

## ***Interactive comment on “Change in coccolith morphology by responding to temperature and salinity in coccolithophore *Emiliana huxleyi* (Haptophyta) isolated from the Bering and Chukchi Seas” by K. Saruwatari et al.***

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Reply by authors (Jan. 29, 2016) to Anonymous Referee #1 (Received and published: 27 November 2015)

[General Comment] The authors reported interesting observations; clonal culture strains of *E. huxleyi* changed morphology and size of coccoliths in relation to change in temperature and salinity in laboratory culture experiments. Their observations are very interesting, however, I cannot evaluate accuracy of their experiments at this moment,

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since they did not describe details of their experiments in the Materials and Methods. Authors did not describe timing of measurements of cell growth rate and of size of coccoliths in their culture experiments, despite it is well known that growth rate of culture strain usually differs greatly between exponentially (logarithmic) and stationary growth phases, and size of coccoliths of *E. huxleyi* changes in relation to growth phase (growth rate); *E. huxleyi* make smaller coccoliths in the exponentially growth phase and make larger coccoliths in the stationary phase (Young and Westbroek, 1991). So I am unsure whether the observations on change in coccolith size in this study actually reflect change in temperature and/or salinity, or just reflect change in growth phase. Another problem; there are too many mistakes in citations. I would recommend authors add detailed information of experiments to materials and methods, reread related papers, and rewrite manuscript with correct references for resubmission.

[Reply] Thanks for the Referee to have pointed out the important fact that we did not clearly describe the timing of the cell growth and of the size of coccoliths. Now, we added the experimental results on cells from both early and late logarithmic growth for MR57N strains as Figure 7 (newly added). Both cell and coccolith sizes at the early logarithmic growth stage were larger than those at the late logarithmic growth stage. In addition, it was difficult to observed coccoliths on the cell at the stationary phase for those species. Therefore, we only compared the cell and coccolith sizes at the early logarithmic conditions. Another point that the referee gave is also important, so that we did reread the papers and rewrite the manuscript.

Followings are my other comments; [Comment 1]: Abstract has too much detailed information. The information on the name of the ship used for sampling (line 2), latitude and longitude of sampling localities (line 8), and explanations of classification of morphotype (lines 15–18, the sentence started from According: : :) are unnecessary here. [Reply 2] We revised the Abstract as suggested by referee.

[Comment 2]: Lines 23–25 of the page 17752 (p.2). Authors wrote “This indicates that subarctic and arctic coccolithophore strains can survive in a wide range of seawater

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temperatures and at lower salinities due to their marked morphometric adaptation ability” without explaining how ‘morphometric adaptation’ helps adaptation of *E. huxleyi* to various temperature/salinity conditions. Please explain it in the Discussion.

[Reply 2] Sorry for such misleading sentence. Now, we changed to: “This indicates that subarctic and arctic coccolithophore strains can survive in a wide range of temperatures and low salinities with change in their morphology.” (p. 2)

[Comment 3]: Lines 14-16 of the page 17753 (p. 3); Please describe the definition of ‘warm water’ and ‘cold water’ in the studied area (Arctic Ocean and Bering Sea).

[Reply 3] We added the information requested. So, “warm water” is  $\sim 5^{\circ}\text{C}$ , while “cold water” is  $< 0^{\circ}\text{C}$ . (p. 3)

[Comment 4]: Line 18 of the page 17753 (p. 3); Prymnesiophyceae not Prymneophyceae. More correctly, *E. huxleyi* belongs to the Family Noelaerhabdaceae, Order Isochrysidales, Class Prymnesiophyceae not to Prymneophyceae family.

[Reply 3] Thanks. We corrected to “Prymnesiophyceae”. (p. 3)

[Comment 5]: Lines 1-9 of the page 17754 (p. 4); Citations in these sentence are wrong. Authors wrote “Hagino et al. (2011) classified coccolith morphotype into four groups: (1) Type A and Type R with: : :.”. Correctly, Hagino et al. (2011) classified *E. huxleyi* into seven groups! Therefore, the all explanations concerning morphotypes of Hagino et al. (2011) in these sentences are inaccurate.

