Change in coccolith morphology by responding to temperature and salinity in coccolithophore *Emiliania huxleyi* (Haptophyta) isolated from the Bering and Chukchi Seas

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Abstract

Strains of the coccolithophore *Emiliania huxleyi* (Haptophyta) collected from the subarctic North Pacific and Arctic Oceans during the R/V *MIRAI* cruise in 2010 (MR10-05) were established as clone cultures and have been maintained in the laboratory at 15 °C and 32 ‰ salinity. To study the physiological responses of coccolith formation to changes in temperature and salinity, growth experiments and morphometric investigations were performed on two strains of MR57N isolated from the northern Bering Sea (56°58′ N, 167°11′ W) and MR70N at the Chukchi Sea (69°99′ N, 168° W). This is the first report of a detailed morphometric and morphological investigation of Arctic Ocean coccolithophore strains. The specific growth rates at the logarithmic growth phases in both strains markedly increased as temperature was elevated from 5 to 20 °C, although coccolith productivity (the percentage of calcified cells) was similar at 10–20 % at all temperatures. On the other hand, the specific growth rate of strain MR70N was affected less by changes in salinity in the range 26–35 ‰, but the proportion of calcified cells decreased at high and low salinities. According to scanning electron microscopy (SEM) observations, coccolith morphotypes can be categorized into Type B/C on the basis of their biometrical parameters, such as length of the distal shield (LDS), length of the inner central area (LICA), and the thickness of distal shield elements. The central area elements of coccoliths varied from grilled type to closed type when temperature was increased or salinity was decreased, and coccolith size decreased simultaneously. Coccolithophore cell size also decreased with increasing temperature, although the variation in cell size was slightly greater at the lower salinity level. This indicates that subarctic and arctic coccolithophore strains can survive in a wide range of seawater temperatures and at lower salinities due to their marked morphometric adaptation ability. Because all coccolith biometric parameters followed the scaling law, the decrease in coccolith size was caused simply by the reduced calcification. Taken together, our results suggest that calcification productivity may be used to predict future oceanic environmental conditions in the Polar Regions.
1 Introduction

Sea-ice reduction due to global warming has become a major concern in the Arctic and Subarctic regions due to its induction of various environmental changes (e.g., Post et al., 2013; Wassmann et al., 2011). As a constituent of oceanic ecosystems, phytoplankton is an important primary producer and a key marker for understanding changes in the oceanic environment (e.g., Fujiwara et al., 2014; Harada et al., 2012). A large-scale change in the oceanic environment was observed as a climatic regime shift in the subpolar Pacific region, such as the Bering Sea, in 1976–1977 (Mantua et al., 1997). Siliceous diatoms are the dominant primary producers in that location (Tsunogai et al., 1979), but an increase in the population of the calcareous haptophyte *Emiliania huxleyi* is suggested by the alkenone biomarkers preserved in the oceanic sediments (Harada et al., 2012). The reduction of sea ice in the northern Chukchi Sea from 2008 to 2010 has influenced the phytoplankton distribution pattern (Fujiwara et al., 2014). The shorter sea ice retreat in 2008 resulted in haptophyte dominance in warm water, while the longer sea ice retreat in 2009 and 2010 led to prasinophytes predominating in cold water. Thus, the composition of marine phytoplankton communities is sensitive to environmental changes in oceanic environments.

The coccolithophore *E. huxleyi*, which belongs to the Prymneophyceae family in the Haptophyta, is one of the most investigated phytoplankton species because of its marked ability to fix carbon dioxide, which enables it to produce considerable quantities of biomass during blooms, having a marked impact on the global climate. It is broadly distributed from the equator to subpolar oceans (e.g., Beaufort et al., 2011; Hagino et al., 2011; Liu et al., 2009), and produces calcified scales called coccoliths. The distal and proximal shield elements, central opening size, and calcite crystals of coccoliths exhibit complex morphologies.

Young et al. (2003) systematized the morphotypes of coccoliths of coccolithophores. In *E. huxleyi*, three well-established morphotypes (Types A, B, and C) and two additional morphotypes (Types B/C and R) were categorized in addition to *E. huxleyi* var.
Coccolith morphology of Arctic Emiliania strains
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Introduction

According to McIntyre and Be (1967), Type A and Type C likely correspond to warm- and cold-water types, respectively, although Hagino et al. (2011) reported that Type C has not always been reported in cold-water environments. Recent studies performed in the Southern Ocean also suggest that coccolith morphotypes are distinct ecotypes in the coccolithophore E. huxleyi because Type A is abundant in warm and nutrient-poor water while Type B/C is abundant in cold and nutrient-rich water (Poulton et al., 2011).

