INTRODUCTION

Links between seawater temperature and the skeleton mineralogy of marine animals were first suggested by Clarke & Wheeler (1922), when correlations were observed between the inorganic constituents of marine organisms and local habitat temperature. In 1954, increased magnesium incorporation in calcium carbonate was recognised as a proxy for increasing seawater temperature by Chave (1954). In the same year, Lowenstein (1954) identified a positive correlation between seawater temperature and aragonite content in bimineralic animals. Substitution of Mg$^{2+}$ for Ca$^{2+}$ in calcite has been studied through inorganic precipitation studies, which demonstrate that incorporation of magnesium within the calcite lattice is thermodynamically more favourable in warmer waters (Chilanger 1962, Katz 1973, Burton & Walter 1987, Mucci 1987, Oomori et al. 1987). Similarly, aragonite is more energetically expensive to build than calcite (Anderson &
Magnesium (Mg) is naturally incorporated within bryozoan calcite and this can range from low (≤4 wt% MgCO₃), through intermediate (>4 wt% MgCO₃ to <8 wt% MgCO₃) to high (≥8 wt% MgCO₃) (Smith et al. 2006, Smith & Girvan 2010), with most bryozoans featuring low or intermediate Mg-calcite (Smith et al. 2006, Lombardi et al. 2008, Kuklinski & Taylor 2009, Taylor et al. 2009). Skeletal variability is exhibited at all taxonomic levels within the phylum with the majority of variability occurring between different clades and species. In all species, some mineralogical variation can be seen between different specimens and often within colonies.

Both biological and environmental controls are believed to influence bryozoan aragonite and Mg-calcite content within species, although the relative influence of these controls on skeleton composition is much debated. Biological control refers to factors such as astogeny (Smith et al. 1998, Kuklinski & Taylor 2009, Smith & Girvan 2010), growth rates (Smith 2007, Kuklinski & Taylor 2008b), food availability (Bone & James 1993), physiological ‘wellness’ (Stanley & Hardie 1998) and directed adaptation to ecological niches in isolated populations (Cheetham et al. 1994). Environmental control indicates that skeletal mineralogy is driven by the seawater within which the bryozoan lives with little or no physiological involvement from the animal itself. Environmental factors influencing mineralogy may include salinity (Bohaty et al. 2012) and carbonate chemistry (Müller et al. 2014) in addition to temperature; however, temperature is assumed to be the dominant control when using wt% MgCO₃ in calcite or wt% aragonite as paleoclimatic proxies (Chave 1954, Lowenstam 1954, Weiner & Dove 2003).

Following the first reports of relationships between mineralogy and environmental conditions, it was observed that not all mineral deposition occurred in isotopic equilibrium and that ‘the presence of a physiological effect in the case of certain groups of animals such as the echinoderms and corals, and plants, such as coralline algae, has seemed probable’ (Epstein et al. 1951). This biological effect on mineralogy subsequently came to be known in biogeochemistry as the ‘vital effect’. A strong vital effect can override, or mask, mineralogical responses to environmental factors such as temperature, compromising the usefulness of a species, or phylum, as a paleoclimatic recorder (Weiner & Dove 2003). Past bryozoan studies have alluded to the existence of a vital effect (Schafer & Bader 2008, Kuklinski & Taylor 2009, Taylor et al. 2009, Smith & Girvan 2010, Loxton et al. 2014); however, the relative influence of biological and environmental control on skeletal mineralogy has not yet been established. Past mineralogical studies on temperate bryozoans have been constrained by limited availability of samples and environmental measurements, and as a consequence, it has not been possible to clearly determine the usefulness of the Bryozoa as paleoenvironmental recorders.

The overall aim of this study was to conduct a high-replicate, multi-site study on the skeletal mineralogy of selected temperate bryozoans to allow compa-
isons to be drawn between the mineralogy and range of skeletal variability between species, and to investigate the relative influence of environmental and biological factors on skeleton chemistry. The study was conducted over a relatively small spatial scale of 12 km, and the reported environmental homogeneity of the study location (Bennett & Covey 1998) leads us to predict that no environmentally driven differences in skeletal mineralogy between sites will be observed. Any differences in skeletal Mg-calcite, or aragonite, which do occur between sites, could therefore be attributed to biological control of skeletal mineralogy.