[Reply 5] Thanks for valuable comments and sorry for such a mistake. Now, we revised the text to “Hagino et al. (2011) classified coccolith morphotype into seven types, and further grouped into the four cross-sectioned types”. (p. 4)

[Comment 6]: Line 4 of the page 17754; ‘corona’ should be written in italic. [Reply 6] Thanks. We revised it on the line 4 and 8, as suggested.

[Comment 7]: Line 14 of the page 17754; ‘McIntyre and Bé’ not ‘McIntyre and Be’

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[Reply 7] We collected it in all parts of the text and reference.

[Comment 8]: Lines 14-16 of the page 17754; Citations in these sentences are completely wrong. Authors wrote “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”. Correctly, McIntyre and Bé (1967) just described warm and cold types of *E. huxleyi*. Young and Westbrook (1991) renamed warm and cold types of McIntyre and Bé (1967) as Types A and C, respectively. They renamed the morphotypes of *E. huxleyi* since Winter (1987) mentioned cold type (= Type C in Young and Westbrook 1991) was not always related to low temperature. Hagino et al. (2000) and Hagino et al. (2006) reported type C from tropical area, but Hagino et al. (2011) did not. Hagino et al. (2011) just introduced observation by Winter (1987) and interpretation by Young and Westbrook (1991).

[Reply 8] Thanks for valuable comments and sorry for such a misleading description. Now, we revised the sentences to: “Concerning the oceanographic distribution of Type A and Type C, defined by Young and Westbrook (1991), approximately correspond to warm- and cold-water types, described by McIntyre and Bé (1967), respectively, although Type C has not always been reported in cold-water environments (Young and Westbrook, 1991; Hagino et al., 2011).”

[Comment 9]: 2. Materials and methods; Please provide more detailed information on materials and methods of experiments.

[Reply 9] As requested, we added the more detailed information of the timing of the growth and the measurements of the sizes in “Materials and methods” and also “Results”.

[Comment 10]: Line 12 of the page 17755; How did you collect ‘samples’ that yielded your culture strains? Please show in situ seawater temperature and salinity of the water samples that yielded culture strains used in this study.

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[Reply 10] We added data on in situ seawater temperature and salinity of the water samples like the below database:

[Comment 11]: Line 17 of the page 17755; How did you establish clonal culture strain from your 'samples'?

[Reply 11] We added detailed description in the text (see Materials and methods) and also added a reference, Satoh et al., 2013.

[Comment 12]: Lines 20-21 of the page 17755; Authors wrote "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." I think the growth rate of *E. huxleyi* is usually changes during culture experiments. Please describe the detailed method used for monitoring of growth rate in this study, and provide information of growth phase of each culture strain at the timing of sampling for the studies of growth rate.

[Reply 12] As shown in Fig. 1, growth profiles in triplicate experiments were quite similar. To determine the timing of sampling for SEM observation, we firstly compared cellular SEM images between the early and the late logarithmic growth phase using MR57N stains (see Supplemental data or Fig. 7 (new)). The results indicated that both cell and coccolith sizes were larger in cells harvested at the early than the late logarithmic growth phase. Thus, we used cells harvested at the early logarithmic growth phase even for other strains.

[Comment 13]: Line 26 of the page 17755; How did you know the strain MS1 is type A?

[Reply 13] Hagino et al. (2011) described that MS1 is Type A. In addition, we also observed the curved central in MS1 (same as RCC1226) under SEM. Now, we revised the text by adding a reference of Hagino et al. (2011) in the text.

[Comment 14]: Line 26 of the page 17755; The strain code of MS1 in the Roscoff

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culture collection is RCC 1226 not D2801-5.

[Reply 14] D2801-5 is another name of MS-1, so that we changed to strain number "RCC1226" in the text.