The relationships between coccolith size and various environmental factors, such as temperature, salinity, and nutrients, have been investigated using E. huxleyi cultures (e.g., Watabe and Wilbur, 1966; Paasche, 2001; Fielding et al., 2009). Watabe and Wilbur (1966) reported that coccolith size decreased with increasing temperature; other authors have reported similar results for coccolithophore cell size (Sorrosa et al., 2005; De Bodt et al., 2010). Regarding the effects of salinity, Paasche et al. (1996) first reported that lower salinity was associated with a decrease in the length of the distal and proximal shield elements. Fielding et al. (2009) reported a linear correlation between salinity and the length of the distal shield. Phosphorous deficiency may induce...
over-calcification, while nitrogen limitation may result in the production of less-calcified coccoliths (Paasche, 1998).

In this study, the effects of temperature and salinity on coccolithophore growth and coccolith morphology investigated by SEM photometry were examined in two newly established strains of *E. huxleyi* isolated from the Bering Sea and Chukchi Sea during the *MIRAI* cruise (MR10-05) in 2010. There were marked changes in coccolith size and productivity (i.e., the percentage of calcified cells); we discuss the implications of this in relation to calcification productivity under future oceanic environments in the Arctic Ocean.

2 Materials and methods

Strains of the coccolithophore *E. huxleyi* (Lohman) Hay and Mohler were isolated at 56°58′ N, 167°11′ W, and 4 m water depth (MR57N) in the Bering Sea and at 69°99′ N, 168° W, and 10 m water depth (MR70N) in the Chukchi Sea. The samples were taken during the R/V *MIRAI* Arctic Ocean research cruise (MR10-05) organized by the Japan Agency for Marine-Earth Science and Technology (JAMSTEC) in August–October 2010. The strains were established as clones, but not axenic cultures at the University of Tsukuba, Japan. Currently, both strains are stored in the algal culture collection of the National Institute for Environmental Studies (NIES), Tsukuba, Japan (strain numbers: NIES3366 and NIES3362, respectively).

Stock cultures of the MR57N and MR70N strains were maintained in MNK medium (Noël et al., 2004) in a 100 mL glass Erlenmeyer flask with an air-permeable, porous, silicone cap under a light/dark regime of 16 h/8 h. Temperature was maintained at 4 °C in a water bath equipped with a thermocontroller. The cultures were illuminated by a white 20 W fluorescent lamp at a light intensity of about 40 µmol photons m⁻² s⁻¹. As controls, two other strains of *E. huxleyi* obtained from the culture collections were used. One was strain MS1 of coccolith morphotype A (alternative strain number, D2801-5), obtained from The Roscoff Culture Collection (Station Biologique De Roscoff, Roscoff,
France). The second was strain NIES1311 of coccolith morphotype O (Culture Collection of the National Bioresource Project in NIES at the Bering Sea in August 2002). Stock cultures of both strains were maintained at 15 °C in an incubator (MLR-350T; Panasonic Healthcare, Tokyo, Japan) under fluorescent lamps at a light intensity of 32–34 µmol photons m$^{-2}$ s$^{-1}$ before use in experiments.

Algal cells were transferred from stock cultures to pre-cultures and then grown to the stationary phase under the same conditions used for the subsequent experimental culture. Cultures involved three cycles of dilution and growth (three generations) to enable cells to acclimate to the experimental temperature or salinity conditions. Growth experiments were independently performed in triplicate in 200 mL glass conical flasks containing 100 mL culture medium. The culture medium was an artificial seawater Marine Art SF-1 enriched with ESM micronutrient-enrichments in which soil extracts were replaced with 10 nM (final concentration) sodium selenite (Danbara and Shiraiwa, 1999). Salinity was adjusted to 26, 32, or 35 ‰, while pH was fixed at 8.2. Final concentrations of nitrate and orthophosphates in the medium were 1.4 mM and 28.7 µM, respectively. Temperature was set at various values using an incubator (TG-180-5L, Nippon Medical and Chemical Instruments, Osaka, Japan). The culture was illuminated using fluorescent lamps under an incident photon flux density of 100 µmol photons m$^{-2}$ s$^{-1}$ with a light/dark regime of 16 h/8 h. The growth rate at each temperature was calculated as the average value of triplicate experiments, and the error bars indicated the minimum and maximum values.

At intervals, 1.5 mL cell suspension was harvested after gentle shaking every 2 days during the light period for enumeration of cells and preparation of samples for SEM observation. Cell counts were performed twice under a polarized microscope (BX-50, Olympus, Tokyo, Japan). The total number of cells, including both calcified and non-calcified (naked) cells, was determined. Samples for SEM observation were prepared by dropping 100 µL algal suspension on polycarbonate filters. After removing salts from the medium by washing with distilled water, the polycarbonate filters were dried on Whatman Nucleopore™ filters (GE Healthcare Japan, Tokyo, Japan). The polycarbon-
ate filters with attached cells were mounted on SEM holders using carbon paste and then coated with Pt–Pd (E-1045, Hitachi Power Solutions, Ibaraki, Japan) for SEM observation (6330F, JEOL, Tokyo, Japan).

