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**Coccolithophores
and CaCO₃
saturation of Baltic &
Black Seas**

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Coccolithophores and calcite saturation state in the Baltic and Black Seas

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Abstract

The Baltic and Black Seas are both brackish, that is to say both have salinities intermediate between freshwater and seawater. The coccolithophore *Emiliana huxleyi* is abundant in one, the Black Sea, but absent from the other, the Baltic Sea. Here we present summertime coccolithophore measurements confirming this difference, as well as data on the calcium carbonate saturation state of the Baltic Sea. We find that the Baltic Sea becomes undersaturated (or nearly so) in winter, with respect to both the aragonite and calcite mineral forms of CaCO_3 . Data for the Black Sea are more limited, but it appears to remain strongly supersaturated year-round. The absence of *E. huxleyi* from the Baltic Sea could therefore potentially be explained by dissolution of their coccoliths in winter, suggesting that minimum annual (wintertime) saturation states could be most important in determining future ocean acidification impacts. In addition to this potential importance of winter saturation state, alternative explanations are also possible, either related to differences in salinity or else to differences in silicate concentrations.

1 Introduction

Despite many decades of research, the ecological niche for coccolithophores remains poorly understood. By analogy with diatoms, which take up dissolved silicate for construction of their silica frustules (shells), and as a consequence become uncompetitive when levels of dissolved silicate run out (Egge and Aksnes, 1992; Brzezinski et al., 1998), coccolithophores might be expected to be affected by the availability of dissolved substances from which to build their calcium carbonate coccoliths (shells) (Merico et al., 2006). As far as inorganic precipitation and dissolution of calcium carbonate are concerned, the tendency for these two processes to take place is governed by the saturation state:

$$\Omega = [\text{CO}_3^{2-}] \cdot [\text{Ca}^{2+}] / K_{sp} \quad (1)$$

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with inorganic precipitation increasingly likely and/or rapid as Ω increases further above 1.0, and dissolution increasingly likely and/or rapid as Ω decreases further below 1.0.

If biogenic calcification were to be controlled in the same way as inorganic calcification, and if coccolithophore ecological success is contingent on being able to build coccoliths (the function of coccoliths remains poorly understood) then coccolithophores will become uncompetitive when levels of carbonate ion and/or calcium ion run low. It is still unclear whether this is actually the case. Two considerations suggest otherwise: (1) coccolith formation takes place in an internal vesicle, the chemistry of which need not be intimately connected to external seawater chemistry (Brownlee and Taylor, 2004), and (2) it is unclear whether bicarbonate ion (HCO_3^-) or carbonate ion (CO_3^{2-}) (isotopic constraints rule out CO_2 – Rost and Riebesell, 2004) is taken up as the substrate for calcification (Paasche, 2002).

This question of a possible connection between Ω and coccolithophore success is of considerable importance because of the current ongoing acidification of the oceans (Caldeira and Wicket, 2003, 2005; Orr et al., 2005). Falling pH is expected (depending on emissions) to lead to a halving of carbonate ion concentration by the year 2100, compared to pre-industrial levels, and therefore to a halving of saturation state (Zeebe and Wolf-Gladrow, 2001; Caldeira and Wicket, 2005; Orr et al., 2005). Here we investigate the likely impact of acidification on coccolithophores through comparison of their success in two seas of differing saturation state.

2 Absence of coccolithophores in the Baltic Sea

The first ever recorded observations of coccoliths were made by Christian Gottfried Ehrenberg in 1836. They were made in the Baltic Sea, although not in the water itself but rather from examination of fossil coccoliths in samples from the cliffs of the Isle of Rugen (Siesser, 1994). This island, in the southern Baltic Sea off the north coast of Germany, has distinctive white chalk cliffs, dating from the Cretaceous period when sea-levels were very much higher and the Baltic was not more or less cut off from the

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open sea as it is today. It is somewhat ironic that the first observations of coccoliths were made in ancient Baltic Sea sediments because today planktonic calcifiers are thought to be largely absent from the Baltic Sea in modern times. We tested this received knowledge (HELCOM, 2002) with direct targeted sampling for coccolithophores in the Baltic Sea proper (eastern Gotland Sea, ~55° N, 16° E) during a cruise on R/V Gauss in the period 19–27 July 2006.