To investigate this, samples of 3 temperate bryozoan species were collected and examined from 5 sites in Scapa Flow, Orkney, Scotland. Multi-year measurements of seawater temperature in Scapa Flow were recorded at 2 depths to establish interseasonal variability and any effect of depth. Further environmental parameters of seawater temperature, depth and salinity were recorded at each site at the time of specimen collection. Water samples were collected at each site for laboratory analysis and calculation of carbonate system parameters. The resulting datasets were examined to (1) compare the mineralogy and range of skeletal Mg-calcite and aragonite variability between the 3 temperate species, (2) investigate the relationship between bryozoan wt% MgCO$_3$ in calcite, wt% aragonite in total CaCO$_3$ and environmental factors, and (3) identify potential biological factors influencing temperate bryozoan mineralogy. This study will establish a mineralogical baseline for 3 common British bryozoans for comparison in future studies.

**MATERIALS AND METHODS**

**Study area**

Orkney is an archipelago of islands situated 16 km northeast of mainland Scotland (58° to 59° N, 2° to 3° W), at the point of confluence of North Atlantic and North Sea waters (Fig. 1A). At the centre of this island group lies Scapa Flow, a semi-enclosed marine basin covering an area of ~130 km$^2$ (Joyce 2004), where water depths reach a maximum of 70 m. Tides and currents circulate through Hoy Sound, in the northwest, and Hoxa Sound in the south, which open to the Atlantic Ocean and Pentland Firth, respectively. Benthic sea temperature is highly variable throughout the year with a reported range from 5 to 14°C (Bennett & Covey 1998). The salinity regime of near bottom waters varies annually from 30 to 35 psu (Bennett & Covey 1998). More detail on the profile of the Orkney marine environment was published by Jones (1975).

**Sampling and specimen processing protocol**

Between October 2010 and March 2013, hourly seawater temperature measurements were recorded using a HOBO UA-002-64 temperature logger at 6 and 12 m benthic sites in Scapa Flow (6 m site: 58° 55.300’ N, 3° 06.579’ W; 12 m site: 58° 55.273’ N,
Logger accuracy is reported to be 0.5°C for temperature (measurable range −20 to 70°C) (HOBO® Pendant Temperature/Light Data Logger (UA-002-xx) manual, p 1–3).

All sample collecting took place by SCUBA in a single week in May 2012. Five collection sites (1 to 5) were selected within Scapa Flow with similar depths (9.5 ± 3.5 m) and substrate (boulders and cobbles) (Fig. 1B). All sample sites had similar exposure and geomorphology. At each site, specimens were collected on pebbles from below the kelp line from within a 5 m² area at constant depth. Sites were an average of 6 km apart (range: 1.7 to 11.8 km), and were within an area of 28.3 km². Conductivity and temperature were measured at each site during collection using a HOBO U24-002 salt water conductivity logger; instrument accuracy is reported to be 0.1°C for temperature (measurable range −2 to 36°C) and 3% of conductivity reading (HOBO® U24 Conductivity Logger (U24-002-C) manual, p 2–7). Conductivity was converted to practical salinity units, as defined on the practical salinity scale of 1978 (Mcdougall et al. 2010) using the Gibbs function of seawater. In situ water samples were collected by SCUBA from each site in 500 ml borosilicate glass bottles. To prevent any further biological activity in the stored sample, each sample was poisoned within 1 h of collection with a saturated solution of mercuric chloride (7 g per 100 ml deionised water) in a 0.02% volume ratio. Carbonate analysis was undertaken at the National Oceanography Centre, Southampton following the protocols of Dickson et al. (2007). Dissolved inorganic carbon (DIC) was determined using coulometric titration; total alkalinity (TA) was determined using closed-cell titration. The precision of DIC and TA measurements obtained using this method is reported as ±0.05% or better (Dumousseaud et al. 2011). The measurement of DIC, TA, temperature and salinity allows the calculation of the other variables of the carbonate system through the use of thermodynamic constants. The CO2SYS program (Lewis & Wallace 1998) was used for the recalculation of the carbonate system variables using the thermodynamic constants of Mehrbach et al. (1973) refitted by Dickson & Millero (1987). Carbonate analysis was conducted according to the methods of Dumousseaud et al. (2011).