[Comment 15]: Line 1 of the page 17756; How did you know the NIES 1311 is type O?

[Reply 15] Hagino et al. (2011) reported that NIES 1311 is type O. According to our SEM observation, the central area of NIES1311 is opened and agreed with Hagino et al. (2011). Now, we revised the text by adding the reference, Hagino et al. (2011).

[Comment 16]: Line 24 of the page 17756; How did you prepare sample for measurement of cell density in a polarized microscope?

[Reply 16] We added more detailed procedure in the text; "The numbers of cells in 10  $\mu$ L cell suspension, including both calcified and non-calcified (naked) cells, were determined using cell counting glass plate under a polarized microscope equipped with camera system and a calibration curve guaranteed by cell counting using the Thoma's haemocytometer. "

[Comment 17]: Lines 26-29 of the page 17756; Please describe pore size, diameter, and product name of the polycarbonate filter.

[Reply 17] We used a Isopore membrane filter with 0.8  $\mu$ m pore size, a Millipore product, ATTP04700. These information was added to the text.

[Comment 18]: Young and Westbroek (1991) reported size of coccoliths of *E. huxleyi* changes in culture experiments in relation to growth phase. Please provide information of growth phase of each culture strain at the timing of sampling for morphometric studies under SEM.

[Reply 18] The timing of sampling for morphometric studies were performed at the early logarithmic growth stage, mostly the 3rd day of the growth experiments, because of the below two reasons. Firstly, the reason is that the size of the coccolith and cell

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were larger at the early logarithmic stage for MR57N strains. Secondly, it is difficult to observe the coccoliths on the cell at the later steady growth stage using SEM.

[Comment 19]: Lines 2-10 of the page 17759; Young and Westbrook (1991) and Young et al. (2003) mentioned that central area of Type A consists of 'curved elements', while that of Type B (and B/C) consists of 'lath-like elements'. Authors classified their culture strains into type B/C without description of morphology of central area elements. So I am unsure if their strains are actually type B/C or not. Please describe morphology of central area elements of the culture strains used in this study.

[Reply 19] MR strains show "lath-like" central area, so that it is consistent with Young and Westbrook (1991) and Young et al. (2003).

[Comment 20]: Lines 10-11 of the page 17762; What is the 'scaling low'?

[Reply 20] "Scaling low" means that the morphometrical parameters are directly proportional.

[Comment 21]: Lines 22-23 of the page 17762 "On the other hand, Types A and B were found around the Southern Subtropical Front in a warm-water areas." Please provide information on papers that reported Type B from warm-water area.

[Reply 21] we added the reference, Patil et al. (2014).

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

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Abstract Strains of the coccolithophore *Emiliana huxleyi* (Haptophyta) collected from the subarctic North Pacific and Arctic Oceans in 2010 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, namely MR57N isolated from the northern Bering Sea and MR70N at the Chukchi Sea. 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°C to 20°C, although coccolith productivity (estimated as the percentage of calcified cells) was similar at 10–20% at all temperatures. On the other hand, the specific growth rate of 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. The central area elements of coccoliths varied from thin lath type to well-calcified lath-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 with change in their morphology. 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

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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 ( $\sim 5^{\circ}\text{C}$ ), while the longer sea ice retreat in 2009 and 2010 led to prasinophytes predominating in cold water ( $< 0^{\circ}\text{C}$ ). Thus, the composition of marine phytoplankton communities is sensitive to environmental changes in oceanic environments. The coccolithophore *E. huxleyi*, which belongs to the Family Noelaerhabdaceae, Order Isochrysidales, Class Prymnesiophyceae 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. *corona*. Hagino et al. (2011) classified coccolith morphotype into