For each of seven stations sampled, up to 1 litre (l) of seawater was filtered through a 47 mm diameter 0.4 µm pore size Millipore® Isopore membrane filter under low vacuum. After rinsing with alkaline water (pH~9.7) each filter was left to air-dry. Filters were stored in dark and dry conditions until analysis, at which point a small piece (about 1 cm²) of each membrane filter was cut radially, placed on a stub and gold-coated. Counting of the coccospheres under a Leo 1450VP (variable pressure) scanning electron microscope (SEM) was carried out with a computer-controlled stage and a fully automated image-capturing system. The software SmartSEM V05-01 was used to capture and store all images collected for each sample along a predefined meander-shaped transect. Each transect consisted of 225 view fields (total area of 2.5 mm²) at a magnification of x3000 and all images were stored on a CD-ROM. An example field-of-view, showing a portion of one of the filter papers as seen under the SEM, is displayed in Fig. 1.

Despite visual examination of more than 1500 single images (fields of view), not one coccolithophore or coccolith was seen on the filter papers. Numerous pennate diatoms were observed. Many collapsed non-mineralised cells were also seen, of unknown taxonomic affiliation. The sampling location was offshore (coccolithophores tend to be more successful in offshore than in coastal locations) and the time of year was also one at which coccolithophores are found to be present in high numbers in the open North Atlantic (Brown and Yoder, 1994; Holligan, 1993a; Raitso et al., 2006).

Our Baltic Sea analyses found no evidence of coccolithophores being present, in agreement with the received knowledge.

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3 Calcite saturation state in the Baltic Sea

Coccolithophores are abundant in the sub-polar North Atlantic (e.g. Holligan et al., 1993a; Brown and Yoder, 1994; Raitso et al., 2006), the Barents Sea (Smyth et al., 2004), the northern North Sea (Holligan et al., 1993b; Burkill et al., 2002) offshore and within the fjords of south-western Norway (e.g. Kristiansen et al., 1994) and even in the Skagerrak (Pingree et al., 1982; <http://www.noc.soton.ac.uk/soes/staff/tt/eh/pics/sat/skag2.jpg>), where the Baltic Sea meets the North Sea. It is therefore somewhat puzzling that they are absent from the Baltic Sea east of the transition area to the North Sea, given that they thrive in nearby waters at similar latitudes. Low salinity of the Baltic Sea may be considered as an obvious explanation for their absence; we argue below, however, that this may not be the critical reason.

In light of the current concerns over possible consequences of ocean acidification, we also consider whether the absence from the Baltic Sea could be explained by differences in CaCO_3 saturation state (Ω).

Comprehensive studies on the Baltic Sea CO_2 system have been performed during past years. These investigations concentrated on surface waters and were based mainly on measurements of the CO_2 partial pressure ($p\text{CO}_2$) and the total CO_2 concentration (C_T) (e.g. Schneider et al., 2003; Schneider and Kuss, 2004). Here we use this same data to determine the calcite/aragonite saturation. The CO_3^{2-} concentrations were calculated from the $p\text{CO}_2$ and C_T data using the equilibrium constants for the marine CO_2 system proposed by Roy et al. (1993) and the CO_2 solubility constants given by Weiss (1974). The Ca^{2+} concentrations (mmol kg^{-1}) were estimated from Ca/salinity relationships for the central Baltic Sea (Dyrssen, 1993: $\text{Ca}=0.331 \times \text{S}+0.392$) and for the Bothnian Bay in the uppermost north of the Baltic Sea (Gripenberg, 1960: $\text{Ca}=0.375 \times \text{S}+0.0368$). Although magnesium is less abundant in the Baltic Sea than in oceanic seawater, the molar Mg/Ca ratios are similar (~ 5 in both cases, Gunnars et al., 2004 and references therein) and therefore we estimate a minimal effect on K_{sp} . To calculate the solubility products for calcite and aragonite as

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a function of salinity and temperature, we used the algorithms given by Mucci (1983). The calcite/aragonite saturation was then determined on the basis of CO_2 measurements performed in 1997 (November) and 2001 (March, May, July) in a station grid (26 stations) in the central Baltic Sea (Gotland Sea). Data for March 2000 were also available for the Gulf of Riga and the Bothnian Bay. The results (Fig. 2) indicate a distinct seasonality of the mean Ω in the central Baltic Sea for both calcite and aragonite. The minimum is observed in March during the culmination of the convective mixing with Ω of about 1.0 and 0.6 for calcite and aragonite, respectively.

