundance values. Over the extended time period (1902-1912, 1958-2015), SST had a stronger influence  
377 on zooplankton communities than phytoplankton, in both the Northern and Central/Southern North Sea areas. In  
378 particular, it is known that temperature is an important structural variable for zooplankton communities and is a key  
379 determinant of the limits to distributions (Richardson, 2008). In contrast, although SST was a significant driver of  
380 community composition in phytoplankton in both the Northern and Central/Southern North Sea, no single taxa

381 showed significant contributions to the overall community effect. Previous studies have suggested the importance of  
382 physical variables other than SST directly influencing phytoplankton community composition including salinity and  
383 wind stress (Hinder et al., 2012).

384 Multiple zooplankton taxa in the Central/Southern North Sea area showed significant univariate responses to SST  
385 change, with *Centropages typicus* and the multi-species group *Bivalvia* showing the largest responses. A positive  
386 association between the abundance of *Centropages typicus* and SST has previously been identified in the North Sea  
387 (Lindley and Reid, 2002), and this pattern is also shown here when examining the CPR time-series at a  
388 presence/absence resolution. The lack of a large difference in relative occurrence frequency between the beginning  
389 of the 20<sup>th</sup> century and the 1960s found here however suggests that the response of *Centropages typicus* to SST  
390 occurred since the 1960s. In contrast, the larger difference in the occurrence frequency of *Bivalvia* found here  
391 between the beginning of the 20<sup>th</sup> century and the 1960s suggests the decline in the abundance of bivalve larvae  
392 previously identified in the North Sea (Kirby et al., 2008) occurred over a longer time scale. Kirby et al. hypothesise  
393 that the long-term decline in bivalve larvae found through the CPR survey is a result of predation from increasing  
394 abundance of decapod larvae, also observed through the CPR survey, and the increase in decapod larvae is  
395 associated with increasing SST (Lindley et al., 2010). In this study, decapod larvae in the Central/Southern North Sea  
396 had a significant response to SST, and increased in relative occurrence between 1902-1912 and the 1960s,  
397 suggesting that trophic amplification of a climate signal could explain the decrease in bivalve larvae also at the  
398 century scale. The differences in whether the taxa with strong overall responses to SST also showed large differences  
399 in occurrence frequency between time periods suggests that the temporal scale of responses to SST change, and  
400 temporal scale of baseline shifts, is variable between individual taxa.

#### 401 402 **4.1 Conclusions and policy implications**

403 Through integrating and directly comparing the CPR dataset to the ICES historical database, important  
404 considerations have been identified for using disparate plankton datasets together, with applications for large scale  
405 assessment and integrated monitoring programmes, such as regional scale assessments undertaken at the OSPAR  
406 level (OSPAR 2017). Particularly, zooplankton taxa varied greatly in the taxonomic resolution in which they were

407 recorded between surveys, and much attention needs to be drawn to this when designing integrated monitoring  
408 programmes constructed from different surveys. However, we have shown that a subset list of shared common taxa  
409 can inform on community change when combining data from disparate sources. Furthermore, occurrence frequency  
410 seems to be a relevant proxy for abundance, when abundance data is non-comparable, for example occurrence  
411 frequency resolution still revealed strong seasonality signals. As sampling and analysis biases cannot ever be fully  
412 reconciled in contemporary comparisons of rescued historical datasets, such as varying mesh sizes, often resulting in  
413 low quantitative resolution, we suggest that 'rescued' historical datasets can be useful as an additional contextual  
414 tool for understanding climate change effects on plankton communities, but caution should be employed when  
415 using disparate historical datasets as robust evidence bases on their own.

416 A stable historical baseline, from which plankton communities are assessed for impacts of direct anthropogenic  
417 pressures, may be hard to define in the North Sea, as the plankton communities vary on inter-annual, multi-decadal  
418 and, suggested here, century-wide scales in response to environmental change. Phytoplankton community  
419 composition may show less directional change in community composition, in terms of the relative occurrence  
420 frequency of common taxa, over multi decadal time scales than zooplankton communities. Although statistically  
421 significant changes were observed in particular individual taxa between time periods, and across the wider time-  
422 series in response to SST, this does not necessarily inform on the ecological significance of changes. When formally  
423 assessing change in North Sea plankton communities under policy drivers, it is important to consider the functional  
424 consequences of community change, as well as the century-scale shifts in community composition baselines.

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