A total of 480 specimens of 3 bryozoan species were collected for mineralogical examination. The species selected for analysis from Orkney were the cheilostomatous bryozoans Microporella ciliata, sensu stricto (Pallas, 1766), Membraniporella nitida (Johnston, 1838) and Escharella immersa (Fleming, 1828).

The study species were selected on the basis of being well-known, common species with distinctive features for identification (Fig. 2). The 3 species represent both calcitic and bimineralic mineralisers and were present in sufficient quantities for sampling during the experimental period. M. ciliata and M. nitida are both reported to have highest levels of settlement in the spring and summer (Maughan & Barnes 2000, Denitto et al. 2007), while, between 2010 and 2013, the authors observed E. immersa as having a predominantly winter breeding cycle. Growth rate data are not available for E. immersa or M. nitida; however, Ball et al. (1995) estimate M. ciliata to grow at a rate of 1 mm mo⁻¹.

Specimens were identified to species level under a dissection (stereo) microscope (Zeiss) using the monographs of Hayward & Ryland (1998, 1999) and the re-description of Microporella by Kuklinski & Taylor (2008a). All specimens were alive when collected and were subsequently rinsed in fresh water and air dried for a minimum of 1 mo before samples were extracted for analysis. To minimise any potential mineralogical variability caused by season or specimen age, individual colonies of similar diameter were selected for sampling. Each sample consisted of ~5 zooids extracted from the outermost growing edge of a colony, containing the most recently deposited skeletal material. As far as possible, care was taken to ensure that no substrate (e.g. coralline algae) or epibionts were included within the sample, as they could potentially contaminate results through their added mineralogies.


**Analysis techniques**

Mineralogical analyses were conducted at the EMMA unit (NHM London) using an Enraf-Nonius X-ray diffractometer (XRD) with an INEL CPS-120 curved position-sensitive detector and cobalt X-ray source. Tube operation conditions were 40 kV and 40 mA. A primary germanium 111 monochromator (INEL) with slit settings of 0.14 x 5 mm was used to confine the X-ray beam to pure Co Ka1 radiation. Samples were measured in reflection using asymmetric flat-plate geometry. Diffracted intensities were collected simultaneously over a 2-θ range of...
120° without angular movement of tube, sample or
detector. The angle between the incoming mono-
chromatic beam and sample holder was kept con-
stant at 5.9°. The sample holder was rotated to in-
crease the number of crystallites and the randomness
of their orientations in the X-ray beam. The angular
linearity of the position-sensitive detector was cali-
brated using silver behenate (AgC_{22}H_{43}O_{2}) and NIST
silicon powder (SRM 640) as external standards and
the calibration curve was fitted using a least-squares
cubic spline function.

Bryozoan samples were powdered using an agate
pestle and mortar and affixed using a drop of acetone
to single quartz crystal substrates. Quantitative XRD
analysis was undertaken to determine the dominant
calcium carbonate polymorph, the proportions of cal-
cite and aragonite in bimineralic species, and the Mg
content of the calcite.

To determine the proportions of aragonite and
calcite in bimineralic species, peak intensities were
fitted to standard patterns generated from 100% aragonite and 100% calcite. Results are presented
as wt% aragonite, which is the proportion of the
total calcium carbonate that is represented by the
polymorph, aragonite. The error associated with this
method is estimated to be 2% based on repeatability
studies of samples with a known aragonite propor-
tion (Kuklinski & Taylor 2009). To calculate wt%
MgCO_{3} in calcite, the position of the d104 peak was
measured, assuming a linear interpolation between
calcite (CaCO_{3}) and Mg-calcite (MgCO_{3}). Wt%
MgCO_{3} in calcite is the proportion of the total cal-
cite (Ca_{(1-x)}Mg_{x}CO_{3}) within which the Ca^{2+} ion has
been substituted for an Mg^{2+} ion. Mackenzie et al.
(1983) validated the linear relationship between
d104 and Mg content as accurate in the composi-
tional range up to 17.4 wt% MgCO_{3} in calcite;
above this range there is nonlinear behaviour. Pre-
vious measurements of bryozoan MgCO_{3} in calcite
have ranged between 0 and 13.7 wt% (Smith et al.
2006) — within the validated linear range. Composi-
tion information is considered accurate to within 2%
on a well-calibrated instrument (Kuklinski & Taylor
2009). Data evaluation was undertaken using WinX-
Pow software.