3.1 Cause of low wintertime Ω

Such low winter values do not occur in oceanic surface waters at similar latitudes, which are significantly oversaturated with regard to both calcite and aragonite throughout the year. This is a consequence of lower CO_3^{2-} and Ca^{2+} concentrations in the central Baltic Sea that lower Ω (see Eq. 1) despite the decrease of K_{sp} by a factor of about 7 at a salinity of 7 (Mucci, 1983). The state of the marine CO_2 system is controlled by a complex interplay between physical and biogeochemical variables. Lower salinities and winter temperatures in the Baltic Sea ($S \approx 7$, $T \approx 0^\circ\text{C}$) lead to dissociation constants for carbonic acid that are considerably below those in mid-latitude oceanic regions ($S \approx 35$, $T \approx 8^\circ\text{C}$) and thus cause reduced CO_3^{2-} concentrations (Baltic Proper average of $126 \mu\text{mol kg}^{-1}$ in summer, $29 \mu\text{mol kg}^{-1}$ in winter). This effect is compounded by the alkalinity that in most regions of the Baltic Sea is lower ($\approx 1600 \mu\text{Equiv kg}^{-1}$ at $S=7$) than in oceanic surface waters ($\approx 2320 \mu\text{Equiv kg}^{-1}$ at $S=35$). Finally, deep convective mixing during winter generates surface CO_2 partial pressures in the Baltic Sea that are above $400 \mu\text{atm}$ whereas CO_2 in comparable oceanic waters is close to equilibrium with the atmosphere. In total, these differences produce CO_3^{2-} concentrations in the Baltic Sea surface water that during winter are about 5 times lower than those in the North Atlantic ($\approx 140 \mu\text{mol kg}^{-1}$). Additionally, Ca^{2+} concentrations in the Baltic Sea ($\approx 3 \text{mmol kg}^{-1}$ at $S=7$) are about 4-fold lower than in oceanic waters and further reduce

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Ω to values of about 1 for calcite and 0.5 for aragonite.

3.2 Cause of large seasonal variation in Ω

The seasonal amplitude of Ω is also unusually high in the Baltic Sea (range of about 4, Fig. 2) in comparison to open ocean locations (range of between 1 and 2: Findlay et al., 2007). This is a consequence of contrasting seasonality of temperature, differences in water chemistry, and differences in the dominant phytoplankton.

In the Baltic Sea the temperature difference between the minimum in February and the maximum in July is about 20°C whereas the corresponding difference is less than half as great as this in mid-latitude oceanic waters. The consequence is a larger seasonality of the CO_3^{2-} concentration in the Baltic Sea.

Additionally, the removal and addition of CO_2 by production and decomposition of organic matter affects saturation states much more strongly in the Baltic Sea. Rather similar phosphate drawdowns occur from winter to summer ($0.5 \mu\text{mol kg}^{-1}$) in the Baltic Proper and in the North Atlantic at 50–60° N, 20° W (Tyrrell and Taylor, 1996). Assuming similar $\text{C}_{\text{org}}:\text{P}$ at the two locations, PO_4 removal would be accompanied by similar drawdowns of C_T . According to Eq. (1) above, saturation state is equal to $([\text{Ca}^{2+}]/K_{sp})$ multiplied by $[\text{CO}_3^{2-}]$. As just discussed, K_{sp} is about 7-fold lower in the Baltic, but $[\text{Ca}^{2+}]$ is only about 4-fold lower. Therefore, because the term $([\text{Ca}^{2+}]/K_{sp})$ in the Baltic is almost double the value in the Atlantic, similar absolute changes in $[\text{CO}_3^{2-}]$ (e.g. driven by carbon removal during spring blooms) produce much greater absolute changes in saturation state in the Baltic. By a similar reasoning, the effect of anthropogenic CO_2 on seawater Ω will be much stronger in the Baltic Sea than in oceanic waters.

$\text{C}_{\text{org}}:\text{P}$ is, however, not similar at the two locations, contrary to the assumption above. In the Baltic Sea about 50% of the phosphate is used by nitrogen-fixing cyanobacteria because there is a high excess of phosphate over nitrate (relative to the Redfield ratio) in the Baltic Sea winter water. Several studies have shown that cyanobacteria produce

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organic matter with high $C_{\text{org}}:P$ ratios (e.g. Schneider et al., 2003: about 300 and Larsson et al., 2001: about 400, compared to the Redfield ratio of 106), thus removing much more C_T , leading to greater seasonality in $[\text{CO}_3^{2-}]$.

3.3 Cause of regional variability in Ω

5 The previous considerations referred to the central Baltic Sea which can be considered as a mixing chamber for inflowing water from the North Sea and from the various rivers. Differing conditions with regard to the state of the CO_2 system can be found in semi-enclosed estuarine areas of individual rivers. A prominent example is the Gulf of Riga where water from the River Daugava and from the central Baltic Sea are mixed.
10 River Daugava water originates from the limestone-rich watershed area in continental Europe and has alkalinities of up to $3000 \mu\text{mol kg}^{-1}$. As a consequence, increased CO_3^{2-} concentrations are observed in most areas of the Gulf and cause calcite oversaturation (Fig. 2: $\Omega=1.3$ at $S\approx 5$) even during wintertime. A contrasting example is the Bothnian Bay which is influenced by rivers that have crossed the Scandinavian shield.
15 This watershed area consists mainly of igneous rocks which hardly contribute to river water alkalinity ($200 \mu\text{mol kg}^{-1}$). Hence, the Gulf of Bothnia water is characterized by low CO_3^{2-} concentrations resulting in a distinct calcite undersaturation during wintertime (Fig. 2: $\Omega=0.2$ at $S\approx 5$).