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seven types, and further grouped into the four cross-sectioned types : (1) Type A and Type R with moderate to heavily calcified distal shields that are larger than the proximal shields, a grilled central area, and a length of distal shield (LDS) less than  $4\ \mu\text{m}$ ; (2) *E. huxleyi* var. *corona*, whose distal and proximal shields and central area are similar to those of Group (1) but whose central tube elements are elevated and whose LDS is  $3.5\text{--}4.5\ \mu\text{m}$ ; (3) Type B, Type B/C, and Type C, with lightly calcified distal shields that are smaller than the proximal shields and a fully calcified central area but their LDSs change from larger ( $> 4\ \mu\text{m}$ ) to smaller ( $< 3.5\ \mu\text{m}$ ); and (4) Type O, whose distal and proximal shields are similar to those of Group (3) but the central area is opened and lacks calcification. Young and Ziveri (2000) and Poulton et al. (2011) estimated the calcite contents of Types A, B, and B/C. Because the estimation is proportional to the cube of the coccolith shield length, calcite contents were in the following order from highest to lowest: Type B, Type A, and Type B/C. Concerning the oceanographic distribution of Type A and Type C, defined by Young and Westbroek (1991), approximately correspond to warm- and cold-water types, described by McIntyre and Bé (1967), respectively, although Type C has not always been reported in cold-water environments (Young and Westbroek, 1991; Hagino et al., 2011). 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 growth phase, temperature, salinity, and nutrients, have been investigated using *E. huxleyi* cultures (e.g., Watabe and Wilbur, 1966; Young and Westbroek, 1991; Paasche 2001; Fielding et al., 2009). Young and Westbroek (1991) investigated the size of coccolith at the end of growth phase, resulting that Type A coccolith is normally smaller than Type B coccolith. However, both types showed an overlapping size distribution and also a Type A strain (Strain L) unusually produces large coccolith in the late stationary growth phase. Watabe and Wilbur (1966) reported that coccolith size decreased with increasing temperature at the end of growth phase; other authors have reported similar results

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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 growth phase, 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** 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. Strain names established as clones of the coccolithophore *E. huxleyi* (Lohman) Hay & Mohler were MR57N and MR70N, respectively. Those strains, MR57N and MR70N, were isolated from seawater samples obtained at 56°58'N, 167°11'W (Station: s15), and 4 m water depth in the Bering Sea (sampling date: October 15th, 2010; in situ temperature and salinity: not recorded exactly, but SST at the nearest point determined on October 14th, 2010 is 3.6°C) and at 69°99'N, 168°W (Station: 166), and 10 m water depth in the Chukchi Sea (in situ temperature and salinity: 5.73°C and 31.22 ‰, respectively), respectively. Water samples were collected by a water-sampling system with CTD (Conductivity-Temperature-Depth profiler, 12 liters x 36 bottles, SBE911 Plus/Carousel, Sea-Bird Electronics, Inc., USA) and also a continuous monitoring system set at sea surface level in the monitoring laboratory on R/V MIRAI. Water samples were filtrated through a 300- $\mu$ m nylon mesh and then the filtrate water was used for preparing seawater for algal culture by mixing with seawater enriched with Erd–Schreiber's medium (ESM)

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containing 10 nM sodium selenite, instead of soil extracts usually contained (Danbara and Shiraiwa, 1999). Those water samples had been maintained under weak illumination with a regime of light /dark (16/8 h) at light intensity of 10  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup> and at 4°C on board. For isolation of coccolithophores, algal samples highly diluted with ESM-seawater had been maintained in microplates for about two months on board according to so-called the dilution method. Afterwards, tens of single cells of coccolithophores were isolated from sea water sample by picking up under microscope. The strains were established as clones according to our previous report (Satoh et al. 2013) at the University of Tsukuba, Japan, as described above, but those are not axenic cultures. 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  $\mu$ mol photons m<sup>-2</sup> s<sup>-1</sup>. As controls, two other strains of *E. huxleyi* obtained from the culture collections were used. One was strain MS1 of coccolith morphotype A (Hagino et al., 2011), obtained from The Roscoff Culture Collection (RCC1226; Station Biologique De Roscoff, Roscoff, France). The second was strain NIES1311 of coccolith morphotype O (Hagino et al, 2011), obtained from 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  $\mu$ mol photons m<sup>-2</sup> s<sup>-1</sup> 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