Statistics and data analysis

The 6 and 12 m long-term temperature data series
were statistically compared using a general linear
model ANOVA.
All mineralogical measurements underwent weighted average transformation in the context of beta regression following the methodology of Smithson & Verkuilen (2006). This initial transformation, \((y(n-1) + 0.5)/n\), ensured that no data points equalled 0 or 1 and prepared the data for subsequent transformation using the logit function. The logit function, \(\text{logit}(p) = \ln(p/[1 - p])\), accounts for the mineralogical measurements being proportional data (\(p\)) (Warton & Hui 2011).

Mineralogy data were tested for normality using Anderson-Darling normality tests and homogeneity of variance using Levine’s test of equal variance. Mineralogy measurements were found to be normally distributed in most cases, but failed Levine’s test of equal variance. The criteria for parametric testing were not satisfied for the mineralogical measurements; therefore, data were analysed using non-parametric tests. Variance was tested using Kruskal 1-way ANOVA with post hoc pair-wise testing (Mann-Whitney U-test). Correlations with environmental variables were explored using the Kendall correlation method. P-values for all analyses were calculated based on the probability of error of 0.05. All statistical data analysis was conducted in the R programming environment (R Core Team 2013).

**RESULTS**

**Environmental analysis**

The long-term data series showed no statistical difference (ANOVA) between 6 and 12 m depths (Fig. 3).

Temperature data measured during sample collection at the sites was compared with data from the same date and time (±2 h) extracted from the long-term data series at 12 m depth. The site data was found to deviate from the long-term data by a mean of just 0.12°C (maximum deviation = 0.208°C, minimum deviation = 0.054°C).

Environmental measurements taken from the individual sites also show little seawater temperature variability between sites (mean = 8.23°C, SD = ±0.07°C) with only slightly higher variability in salinity (mean = 33.01 psu, SD = ±0.49 psu) (Table 1).

**Species mineralogy**

In this study, the wt% MgCO\(_3\) in calcite and wt% aragonite in CaCO\(_3\) of 480 specimens of 3 bryozoan species was quantified. All of the species, Membraniporella nitida (n = 139), Microporella ciliata (n = 145) and Escharella immersa (n = 146), were found to have intermediate Mg-calcite (4 to 8 wt% MgCO\(_3\) in calcite) with a mean wt% MgCO\(_3\) in calcite (±SD) of 6.25 (±0.648), 6.88 (±0.638) and 5.66 (±0.612), respectively (Fig. 4).

One of the species, M. nitida, was found to be entirely calcitic. M. ciliata and E. immersa were both found to be calcite-dominated bimineralisers with varying volumes of aragonite present. M. ciliata was found to contain levels of aragonite ranging between 1 and 66 wt% (mean = 22.7 wt%, SD = ±18.96 wt%) of total CaCO\(_3\) present. E. immersa was likewise found to contain levels of aragonite ranging between 1 and 75 wt% (mean = 31 wt%, SD = ±18.69 wt%) of total CaCO\(_3\).
ANOVA analysis showed a statistically significant difference between species for both the wt% MgCO₃ in calcite (p < 0.001, \(F = 134.21\)) and the proportion of aragonite in total CaCO₃ (p < 0.001, \(F = 143.3\)).

Mineralogical variability between sites

Results of a Kruskall-Wallis 1-way ANOVA analysis showed that (a) the mean skeletal wt% MgCO₃ in calcite and (b) the mean skeletal wt% aragonite in total CaCO₃ were different between sites for all 3 species (Table 2). Post hoc Mann-Whitney U-tests showed that (a) wt% MgCO₃ in calcite and (b) the mean skeletal wt% aragonite in total CaCO₃ showed exactly the same variability, being both statistically significantly different between sites in 18 out of 30 cases (Table 3).