4 Discussion

20 4.1 Comparison to other phytoplankton sampling in the Baltic Sea

We checked the findings of our work, and of HELCOMM (2002), against other work. In his book on Baltic Sea algae, Helmut Pankow (1990) lists 9 genera of coccolithophores occurring in the Baltic Sea (*Acanthoica*, *Balaniger*, *Calciopappus*, *Discosphaera*, *Emiliana*, *Hymenomonas*, *Pappomonas*, *Papposphaera*, and *Trigonapsis*). For most of

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these genera, the occurrence is limited to the Danish Seas, which is likely to reflect an inflow from North Sea waters. *Emiliana huxleyi* is also reported from the Kiel Bight, but again this may correspond to inflow of North Sea water. No indication is given of abundances or frequencies of occurrence. In terms of waters unaffected by Atlantic influence, *Pleurochrysis carterae*, *Hymenomonas roseola*, *Balaniger balticus* and *Pappomonas virgulosa* have been found in the northeastern Baltic Sea. In his comprehensive listing of Baltic Sea phytoplankton species, Hällfors (2004) lists only four species (*Coccolithus pelagicus*, *Pleurochrysis carterae*, *Pappomonas lepida* and *Balaniger balticus*) as occurring in the Baltic Sea (excluding the Kiel Bight, where many more species have been seen). *C. pelagicus* is unusual in that it is the only coccolithophore species that has been seen in the Baltic Sea proper; it has also been found in the Bothnian Sea and Bothnian Bay as well as in the Gulf of Finland. Interestingly, in laboratory culture experiments *C. pelagicus* appears unusually robust in as much as its calcification is hardly affected by the low saturation states associated with high CO₂ (Langer et al., 2006).

Samples taken in Kiel fjord occasionally contain individual cells of coccolithophores, but always in very low numbers (U. Sommer and A. Stühr, personal communication). In routine weekly phytoplankton countings by Jeanette Goebel (personal communication) of the Landesamt für Natur und Umwelt in Schleswig-Holstein, blooms of *E. huxleyi* were recorded in the mid 1980s and again in the early 1990s. These were probably confined to the Danish waters and the Kiel Bight.

An examination of the literature thus slightly modifies the conclusion we obtained from our own sampling. Rather than being completely absent, it seems that coccolithophores are instead occasionally present (although extremely scarce) in the Baltic Sea. To our knowledge there have never been any observations of coccolithophore blooms or high abundances in the Baltic Sea proper, although coccolithophores may occasionally be missed due to the common practice of preservation with acidic Lugols (which dissolves coccoliths). They have always been found to be either completely absent or else a very minor component of the community. The only exception is in the

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Danish Seas and Kiel Bight, in waters potentially affected by the inflow from the North Sea.

4.2 Comparison to other carbonate system measurements

4.2.1 Global

5 The GLODAP dataset (Key et al., 2004) is a global dataset of carbonate chemistry (DIC and Alkalinity) measurements from open ocean cruises, much of it collected as part of the World Ocean Circulation Experiment (WOCE) and Joint Global Ocean Flux Study (JGOFS) programmes during the 1990s. The data points are combined together to give a global distribution of the carbonate chemistry. Seasonal cycles are not considered.

10 Only an average annual distribution is presented, and possible biases due to sampling of many locations at only one time of year (e.g. scarcity of winter measurements at high latitudes) are ignored. For the open North Atlantic at 60° N, 20° W (at a similar latitude to the Baltic Sea proper), the carbonate ion concentration derived from the GLODAP dataset (together with temperature and salinity taken from Levitus climatologies) is

15 about 170 $\mu\text{mol kg}^{-1}$, leading to a calcite saturation state of about 4. This compares to global (70° S–60° N) ranges of about 80–300 $\mu\text{mol kg}^{-1}$ for carbonate ion (Orr et al., 2005) and about 2–7 for calcite saturation state.