For the photometric analyses, about 100 coccolithophore cells were observed by SEM per sample, and image analyses were performed using Image J (Image Processing and Analysis of Java: http://rsb.info.nih.gov/ij/).

3 Results

The MR57N and MR70N strains showed similar growth properties at 5 to 20°C (Fig. 1, Table 1). The final cell densities obtained at the stationary growth phase were about $1 \times 10^7$ cells mL$^{-1}$ for all *E. huxleyi* strains, suggesting that growth limitation during the stationary growth phase was due to nutrient depletion (Fig. 1, Table 1). The specific growth rate ($\mu$ value) of MR70N increased linearly with temperature from 5 to 2°C. The $\mu$ value at 5°C ($\mu = 0.31–0.29$ d$^{-1}$) was about 40% lower than that at 20°C ($\mu = 0.78–0.86$ d$^{-1}$). The $\mu$ value at 20°C was similar to that of other strains, such as MS1 and NIES1311, isolated from the North Sea of the Atlantic Ocean and the Bering Sea and which exhibited values of 0.76 and 0.63 d$^{-1}$, respectively. However, both the MS1 and NIES1311 strains did not grow at <10°C (data not shown). The growth rates of whole cells of the MR70N strain at salinities of 26 and 35‰ at 15°C were higher ($\mu = 0.6$ and 0.58 d$^{-1}$) than those at 32‰ ($\mu = 0.53$ d$^{-1}$) (Table 2). The growth rate of calcified cells increased with decreasing salinity from 0.32 to 0.42 d$^{-1}$.

The effect of temperature on calcification, namely coccolith productivity, was examined by monitoring the number of calcified and non-calcified cells. Interestingly, the numbers of calcified cells in cultures of strains MR57N and MR70N were lower than those of non-calcified (naked) cells with the approximate proportion of 8 to 26% (Fig. 1 and Table 1). Compared to the MR57N and MR70N strains, about half (56–41%) of MS1 and NIES1311 cells were calcified, indicating that *E. huxleyi* MR strains were
less extensively calcified under the culture conditions. The numbers of calcified cells decreased markedly to 1% at both lower and higher salinities (Table 2).

Morphometric parameters and the morphological properties of the newly established Bering and Chukchi strains MR57N and MR70N changed during culture under various conditions (Figs. 2–3). All measured parameters of cells and coccoliths of the MR57N and MR70N strains increased with decreasing temperature (Fig. 2d–g). The MS1 and NIES1311 strains cultured at 20°C showed similar morphometric parameters, with the difference that the number of distal shield elements in MS1 was slightly lower than that in NIES1311 (Fig. 2g). MR57N and MR70N cells exhibited reductions in size of 5.3–5.5 to 4.4–5.0 µm as temperature increased from 5 to 20°C (Fig. 2d). Moreover, average LDS values decreased from 4.10–4.15 µm at 5°C to 3.09–3.32 µm at 20°C (Fig. 2e). The LDS values of the MS1 and NIES1311 strains at 20°C were similar to those of MR70N, whereas MR57N exhibited slightly higher values (Fig. 2e). The LICA values of the MR57N and MR70N strains were almost identical and decreased with increasing temperature. The LICA values of the MS1 and NIES1311 strains were identical (about 1.4 µm on average), but smaller than those of the MR strains (1.6–1.7 µm on average) at 20°C (Fig. 2f). The number of distal shield elements decreased with increasing temperature; this trend was similar to the changes in LICA and LDS in the MR57N and MR70N strains. At 20°C, the numbers of distal shield elements in the MR57N and MR70N strains (37 and 35 on average, respectively) were greater than those in the MS1 and NIES1311 strains (30 and 32 on average, respectively) (Fig. 2g). Consequently, cell and coccolith sizes of both MR strains were larger than those of the MS1 and NIES1311 strains at 20°C.

Figure 3 shows the effects of increasing temperature (5–20°C) on the relationship between cell diameters and LDS in E. huxleyi strains MR57N and MR70N cultured at a salinity of 32‰. The sizes of both cells and coccoliths increased linearly with increasing temperature (Fig. 3a). The distribution of coccolith sizes overlapped with those of Types B, B/C, and C, which were defined previously by Young et al. (2003)
and Hagino et al. (2011) (Fig. 3a). Figure 3b is drawn as the schematic model of the correlated cell and coccolith sizes at the higher and lower temperature.

The morphology of coccoliths of both MR strains was characterized by fragile/delicate distal shield elements, a completely calcified or often partially grilled central shield element and a proximal shield element larger than the distal shield element (Fig. 2a–c). In addition, the length of the distal shield element (LDS) was 3–5 µm (3.3–4.3 on average) in cells cultured at various temperatures (Figs. 2e and 3a). Based on these properties, both the MR57N and MR70N strains can be classified as being of the Type B/C morphotype, which was defined previously by Young et al. (2003) and Hagino et al. (2011).