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glass conical flasks containing 100 mL culture medium. The culture medium was 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  $\mu$ M, respectively. Temperature was set at various values using an incubator (TG-180-5L, Nippon Medical & Chemical Instruments, Osaka, Japan). The culture was illuminated using fluorescent lamps under an incident photon flux density of 100  $\mu$ mol photons m<sup>-2</sup> s<sup>-1</sup> 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 numbers of cells in 10  $\mu$ L, including both calcified and non-calcified (naked) cells, were determined using cell counting glass plate under the microscope and then the total numbers of cells were extrapolated from them. Samples for SEM observation were prepared by dropping 100  $\mu$ L algal suspension on polycarbonate filters (ATTP04700, Isopore membrane filter with 0.8  $\mu$ m pore size, Millipore). After removing salts from the medium by washing with distilled water, the polycarbonate filters were dried on Whatman Nucleopore<sup>TM</sup> filters (GE Healthcare Japan, Tokyo, Japan). The polycarbonate 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). As first, the sizes of cell and coccolith of MR57N strain were investigated at the different timing of the growth at each different temperature (see supplement). Based on the first experimental results, the other morphometric experiments of other strains were performed at the early timing of the logarithmic growth condition. For the photometric analyses, about 100 coccolithophore cells were observed by SEM per sample, and image analyses were performed using

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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°C 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<sup>-1</sup> 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°C to 20°C. The  $\mu$ -value at 5°C ( $\mu = 0.31\text{--}0.29$  d<sup>-1</sup>) was about 40% lower than that at 20°C ( $\mu = 0.78\text{--}0.86$  d<sup>-1</sup>). 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 d<sup>-1</sup> and 0.63 d<sup>-1</sup>, 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<sup>-1</sup>) than those at 32‰ ( $\mu = 0.53$  d<sup>-1</sup>) (Table 2). The growth rate of calcified cells increased with decreasing salinity from 0.32 d<sup>-1</sup> to 0.42 d<sup>-1</sup>. 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 (Fig. 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–

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5.5  $\mu\text{m}$  to 4.4–5.0  $\mu\text{m}$  as temperature increased from 5°C to 20°C (Fig. 2d). Moreover, average LDS values decreased from 4.10–4.15  $\mu\text{m}$  at 5°C to 3.09–3.32  $\mu\text{m}$  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  $\mu\text{m}$  on average), but smaller than those of the MR strains (1.6–1.7  $\mu\text{m}$  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 lath-like central area 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  $\mu\text{m}$  (3.3–4.3 on average) in cells cultured at various temperatures (Fig. 2e, 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, Figure 4 shows the relationship between the width of the distal shield elements and LDS. The

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width of the distal shield elements for all strains were less than 0.1  $\mu\text{m}$  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. Since MS1 is categorized as 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. 5-a1 and a2), the central area elements are completely calcified; Type II (Fig. 5-b1 and b2), the central area elements are partially calcified or exhibit lath-like structure similar to the central area of morphotype B or C classified by Young et al. (2003) and Young and Westbroek (1991); Type III (Fig. 5-c1 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. 5-d1 and d2), the central area is open with a hole in the center and the other marginal area is not well calcified, showing lath-like structure; malformed type (Fig. 5-e2), 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. 5-a3-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. 5-a4). 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 sub-morphotypes of coccolithophore cells in *E. huxleyi* MR57N, MR70N, MS1 and NIES1311 strains which were harvested at the early logarithmic growth phase. Strains MR57N and MR70N were nearly 100% Type II cells at 5°C; however, this proportion decreased with increasing temperature, which was accompanied by an 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