4.2.2 High-latitude seas exchanging freely with the open ocean

20 A combined ecosystem/carbon chemistry model of the eastern Bering Sea (Merico et al., 2006) first examined the seasonal cycle of carbonate chemistry. At this high-latitude site (56.8° N, 164° W), the modelling suggested considerable seasonal variation in both carbonate ion (100–150 $\mu\text{mol kg}^{-1}$) and calcite saturation state (2.5–3.5). The values of both parameters were found to be low in winter-time, to rise sharply at the time of the spring blooms, and then to stay at high values through the summer period until

25 values decline in autumn as mixing depths increase and deep waters are mixed up

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into the surface. It was not possible to fully test this model prediction against data from the eastern Bering Sea. Seasonal in-situ data are, however, available from the Norwegian Sea shelf break (Ocean Weather Station “Mike”, 66° N, 2° E), and agree in outline with the model predictions for the eastern Bering Sea. The Norwegian Sea data show a range in carbonate ion of 130–180 $\mu\text{mol kg}^{-1}$ and in calcite saturation state of 3–4, again with a minimum in winter and a maximum in summer (Findlay et al., 2007). The seasonal pattern is caused by the spring diatom blooms, because the intense DIC removal has the side-effect of driving up carbonate ion concentrations (Merico et al., 2006). The seasonal variation in SST is of only second-order importance in determining the seasonal cycle of carbonate ion in the eastern Bering Sea (Merico et al., 2006).

From this comparison to other data it appears that the Baltic Sea exhibits a similar general seasonal pattern to the Norwegian Sea and eastern Bering Sea, in terms of carbonate system parameters. The amplitude of the seasonal range is, however, much larger than in the open-sea locations. Summertime saturation states are high, but wintertime saturation states are extremely low, lower even than in polar oceanic waters. The greater seasonal amplitude derives from the unusual chemistry of the Baltic Sea and the large annual temperature range.

4.2.3 The Black Sea

There is less data from which to calculate CaCO_3 saturation states in the Black Sea. Goyet et al. (1991) measured the carbonate chemistry at a few stations in the Black Sea in 1988, during cruise 134 of R. V. Knorr. Although the measurements were made at the same time of year (June) as the *Emiliana huxleyi* blooms occur, they were made many years before the first satellite observations of *Emiliana huxleyi* blooms in the Black Sea (Cokacar et al., 2004). However, the presence of a thick layer of coccoliths in surface sediments of the Black Sea (Hay, 1988) shows that coccolithophores have also been present before the last few years. Total inorganic carbon (C_T) and total alkalinity (A_T) were measured, and other carbonate system parameters derived from

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them. Simultaneous surface-layer measurements of both C_T and A_T were made at only one offshore station, station 6, at 43° N, 34° E in the central Black Sea. At this station the derived surface water concentration of carbonate ion was $250 \mu\text{mol kg}^{-1}$, and calcite saturation state was 7.5.

5 As discussed above, the seasonal amplitude in saturation state is governed in large part by the intensity of winter mixing (which stirs up nutrients), the consequent seasonal drawdowns of nitrate and phosphate (actually the seasonal drawdown of C_T is critical, but is related to the others through elemental stoichiometry of organic matter), and therefore to the size of the spring blooms. Because it is closer to the equator
10 than is the Baltic Sea, there is a rather muted seasonal cycle in comparison. Only a modest amount of winter mixing and nutrient stirring takes place in winter, with, in consequence, winter/spring blooms that are of small magnitude in relation to those in the Baltic Sea (Oguz et al., 2001).

There is only limited data to confirm relatively high wintertime saturation states, but
15 it can in any case be inferred on the following grounds: (1) relatively high summertime values, and (2) low-amplitude seasonal variation in phytoplankton and nutrients, which will drive only a low amplitude seasonal variation in $[\text{CO}_3^{2-}]$ and Ω (Merico et al., 2006). Surface measurements in May 2001, in offshore waters of the southwestern Black Sea near to the Bosphorus (Hiscock and Millero, 2006), gave rather similar results to Goyet
20 et al. (1991) for C_T , A_T and calcite saturation state (again about 7.5)

4.3 No calcifiers because of winter CaCO_3 undersaturation?

CaCO_3 saturation state generally declines with increasing latitude (Orr et al., 2005; Caldeira and Wickett, 2005), such that undersaturation of surface open ocean waters will be reached first in polar waters, as fossil fuel emissions intensify ocean acidification
25 in the future. However, although lower than summertime values, wintertime saturation states are still considerably supersaturated (i.e. values significantly greater than 1) in: (a) the open Atlantic south of Iceland (calculated from data in Takahashi et al., 1985), (b) in the Norwegian Sea (Findlay et al., 2007), and (c) in the eastern Bering Sea

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according to model calculations (Merico et al., 2006).