To further confirm the morphotype of MR strains, Fig. 4 shows the relationship between the width of the distal shield elements and LDS. The width of the distal shield elements for all strains were less than 0.1 µm that is the range of morphotype B/C determined by Cook et al. (2011). However, the width of the distal shield element in MS1 was larger than that of the other strains. Sincere MS1 is morphotype A, the dashed line might be the boundary between the morphotype A and morphotype B reported by Young and Westbroek (1991).

Because of the SEM observation of several central area morphology, we categorized coccolith and coccolithophore cell morphotypes into four sub-morphotypes (Types I–IV) and malformed types according to their morphological properties observed by SEM of E. huxleyi strains MR70N (Fig. 5). The definitions follow: Type I (Fig. 5a1 and a2), the central area elements are completely calcified; Type II (Fig. 5b1 and b2), the central area elements are partially calcified or exhibit grilled structure similar to the central area of morphotype A classified by Young et al. (2003) and Hagino et al. (2011); Type III (Fig. 5c1 and c2), the central area is open with a hole in the center but the marginal area is well calcified without spaces; Type IV (Fig. 5d1 and d2), the central area is open with a hole in the center and the other marginal area is not well calcified, showing grilled structure; malformed type (Fig. 5e2), the distal shield elements are not well calcified, showing an irregular morphology. Next, we designated “cell morphotypes” according to
coccolith type, which comprised the majority of cells (Fig. 5a3–e3). For instance, Type I cells consisted of about 60–80% of Type I coccoliths and 20–40% of the other types of coccolith; therefore, small amounts of various types of coccolith are produced by a single cell (Fig. 5a4). In contrast, cells with high proportions of coccoliths of various types were defined as “mixed types” to evaluate the proportion of the coccolithophore sub-morphotypes at each experiment (Fig. 6).

Figure 6 shows the proportion of submorphotypes of coccolithophore cells in *E. huxleyi* MR57N, MR70N, MS1 and NIES1311 strains. Strains MR57N and MR70N were nearly 100% Type II cells at 5°C; however, this proportion decreased with increasing temperature, which was accompanied by a increase in the proportion of Type I cells (Fig. 6). At 20°C, Type I cells made up 25 and 35% of strains MR57N and MR70N, respectively. About 10% of cells were classified as malformed or mixed type coccoliths. However, only 7 and 85% were Type I and II cells, respectively, in the MS1 strain cultured at 20°C. On the other hand, 85% of NIES1311 cells were Type O (defined by Hagino et al., 2011), the coccoliths of which have no central shield element. In addition, about 10% were malformed or incomplete coccoliths (Fig. 6).

The effects of salinity on coccolith morphometry and morphotype in strain MR70N at 15°C were shown in Fig. 7. The changes in the average LDS values ranged from 3.38 to 3.53 µm among salinities of 26, 32, and 35‰ (Fig. 7a), but cell diameters were larger at 26‰ salinity (Fig. 7b). Sub-morphotypes of MR70N cells were greatly affected by salinity during growth. The Type I and II subtypes made up about 40 and 25%, respectively, of all cells grown at a salinity of 26‰, but changed to about 2 and 70% at a salinity of 35‰ (Fig. 7c). As shown in Fig. 7d, there was a positive linear relationship between cell diameter and LDS, and cell diameter increased without change in LDS with decreasing salinity. One explanation of this relationship might be caused by the increase of cell diameter due to the increase of coccolith layers surrounding the cell.
4 Discussion

4.1 Effects of temperature on growth rate, coccolith morphometry, and morphology

The MR57N and MR70N strains exhibited growth at 5°C with μ values of about 0.3 d⁻¹ (Fig. 1); in contrast, other strains such as MS1 and NIES1311 did not grow. On the other hand, the μ values at 20°C of the four strains isolated from cold-water areas were identical (0.8 d⁻¹). The ability of microalgae to grow at low temperatures may be mostly due to their cold-water origin, as reported by Conte et al. (1998). Therefore, the ability of both MR strains to grow at 5°C seems to be due to their genetically fixed ability because their cold tolerance was maintained even after long-term storage as stock cultures at 15°C (see Materials and Methods). This temperature dependency of the two MR strains is similar to that of *E. huxleyi* strain L (NIOZ culture collection, Texel; originally isolated from the Oslo Fjord) reported by van Rijssel and Gieskes (2002), although the specific growth rate at 4°C was 0.12 d⁻¹, which is half that of the MR strains. According to Conte et al. (1998), some *E. huxleyi* strains isolated from cold-water regions can grow at 6°C (μ values, 0.3–0.75 d⁻¹), with variation in growth rates among strains. Both MR strains used in this study exhibited marked cold tolerance.