<table>
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<th>Site</th>
<th>South Cava</th>
<th>Holm of Houton</th>
<th>Flotta</th>
<th>Rysa Little</th>
<th>Barrel of Butter</th>
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<tr>
<td>Date</td>
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<td>09/05/2012</td>
<td>10/05/2012</td>
<td>06/05/2012</td>
<td>03/05/2012</td>
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<td>10</td>
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<td>DIC (µmol kg⁻¹ seawater)</td>
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<td>2096.3</td>
<td>2086.9</td>
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<td>(\text{CO}_3^{2-}) (µmol kg⁻¹ seawater)</td>
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<td>143.5</td>
<td>159.8</td>
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<td>(\Omega_{\text{calcite}})</td>
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<td>3.46</td>
<td>3.85</td>
<td>3.55</td>
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Table 1. Physical and environmental characteristics for the 5 study sites at the time of collection. Missing carbonate data for site 5 (–) is due to sample damage during collection. Dates given as d/mo/yr. Seawater scale used for pH measurements.

DIC = dissolved inorganic carbon; \(\Omega_{\text{calcite}}\) = calcite saturation state

**Fig. 4.** Mean wt% MgCO₃ in calcite for the 3 temperate species. The box shows the standard deviation around the mean; the whiskers indicate the range. The scatterplot shows the spread of samples. Previous analyses by Taylor et al. (2009) are indicated with triangles.
Examination of the Mann-Whitney post hoc analysis of both skeletal wt% MgCO₃ in calcite and wt% aragonite in total CaCO₃ revealed that Sites 1 and 2, South Cava Island and Holm of Houton, were the most mineralogically distinct of all the study sites, differing statistically from 9 of the 12 species–site combinations. The least mineralogically distinct for both measures of skeletal mineralogy was Site 5, Barrel of Butter, which differed significantly in only 5 out of 12 species–site combinations.

Variability in skeletal MgCO₃ in calcite and proportion of total CaCO₃ consisting of aragonite between species and sites can be seen in Fig. 5. All species show the same skeletal mineralogical distinctness between sites, differing statistically in 50% (6 out of 12) of sites. Although overall mineralogical distinctness is the same for wt% aragonite in CaCO₃ and wt% MgCO₃ in calcite, the species that are distinct between sites are different for the 2 mineralogical measurements.

The skeletal composition of *E. immersa* showed a statistically significant negative correlation between site temperature and wt% MgCO₃ in calcite and positive correlations between site salinity and TA with wt% MgCO₃ in calcite (Table 4). The skeletal composition of *M. nitida* showed a negative correlation between site salinity and wt% MgCO₃ in calcite and positive correlations between pH, CO₃²⁻ and calcite saturation state (Ωcalcite) with wt% MgCO₃ in calcite (Table 4).

The skeletal composition of *E. immersa* showed a negative correlation between wt% aragonite in CaCO₃ and site salinity, and positive correlations between wt% aragonite in CaCO₃ and pH, CO₃²⁻ and Ωcalcite (Table 4). As salinity decreases, and pH, CO₃²⁻ and Ωcalcite increase, there is an increase in wt% aragonite in skeletal CaCO₃ (and a corresponding decrease in calcite). The skeletal composition of *M. ciliata* showed weak statistically significant correlations between wt% aragonite in CaCO₃ and site salinity and wt% aragonite in total CaCO₃.

![Fig. 5. Variability in wt% MgCO₃ in calcite (left) and wt% aragonite in total CaCO₃ (right) between the 3 temperate species and the 5 study sites. The box shows the standard deviation around the mean; the whiskers indicate the range; the scatterplot shows the spread of samples](image-url)
temperature and depth. Wt% aragonite in CaCO₃ decreases (and calcite increases) with increasing temperature; wt% aragonite in CaCO₃ increases (and calcite decreases) with increasing depth (Table 4).

### DISCUSSION

In this study, a high-replicate, multi-site analysis of the skeletal mineralogy of temperate bryozoans from Orkney was conducted. This analysis allows comparisons to be drawn between the mineralogy and range of skeletal variability between species, and enables investigation of the relative influence of environmental and biological factors on skeleton chemistry.