The Baltic Sea is therefore anomalous in comparison to oceanic sites at similar latitudes in that wintertime surface waters are distinctly undersaturated with respect to aragonite and borderline undersaturated with respect to calcite. Surface saturation states in summer, on the other hand, are higher than in comparable oceanic locations where coccolithophores thrive. We propose that the low wintertime saturation states may be the critical difference underlying the exclusion of coccolithophores from the Baltic Sea proper. If coccolithophores are unable to prevent dissolution of their coccoliths in undersaturated waters, and if the coccoliths are critical to their ecological competitiveness (their function is still unclear: Young, 1984), then an inability to survive through the winter would explain their absence in the Baltic Sea proper.

4.4 Alternative possible explanations

The ecological niche for *E. huxleyi*, and the factors responsible for blooms developing at some times and places but not others, have recently been reviewed (Tyrrell and Merico, 2004; Balch, 2004; Riebesell and Rost, 2004; Paasche, 2002), but are still relatively poorly understood.

4.4.1 Salinity

At first sight the most obvious reason why coccolithophores might be excluded from the Baltic Sea is the low salinity compared to oceanic waters. The surface water salinity in the eastern Gotland Sea, where our sampling was carried out, was about 7. Salinity varies considerably across the Baltic Sea: areas furthest from the North Sea (Bothnian Bay, Gulf of Finland) have lowest salinities (often less than 5), and salinity then increases towards the North Sea, eventually reaching values greater than 18 in the Kattegat, on the way out to the North Sea.

On the other hand, *E. huxleyi* “is one of the most euryhaline and eurythermal coccolithophore species” (Winter et al., 1994). *E. huxleyi* occurs at bloom concen-

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trations in the central Black Sea (e.g. Fig. 3; Cokacar et al., 2001), at salinities of 18–19, approximately half the average oceanic value of ~35. *E. huxleyi* has also been recorded from the northeastern Black Sea at salinity 15–17 (Mikaelyan et al., 2005) and from the Sea of Azov (Pitsyk, 1963) at salinity ~11 (Matishov et al., 2006; <http://www.nodc.noaa.gov/OC5/AZOV2006/start.html>). In both the Black Sea and the Baltic Sea inflows of deep saline water occur, but surface waters typically flow out to the North and Mediterranean Seas respectively.

In culture experiments it was found that one coastal and two oceanic strains of *E. huxleyi* were able to survive when grown at a salinity of 15, whereas another oceanic strain perished (Brand, 1984). Only the coastal strain grew at all rapidly at this salinity. One oceanic strain plus one coastal strain were tested at a salinity of 5, but neither survived (Brand, 1984). At face value these culture experiments suggest a lower salinity tolerance somewhere between 5 and 15. However, only a few strains were tested and others may be more tolerant of low salinities.

Because the salinity of the central Black Sea (~18) is significantly higher than that of the Baltic Sea proper (~7), salinity remains a possible explanation of the data discussed so far, although *E. huxleyi* success in the Sea of Azov (~11) makes this less likely.

4.4.2 Temperature

Coccolithophores are on average most successful (in terms of diversity and fraction of the total phytoplankton community) in warm, oligotrophic, low-latitude waters (Winter et al., 1994). It has been suggested that temperature itself plays a direct role in the success of the group, although there is little hard data in support of this contention. *Emiliania huxleyi* thrives even in the relatively cold waters of the North Atlantic south of Iceland, the Patagonian Shelf and even the Barents Sea (Smyth et al., 2004); in the latter location monthly average temperatures vary between ~2°C in winter and ~8°C in summer (<http://www.nodc.noaa.gov/OC5/barsea/barindex1.html>). Although the Black Sea is indeed warmer than the Baltic Sea, we do not consider this as a likely reason for

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the difference in success between the two seas. Wintertime temperatures in the Baltic Sea proper (e.g. $\sim 0^{\circ}\text{C}$) are not very different from those in the Barents Sea, whereas summertime temperatures in the Baltic proper (e.g. $\sim 20^{\circ}\text{C}$) are in fact much warmer than those in the Barents Sea.

5 4.4.3 Light

Blooms of *E. huxleyi* always occur in shallow mixed layers of maximum depth 20 or 30 m (Nanninga and Tyrrell, 1996; Tyrrell and Merico, 2004 and references therein); high light intensities appear to be a requirement for the development of *E. huxleyi* blooms (Nanninga and Tyrrell, 1996). Although the Black Sea has shallow mixed layers in summer, the same is also true for the Baltic Sea proper, in which a summer pycnocline is typically present at about 10–20 m depth (e.g. Larsson et al., 2001). We therefore discount a difference in light levels as an explanation for the differences in *E. huxleyi* abundance.