The numbers of calcified and non-calcified (naked) cells of strain MR70N increased logarithmically throughout the early stages of growth (Fig. 1). Around 10–20% of MR strains were calcified at all temperatures. This finding is similar to the results of Watabe and Wilbur (1966), who reported that 20–50% of cells were calcified, depending on temperature (a greater proportion of cells were calcified at 24°C compared to at < 24°C) in *Coccolithus huxleyi* strain BT-6 (present name, *Emiliania huxleyi*) isolated from the Sargasso Sea. In contrast to the MR strains, ~50% of cells in cultures of MS1 and NIES1311 were calcified (Fig. 1e and f). Thus, the calcification abilities of the cold-water strains vary, and MR strains are among the least calcified.

The decrease in cell size with increasing temperature is consistent with previous reports of *E. huxleyi* NIES837 (isolated from the Great Barrier Reef, Australia) and *E.
huxleyi AC481 (isolated from Normandy, France) by Sorrosa et al. (2005) and De Bodt et al. (2010), respectively. Calcium uptake in NIES837 strains was higher at lower temperatures (Sorrosa et al., 2005), while *E. huxleyi* AC481 coccolith morphology and morphometry were unaffected by temperature (De Bodt et al., 2010). Watabe and Wilbur (1966) found a correlation between temperature and coccolith size and growth rate, but not cell diameter. Thus the temperature dependence of coccolithophore growth and cell size was consistent among the strains, but coccolith formation differed by morphotype.

Regarding the MR strains, growth rate increased, but cell size and coccolith size decreased, with increasing temperature. All morphometric parameters followed the scaling law. Furthermore, morphology (such as the central area elements) changed from a completely calcified structure (Type I) at higher temperatures to a partially calcified grilled structure (Type II) at lower temperatures. This might be explained by enlargement of the coccolith due to the increased cell diameter (Fig. 3b). Type III and IV coccoliths, which exhibit an open central area) are similar to coccoliths observed in cells grown under P-limited conditions, as reported by Paasche (1998).

According to Young and Westbroek (1991) and Cook et al. (2011), the width of distal shield elements is also a useful parameter for classifying coccolith morphotypes. The relationship between the width of the distal shield elements and LDS was tested in the MR70N strain (Fig. 4). The MR strains had thin distal shield elements, categorized into Types B, B/C, and C. Concerning the ocean-geographical implications of these data, Type C and B/C strains are reported at higher latitudes in cold, sub-Antarctic oceans (Patil et al., 2014). On the other hand, Types A and B were found around the Southern Subtropical Front in a warm-water areas. In the Bering Sea, the lightly calcified Type A was identified during the bloom that occurred in August 2006 (Harada et al., 2012). Coccolith morphology in various *E. huxleyi* strains isolated from various oceanic areas (including in previous reports) is summarized in Table 3. Both the MR57N and MR70N *E. huxleyi* strains can be categorized as Type B/C, although both were isolated from cold waters: the Bering Sea and Arctic Sea, respectively.
4.2 Effects of salinity

Growth rate increased as salinity decreased from 32 to 26‰, which is in part consistent with Passche et al. (1996); however, the growth rates in this study (0.6–0.53 d\(^{-1}\)) were markedly lower than those reported by Passche. On the other hand, Fielding et al. (2009) reported an increase in growth rate from 0.05 to 0.7 d\(^{-1}\) with increasing salinity. The lower growth rate in their study might have been caused by use of a lower light intensity than that used by Passche et al. (1996).

The proportion of calcified MR70N cells cultured at 15°C decreased markedly when salinity was altered from 32‰ to either 26 or 35‰ (Table 2, Fig. 1g and h). The reduced calcification seems to be similar to the results of Fielding et al. (2009), because a salinity < 26‰ did not result in the sufficient production of coccoliths. On the other hand, Passceh et al. (1996) did not observe naked cells, even at 12‰ salinity. The coccolith productivity might be affected by the different light intensity used and also different types of coccolithophore strains.

Cell diameters and coccolith sizes differed slightly (Fig. 7), although there was no correlation between them. The cell diameter was greatest at the lowest salinity, while coccolith size was greatest at the highest salinity; the latter finding is consistent with previous reports (Passche et al., 1996; Fielding et al., 2009). The sub-morphotypes of larger coccoliths (LDS) also changed to Type II from Type I. This is consistent with the results of the temperature experiments, and indicates that sub-morphotype variation might be a strain-specific property.