The 3 study species were found to have statistically different mineralogies to each other with both wt% MgCO₃ in calcite and wt% aragonite in CaCO₃ differing significantly in all cases. Aragonite was present in *Microporella ciliata* and *Escharella immersa*, but absent in *Membraniporella nitida*. All species were found to have intermediate Mg-calcite (4 to 8 wt% MgCO₃ in calcite). Although all species are ascophorans, the calcitic species *M. nitida* is from the family *Cribriilinidae*. Previous mineralogical analysis of *cribrilinid* species have led to the family being reported as calcitic (Smith et al. 2006) and the data presented in this study confirm this. Although the species *E. immersa* and *M. ciliata* are not in the same taxonomic family, recent phylogenetic analysis by Waeschenbach et al. (2012) placed these bimineralic species within the same clade alongside *Escharoides coccinea*, *Oshurkovia littoralis*, *Pentapora fascialis* and *Cryptosula pallasiana*. Previous mineralogical studies on *E. coccinea*, *O. littoralis*, *P. fascialis* and *C. pallasiana* have shown them all to be bimineralic (Smith et al. 2006, Taylor et al. 2009) and the data presented in this study provide further evidence of the bimineralic propensity of this clade. The differences in mineralogy between species are, therefore, most likely to be attributable to their differential taxonomic and phylogenetic affinities.

Analysis showed that mean skeletal wt% MgCO₃ in calcite and wt% aragonite in total CaCO₃ were different between sites for all species. In this study, the relationship between environmental conditions and aragonite deposition was inconsistent between the 2 bimineralic species examined and no species showed the predicted positive relationship between temperature and aragonite deposition (Anderson & Crerer 1993). The predicted positive relationship between seawater temperature and wt% MgCO₃ in calcite was also not exhibited in any of the species examined. The relationships between Mg-calcite and temperature were found to be inconsistent between species, with responses ‘bucking the trend’ of what would be expected from the literature (Chave 1954, Lowenstam 1954) in some or all cases. Carbonate system parameters of pH, CO₃²⁻ and Ωcalcite were shown to have positive correlations with wt% aragonite in *E. immersa* and with wt% MgCO₃ in calcite in *M. nitida*. *E. immersa* also showed a positive correlation between TA and wt% MgCO₃ in calcite. Investigations on cultured planktonic Foraminifera (Lea et al. 1999) and coccoliths (Müller et al. 2014) have reported a pattern of increasing wt% MgCO₃ in calcite with decreasing pH and associated carbonate

<table>
<thead>
<tr>
<th></th>
<th>E. immersa</th>
<th>M. nitida</th>
<th>M. ciliata</th>
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Table 4: Results of Kendall's correlation analysis comparing (1) wt% MgCO₃ in calcite and (2) wt% aragonite in total CaCO₃ for samples of single species to site environmental variables (Table 1). All mineralogy data were logit transformed prior to analysis. Ωcalcite = calcite saturation state; x = no statistically significant correlation detected; p = p-value; τ = Kendall's tau; ‘–’ = missing data.
parameters; however, this pattern was not seen in any of the bryozoan species in this study. An in situ experiment on the bryozoan species *Myriapora truncata* by Lombardi et al. (2010) found wt% MgCO$_3$ in calcite to decrease with decreasing pH. Our study also detected this positive correlation; however, it was only significant in a single species and therefore cannot be considered reliable evidence of a relationship between skeletal mineralogy and carbonate system parameters in the Bryozoa.

As predicted, environmental conditions alone do not adequately explain the differences in wt% MgCO$_3$ in calcite or aragonite found between sites. As reported by Bennett & Covey (1998), and confirmed in this study, Scapa Flow is found to have a high level of homogeneity in environmental parameters. Across the 12 km scale, only minimal variation in temperature and salinity was detected between sites and there was no significant difference between site measurements and the long-term dataset. Carbonate system parameters were also shown to have only a low level of variation between sites. Thermodynamic studies have shown that, where temperature is the only influencing factor, Mg incorporation within calcite can be expected to increase by ~3% per 1°C (Rosenthal et al. 1997). The temperature differences between sites in this study are fractions of a degree and so any increase in wt% MgCO$_3$ in calcite caused by temperature may be below the levels of detection. We suggest that the results presented in this study provide evidence that biological processes are influencing the process of Mg-calcite and aragonite deposition in these temperate bryozoan species.