4.4.4 Silicate

15 Diatoms have faster growth rates than other phytoplankton of similar size (Furnas, 1991) and, if silicate is available, typically outcompete other phytoplankton functional types. They form the major part of blooms in eutrophic locations: spring blooms in temperate latitudes, blooms near river mouths, and blooms in recently upwelled water. It is possible that the ecological niche for coccolithophores may therefore be defined in part by the conditions required to exclude the competitively superior diatoms, i.e. low concentrations of dissolved silicate in conjunction with non-limiting concentrations of other essential nutrients (Tyrrell and Merico, 2004).

25 Scarcity of silicate starts to become limiting to silica production and diatom growth at concentrations below about $3\ \mu\text{mol kg}^{-1}$ (Brzezinski et al., 1998, 2005). Levels of silicate in summer were originally greater than this several decades ago but have subsequently declined, in both the Baltic Sea (Papush and Danielsson, 2006; Rahm et al.,

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1996) and in the Black Sea (Humborg et al., 1997; Konovalov and Murray, 2001), probably because of river damming (e.g. Humborg et al., 1997, 2006) and eutrophication (e.g. Papush and Danielsson, 2006). At the present time in the Black Sea, summer silicate levels are sufficiently low as to potentially exclude diatoms from dominance (Humborg et al., 1997; Konovalov and Murray, 2001). In the Baltic Sea proper, on the other hand, although levels of silicate in the summer are lower than in the winter (Wulff and Rahm, 1988), it appears that spring blooms consume only a fraction of the winter silicate, leaving much behind to persist through into the summer (Wasmund et al., 1998). Year-round availability of dissolved silicate in the Baltic Sea might therefore be a plausible explanation for why coccolithophores have been unable to colonise it. However, given that the Black Sea appears to have only recently become silicate-poor in summer (Humborg et al., 1997; Konovalov and Murray, 2001), whereas the sediments demonstrate a much longer record of *Emiliana huxleyi* success (Coolen et al., 2006; Hay, 1988; Bukry, 1974), perhaps high levels of silicate are not automatically prohibitive to coccolithophore success.

5 Conclusions

From the data presented here, we suggest that the presence of the coccolithophore *Emiliana huxleyi* in the Black Sea but not the Baltic Sea could be attributed to the very low winter-time saturation states in the surface Baltic Sea (either at or below the threshold for undersaturation, for calcite and aragonite, respectively). Alternative explanations relate to lower salinity in the Baltic Sea, or to higher silicate concentrations. These findings have potential implications for the fate of coccolithophores under ocean acidification, although more information is needed from environments where saturation states are low but salinity is high and/or silicate is low. Our study highlights the possible significance of minimum annual (wintertime) saturation states in determining success of planktonic calcifiers.

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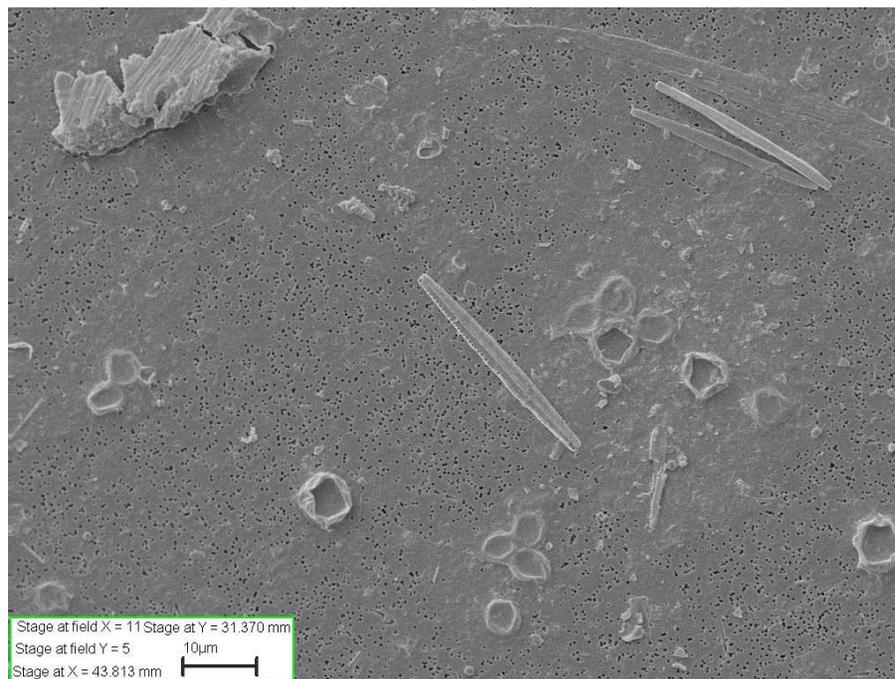


Fig. 1. Typical SEM field-of-view from Baltic Sea samples collected in the vicinity of (55° N, 16° E) during July 2006.