Previous studies (Passche et al., 1996; Fielding et al., 2009) have considered the original oceanic environment of the strains, for example, coastal/marginal seas or oceans. The morphological and morphometric properties, and the relationships between LDS and temperature and salinity, in MR strains as well as other *E. huxleyi* strains were graphed together with findings reported previously (Fig. 8). Strains from the open ocean exhibited a strong correlation between LDS and temperature, while those from marginal waters showed a strong correlation between LDS and salinity.
4.3 Implications for the future polar oceanic environment

Growth rate and coccolith productivity are important oceanic environmental factors because these affect the biological and physical cycles of the ocean. The carbon cycle is particularly highly affected (Rost and Riebesell, 2004).

Global warming results in an increase in ocean temperature in the polar region, leading to melting of sea ice. This may lead to two scenarios in terms of *E. huxleyi* assemblages, as discussed by Bach et al. (2012). First, the present MR strains may remain dominant in these regions and respond physiologically to the environmental changes. Because two MR strains exhibited growth at 20°C to a degree comparable to the other strains and morphotypes (MS1 and NIES1311), this scenario is feasible. In this case, the present data can be directly applied to predict future conditions in the warmer polar region. An increase in the growth rate will result in higher biological activities in this region. Concerning calcification ability, temperature did not affect the proportion of calcified cells (Table 1), but all coccolith morphological parameters decreased with increasing temperature, and followed the scaling law. Thus an increase in oceanic temperature will result in a reductions in coccolith volume and calcification in this region. The reduced salinity caused by melting sea ice in the Arctic Ocean will facilitate growth of MR strains, the calcification abilities of which will be decreased by the reduction in coccolith production. Thus, higher temperatures and lower salinities will lead to reduced calcification by MR strains in this region.

The second scenario is that warmer-type strains or lower salinity-type strains other than MR strains become dominant in this region. According to their morphotype, the Bering Sea and Chukchi Sea *E. huxleyi* strains (MR57N and MR70N, respectively) can be classified predominantly as Type B/C. Moreover, the majority is of the Type II subtype when cultured at 5°C, but the population of Type II subtype cells decreases gradually and that of Type I subtype cells increases gradually as temperature is increased to 20°C. According to Poulton et al. (2011), the Type B/C morphotype has a lower calcite content (0.011–0.025 pmolC per coccolith) than Type A (0.015–
0.035 pmol C per coccolith). Furthermore, our data indicate that the coccolith productivity of MR strains is lower than that of Type A strains, such as MS1. In the case of the maximum different coccolith productivities between Type A (100% calcification) and MR strains (15% calcification), calcite production of Type A and MR strains are estimated as 0.035 and 0.0016 pmol C respectively. This estimation suggests that the maximum calcification may increase ~20-fold. On the other hand, if the abundance of lower salinity-type strains increases due the melting of sea ice, coccolith size may also decrease, as reported by Fielding et al. (2009). However, coccolith productivity may still affect more than the coccolith size reduction and the calcite production will increase about tenfold from 0.0016 pmol C (MR strains) to 0.015 pmol C (smaller Type A).

Type B/C represents a single, apparently cosmopolitan, population in the Southern Ocean (Cubillos et al., 2007). On the other hand, Triantaphylloue et al. (2010) reported that the size of *E. huxleyi* coccoliths in the Aegean Sea increased during cooler winter and spring periods. Different strains predominated during the different seasons, similar to the second scenario mentioned above. The morphotype population and the predominant strain in the studied area in the polar region are at present unknown. To facilitate the prediction of future environmental parameters, seasonal and morphotype variation in *E. huxleyi* should be elucidated.

## 5 Conclusions

Bering Sea and Chukchi Sea coccolithophore strains of *E. huxleyi* are capable of growth at a wide range of temperatures and salinities, and respond differently to different temperature and salinity conditions. We found that temperature affected the growth rates of both strains, and influenced coccolithophore cell size, coccolith size, and coccolith morphology. The MR70N strain exhibited reduced calcification and higher growth rates at lower and higher salinities, respectively, at 15°C. These results suggest that MR strains can adapt to various environments, including the low temperatures and low
salinities caused by the melting of sea ice in the Pacific Subarctic and Arctic Oceans. If these strains become dominant in this region, coccolith productivity will decrease, leading to an increase in the so-called biological pump. On the other hand, if other morphotypes become dominant in this region, calcification productivity will increase, leading to an increase in the biological pump. Thus, investigations of coccolithophores will enhance our understanding of the future environment in the polar region.

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Table 1. Temperature effect on specific growth rates ($\mu$) of various strains of the coccolithophore *E. huxleyi* isolated from the Bering Sea (MR57N), the Arctic Sea (MR70N), the North Sea (MS1) and the Bering Sea (NIES1311). In the whole culture, numbers of calcified cells with coccoliths and naked cells without coccolith were separately counted under microscope. Salinity of the medium was 32‰. All values are average of 3 separate experiments ($n = 3$).