Two of the study species, *M. nitida* and *E. immersa*, showed contrasting responses of mineralogy to salinity. This dichotomy may be explained by the ecological preferences of the 2 species. *E. immersa* is only found in full salinity waters (Hayward & Ryland 1999), while *M. nitida* has also been reported from brackish sample stations by Winston (1977). At lower salinities, *E. immersa* may be encountering some physiological and metabolic stress, reducing the energy available for skeleton construction and resulting in the reported decrease in aragonite and lower overall Mg incorporation in calcite. Conversely, in low-salinity conditions, *M. nitida* exhibits increased wt% MgCO$_3$ in its skeletal calcite and this may be evidence of the stable metabolic state of this species in diminished salinity conditions.

A potential explanation for the observed differences in mineral deposition between sites may be the hydrodynamic linkages between study locations. Orkney features a highly tidal hydrological regime. The main tidal flows come in and out of Scapa Flow from the northwest and the south, both at an average flow rate of ~0.25 m s$^{-1}$ (Marine Scotland 2012). The island cluster in the west of Scapa Flow causes a complex funneling of the tidal current between these sites. Fara, in particular, creates an almost cyclical flow around the island at all points of the tide and the south of Flotta is prone to eddies, as the southern tidal in-flow meets the ebbing flow from Fara (Orkney Islands Council 2013).

The least mineralogically distinct site was the Barrel of Butter, Site 5, which is in the centre of Scapa Flow and is, therefore, subject to incoming flow, larvae and gametes from both the south and the northwest. It is particularly indistinct from Flotta, Site 3, with no statistically significant difference in any species for either wt% MgCO$_3$ in calcite or wt% aragonite in CaCO$_3$, despite its geographical distance from Flotta (>7.5 km). Lecithotrophic larvae from the temperate bryozoan *Celleporella hyalina*, which are of a comparable size to the species used in this study, were found to be able to swim for up to 4 h, with a preference to settle within 1 h of release (Goldson et al. 2001). If we expect similar planktonic behaviour for our comparably sized larvae, then during periods of peak tide, the dispersal potential would be up to 14.4 km, with distances of over 3.6 km achieved in 1 h. This would allow transport of larvae from Flotta to the Barrel of Butter in just over 2 h, well within the feasible time limit for successful metamorphosis upon settlement (Wendt 2000). It is therefore possible that the mineralogical indistinctness between the Barrel of Butter and Flotta is caused by population interconnectivity. The most mineralogically distinct sites were South Cava (Site 1) and Holm of Houton (Site 2). The cyclical tidal flow around Fara appears strong at all points of the tide (Orkney Islands Council 2013) and may be acting to isolate South Cava and Holm of Houton from the south of Scapa Flow, helping to explain why all species from these sites are mineralogically distinct from Flotta populations. Population genetics and calculation of the level of gene flow between sites would need to be conducted on bryozoans from all sites to test this hypothesis.

**CONCLUSIONS**

Using mineralogical analysis, we detected a high degree of variability in the wt% MgCO$_3$ in calcite and wt% aragonite in the skeletons of marine bryozoans sampled from Orkney, Scotland. Among species differences in mineralogy provide evidence for
the vital effect in some temperate bryozoans. Significant variability was found to occur between sites and, in some cases, was found to be related to seawater temperature, salinity and carbonate system parameters. Detected patterns between mineralogy and environmental conditions were inconsistent, however, and we suggest that differences in population connectivity and physical hydrography between sites may be additionally influencing skeletal mineralogy. Although the underlying mechanisms remain unclear, we see that physiological processes may be overriding or masking environmental patterns in some bryozoan species. Our study emphasizes the importance of sampling replication and accurate measurement of environmental conditions, as well as consideration of species and site-specific biological factors when discussing mineralogical variability. Exploring the role of both biological and environmental factors in the control of calcium carbonate mineralogy should continue to provide a fertile area for research in the future.

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LITERATURE CITED


Chlanger GV (1962) Dependence on temperature of Ca/Mg ratio of skeletal structures in organisms and direct chemical precipitates of sea water. Bull South Calif Acad Sci 61:45−61

Clarke FW, Wheeler WC (1922) The inorganic constituents of marine invertebrates. US Geol Surv Prof Pap 124:34–36


Chlanger GV (1962) Dependence on temperature of Ca/Mg ratio of skeletal structures in organisms and direct chemical precipitates of sea water. Bull South Calif Acad Sci 61:45−61

Clarke FW, Wheeler WC (1922) The inorganic constituents of marine invertebrates. US Geol Surv Prof Pap 124:34–36