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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 area element. In addition, about 10% were malformed or incomplete coccoliths (Fig. 6). When cell growth stage proceeded to the late logarithmic phase, the proportions of sub-morphotypes were changed even in the same strain of *E. huxleyi* MR57N, as shown in Figure 7. Cell diameter and LDS were increased proportionally by decreasing growth temperature, but no obvious change was observed by proceeding growth phase from the early to late logarithmic phases (Fig. 7a). In cells at the early logarithmic phase, Type II morphotypes were dominant at 5°C but substituted gradually with other morphotypes, especially Type I, by increasing growth temperature (Fig. 7b). On the other hand, Type II was dominant at 5°C but substituted by Type IV which became dominant at 20°C in cells at the late logarithmic phase (Fig. 7c). The effects of salinity on coccolith morphometry and morphotype in strain MR70N at 15°C were shown in Figure 8. The changes in the average LDS values ranged from 3.38 to 3.53  $\mu\text{m}$  among salinities of 26‰, 32‰ and 35‰ (Fig. 8a), but cell diameters were larger at 26‰ salinity (Fig. 8b). 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. 8c). As shown in Figure 8d, 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 Effects of temperature on growth rate, coccolith morphometry, and morphology The MR57N and MR70N strains exhibited growth at 5°C with  $\mu$ -values of about 0.3 d<sup>-1</sup> (Fig. 1); in contrast, other strains such as MS1 and NIES1311 did not grow. On the other hand, the  $\mu$ -values at 20°C of the four strains isolated from cold-water areas were identical (0.8 d<sup>-1</sup>). The ability of microalgae to grow at low temperatures may be

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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<sup>-1</sup>, 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 ( $\mu$ -values, 0.3–0.75 d<sup>-1</sup>), 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, 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 (Fig. 2-4) 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 mostly consistent among the strains, but coccolith formation differed by morphotype. Type I was dominant in MR57N and MR70N cells grown at 5°C and MS1 grown at 20°C although

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Type O was highly dominant in NIES1311 strain grown at 20°C (Fig. 6). 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, coccolith morphology (such as the central area elements) changed from a completely calcified structure (Type I) at higher temperatures to a partially calcified lath-like structure (Type II) at lower temperatures (Fig. 5, 6). 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 (Fig. 5), are similar to coccoliths observed in cells grown under P-limited conditions, as reported by Paasche (1998). In this study, morphometric parameters and morphology of whole cells and coccoliths were examined in cells harvested at the early logarithmic growth phase, as described above. However, in cells harvested at the late logarithmic stage, the proportion of Type II was over 60% at 5°C. However, Type IV was increased markedly with increasing temperature, especially high at 20°C, whereas Type I increased up to 25% (Fig. 7). 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, while Types A and B were found around the Southern Subtropical Front in a warmer-water areas (Patil et al., 2014). 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.

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

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(0.6–0.53 d<sup>-1</sup>) 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<sup>-1</sup> 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, 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, Passche 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. 8), 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. 9). 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.

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 increase in ocean temperature in the polar region, leading to melting of sea ice. This may lead to two scenarios in terms

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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 reduction 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 pmol C 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

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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, Triantaphyllou 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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Figure legends 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  $\mu$ -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.

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Figure 3. (a) Changes in cell diameters and LDS in *E. huxleyi* strains MR57N and MR70N grown at 5°C, 10°C, 15°C, 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°C, 10°C, 15°C, 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 lath-like 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. Relationships between LDS and temperature during growth (a) and proportions of morphotypes of coccoliths and coccolithophore cells in *E. huxleyi* strain

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MR57N harvested at the early (b) and late (c) logarithmic growth phases. The number below temperature in (b) and (c) indicate the date harvested after initiating culture. For morphotypes, refer Fig. 5.

Figure 8. 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 9. Relationships between LDS and cell diameter changed during growth (a) and LDS and salinity during growth (b) in various strains of *E. huxleyi*, including MR strains and other strains reported previously.

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