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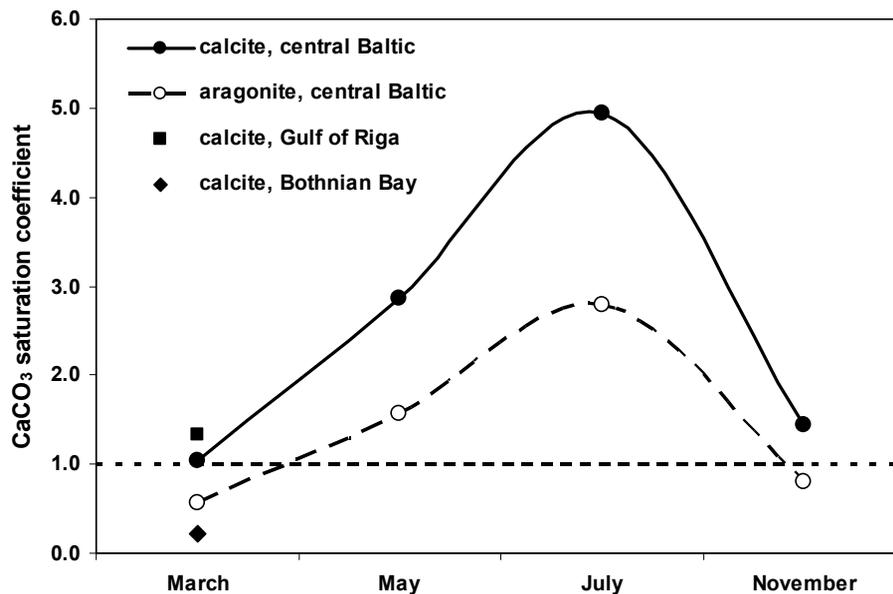


Fig. 2. Seasonality of the calcite and aragonite saturation in the central Baltic Sea (each point represents the average of 26 adjacent measurements), and calcite saturation in both the Gulf of Riga and the Bothnian Bay during March.

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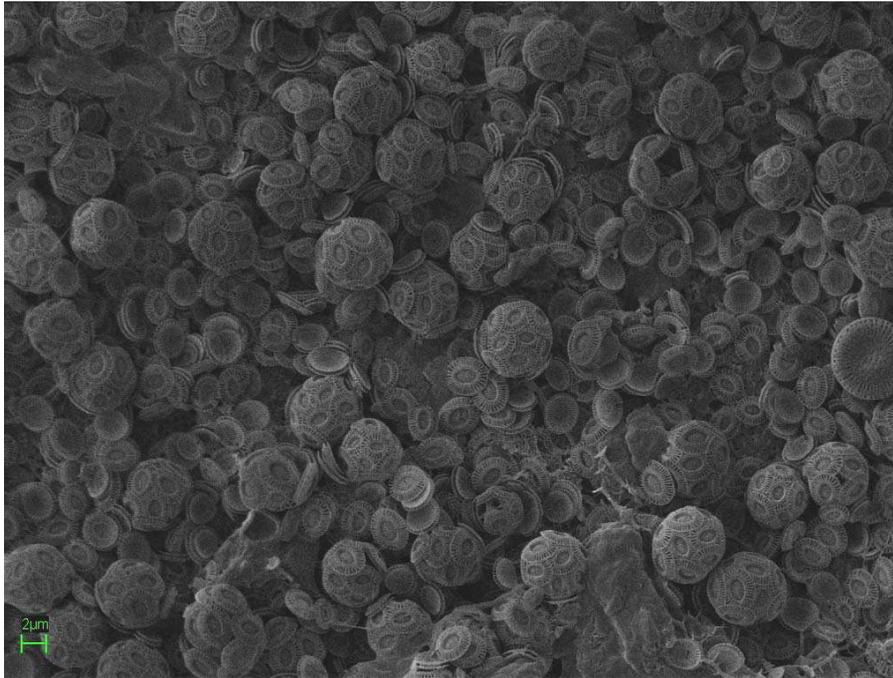


Fig. 3. An SEM image from $\sim(28.5^\circ \text{ E}, 42.8^\circ \text{ N})$ in the Black Sea, June 2006. *E. huxleyi* concentrations of 0.8–2.6 million cells per litre were calculated. Methods were as described earlier for the Baltic Sea, with the exception that up to 3 l were filtered in the Black Sea, and that $0.45 \mu\text{m}$ nitrocellulose filter papers were used.

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