<table>
<thead>
<tr>
<th>Growth temperature Strains</th>
<th>20 °C</th>
<th>15 °C</th>
<th>10 °C</th>
<th>5 °C</th>
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<tr>
<td></td>
<td>MR57N</td>
<td>MR70N</td>
<td>MS1</td>
<td>NIES1311</td>
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<tr>
<td>Whole culture</td>
<td>0.78</td>
<td>0.84</td>
<td>0.76</td>
<td>0.63</td>
</tr>
<tr>
<td>Calcified cells</td>
<td>0.64</td>
<td>0.66</td>
<td>0.65</td>
<td>0.81</td>
</tr>
<tr>
<td>Naked cells</td>
<td>0.82</td>
<td>0.99</td>
<td>0.89</td>
<td>0.52</td>
</tr>
</tbody>
</table>
Table 2. Effect of salinity on growth rates of the coccolithophore *E. huxleyi* isolated from the Arctic Sea (MR70N). For other conditions, see Table 1.

<table>
<thead>
<tr>
<th></th>
<th>Specific growth rate (µd⁻¹) at 15°C with salinity of</th>
<th>26‰</th>
<th>32‰</th>
<th>35‰</th>
</tr>
</thead>
<tbody>
<tr>
<td>Whole culture</td>
<td></td>
<td>0.60</td>
<td>0.53</td>
<td>0.58</td>
</tr>
<tr>
<td>Calcified cells</td>
<td></td>
<td>0.42</td>
<td>0.34</td>
<td>0.32</td>
</tr>
<tr>
<td>Naked cells</td>
<td></td>
<td>0.60</td>
<td>0.60</td>
<td>0.58</td>
</tr>
</tbody>
</table>
### Table 3. List of coccolith morphology of *Emiliania huxleyi* strains in laboratory culture experiments.

<table>
<thead>
<tr>
<th><em>Emiliania huxleyi</em> Strain</th>
<th>Coccolith morphology</th>
<th>Habitat isolated</th>
<th>Temperature</th>
<th>Salinity</th>
<th>Light intensity</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>MR70N</td>
<td>Type B/C</td>
<td>O</td>
<td>5, 10, 15, 20</td>
<td>26, 32, 35</td>
<td>32–34</td>
<td>Present study</td>
</tr>
<tr>
<td>MR57N</td>
<td>Type B/C</td>
<td>O</td>
<td>5, 10, 15, 20</td>
<td>32</td>
<td>*</td>
<td></td>
</tr>
<tr>
<td>BT6 (CCMP373)</td>
<td>Type C&lt;sup&gt;a&lt;/sup&gt;</td>
<td>O</td>
<td>7, 12, 18, 24, 27</td>
<td>35</td>
<td>~ 48</td>
<td>Watabe and Wilbur (1966)</td>
</tr>
<tr>
<td>DWN53/74</td>
<td>–</td>
<td>O</td>
<td>20</td>
<td>24, 34</td>
<td>200</td>
<td>Green et al. (1998)</td>
</tr>
<tr>
<td>G17 779Ga</td>
<td>Type A</td>
<td>O</td>
<td>20</td>
<td>24, 34</td>
<td>200</td>
<td>*</td>
</tr>
<tr>
<td>AC472</td>
<td>Type R</td>
<td>O</td>
<td>18</td>
<td>35</td>
<td>~ 100</td>
<td>Beaufort et al. (2007)</td>
</tr>
<tr>
<td>EHSO 5.30</td>
<td>Type A</td>
<td>O</td>
<td>16</td>
<td>–</td>
<td>70</td>
<td>Suellen et al. (2011)</td>
</tr>
<tr>
<td>EHSO 5.25</td>
<td>Type A</td>
<td>O</td>
<td>16</td>
<td>–</td>
<td>70</td>
<td>*</td>
</tr>
<tr>
<td>EHSO 5.28</td>
<td>Type A</td>
<td>O</td>
<td>16</td>
<td>–</td>
<td>70</td>
<td>*</td>
</tr>
<tr>
<td>EHSO 5.11</td>
<td>Type B/C</td>
<td>O</td>
<td>16</td>
<td>–</td>
<td>70</td>
<td>*</td>
</tr>
<tr>
<td>EHSO 6.17</td>
<td>Type B/C</td>
<td>O</td>
<td>16</td>
<td>–</td>
<td>70</td>
<td>*</td>
</tr>
<tr>
<td>EHSO 8.15</td>
<td>Type B/C</td>
<td>O</td>
<td>16</td>
<td>–</td>
<td>70</td>
<td>*</td>
</tr>
<tr>
<td>PLY B92/11</td>
<td>Type A</td>
<td>M</td>
<td>10, 15, 20</td>
<td>26–40</td>
<td>~ 30</td>
<td>Fielding et al. (2009)</td>
</tr>
<tr>
<td>PLY B92/11</td>
<td>Type A</td>
<td>M</td>
<td>20</td>
<td>24, 34</td>
<td>200</td>
<td>Green et al. (1998)</td>
</tr>
<tr>
<td>OF8</td>
<td>–</td>
<td>M</td>
<td>20</td>
<td>12</td>
<td>200</td>
<td>Paasche et al. (1996)</td>
</tr>
<tr>
<td>SC91</td>
<td>Type A&lt;sup&gt;b&lt;/sup&gt;</td>
<td>M</td>
<td>20</td>
<td>17, 25, 34</td>
<td>200</td>
<td>*</td>
</tr>
<tr>
<td>AC481</td>
<td>Type A&lt;sup&gt;b&lt;/sup&gt;</td>
<td>M</td>
<td>13, 18</td>
<td>35.6</td>
<td>150</td>
<td>De Bodt et al. (2012)</td>
</tr>
</tbody>
</table>

<sup>a</sup> Open (O) or Marginal (M) ocean.  
<sup>b</sup> Determined from SEM image published.
Figure 1. Growth responses of an Arctic strain of *E. huxleyi* (strain MR70N) to changes in temperature and salinity, (a) growth curves of *E. huxleyi* at 20°C and a salinity of 32‰; (b) at 15°C; (c) at 10°C; (d) at 5°C; (e) growth curves of *E. huxleyi* strain MS1 at 20°C; (f) growth curves of *E. huxleyi* strain NIES1311 at 20°C; (g) growth curves of *E. huxleyi* strain MR70N at 26‰ salinity; (h) growth curves of *E. huxleyi* strain MR70N at 35‰ salinity. Solid, gray and white symbols indicate whole culture (naked + calcified cells), non-calcified (naked) and calcified cells, respectively. (i) Effect of growth temperature on the specific growth rates of whole cells of *E. huxleyi* strains MR57N (squares), MR70N (diamonds), MS1 (triangles) and NIES1311 (crosses) at 32‰, and MR70N at 26‰ (asterisks) and 35‰ (circles). For μ values, see graphs (a–h) and Table 1.
**Figure 2.** Effects of temperature on cell morphology. (a) SEM images of strain MR70N grown at 20°C; (b) SEM images of strain MR70N grown at 5°C; (c) definitions of morphometric parameters of *E. huxleyi* cells: (d) cell diameter; (e) longer distal shield length (LDS); (f) long axis length of the inner central area (LICA); and (g) the numbers of distal shield elements in a coccolith. The MR1 and NIES1311 strains grown at 20°C were used as controls. Asterisk (*) and $N$ indicate the average value of each histogram and the number of samples determined, respectively.
Figure 3. (a) Changes in cell diameters and LDS in *E. huxleyi* strains MR57N and MR70N grown at 5, 10, 15, and 20°C, (b) schematic models of images of cell and coccolith sizes according to growth temperature. Descriptions of Type B, B/C, and C indicate the LDS range of coccoliths of the morphotypes defined by Young et al. (2003) and Hagino et al. (2011).
Figure 4. Relationship between the width of the distal shield elements and LDS in *E. huxleyi* strain MR70N grown at 5, 10, 15, and 20 °C and strains MS1 and NIES1311 grown at 20 °C. Area described with Type A indicates an area where sizes of Type A coccoliths distribute in literatures (Young and Westbroek, 1991; Cook et al., 2011).
**Figure 5.** Four sub-morphotypes (Type I to IV) of MR70N coccoliths, coccolithophores, and malformed cells were categorized by morphology on the basis of SEM images. (a1) Schematic drawing of Type I, whose central area elements are completely calcified, similar to the SEM image shown in (a2). (b1) Schematic of Type II, whose central area elements are partially calcified or with grilled spaces similar to the SEM image shown in (b2). (c1) Schematic drawing of Type III, whose central area is opened with a hole in the center with well-calcified marginal area, similar to the SEM image shown in (c2). (e1) Schematic drawing of Type IV, whose central area is opened with a hole in the center and a less-calcified marginal area, similar to the SEM image shown in (e2). An SEM image of the malformed type is shown in (e2); the distal shield elements are not well calcified and show an irregular morphology. (a3) to (e3) are coccolithophore cells of each coccolith type; histograms (a4) to (e4) indicate the proportions of the various coccolith morphotypes (see text).
**Figure 6.** Proportions of morphotypes of coccoliths and coccolithophore cells in *E. huxleyi* strains MR57N, MR70N, MS1, and NIES1311.
Figure 7. Influence of salinity on the morphometric parameters of *E. huxleyi* strain MR70N. (a) LDS; (b) cell diameter; (c) proportion of coccolithophore morphotypes; (d) relationship between cell diameter and LDS.
Figure 8. Relationships between LDS and temperature during growth (a) and LDS and salinity during growth (b) in various strains of *E. huxleyi*, including MR strains and other strains reported previously.