**Diversity and abundance of haptophytes in the East China Sea**

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ABSTRACT: Haptophytes (Prymnesiophytes) are important members of eukaryotic phytoplankton and therefore influence carbon and sulfur cycles in the marine ecosystem. The objective of this study was to investigate the distribution and taxonomic composition of haptophytes in the East China Sea (ECS). Our sampling area covered the ECS continental shelf in the spring and summer of 2009. An abundance of haptophytes was revealed by fluorescent in situ hybridization with an 18S rRNA-specific probe. A haptophyte blooming event reaching $2.2 \times 10^4$ cells ml$^{-1}$ was observed in the Yellow Sea Mixing Water in spring. Phylogenetic analysis of the 18S rRNA gene revealed that the majority of this bloom was composed of *Phaeocystis* sequences that affiliated with *P. jahnii* and *P. cordata*. In contrast, in summer the assemblage composition of haptophytes was dominated by *Chrysochromulina*, with a maximum abundance of $3.1 \times 10^3$ cells ml$^{-1}$ occurring near the edge of the Changjiang Diluted Water. We also noted a relationship between larger size haptophytes and water eutrophication. The contribution of haptophytes to total nanoflagellate abundance at the ECS surface was 32% in spring and 12% in summer.

KEY WORDS: Haptophyte · *Phaeocystis* · 18S rRNA gene

**INTRODUCTION**

Marine phytoplanktonic photosynthesis plays an important role in the global carbon cycle. It is known that eukaryotic phytoplankton contribute more than prokaryotic phytoplankton to biomass and primary productivity in small-size fractions (Worden et al. 2004). Eukaryotic phytoplankton comprise mainly Chlorophyta, Haptophyta, Heterokonta, Cryptophyta and Chlorarachniophyta (Vaulot et al. 2008), of which haptophytes (Prymnesiophytes) sometimes dominate in small-size fractions (cell size <5 µm) based on quantitative data (Cuvelier et al. 2010, Masquelier et al. 2011). Also, an analysis of pigment composition and clone libraries suggests that haptophytes are important constituents of phytoplankton in the ocean (Not et al. 2008, Liu et al. 2009). Furthermore, Jardilier et al. (2010) reported that haptophytes are responsible for a significant percentage of the total carbon dioxide fixation in tropical and subtropical waters. However, the taxonomic composition of haptophytes is less well understood in marine ecosystems.

The haptophytes, which are mostly photosynthetic eukaryotes, are named for their unique haptonema used for attachment or capturing prey. The haptophytes, which include many famous taxa, such as *Phaeocystis*, *Chrysochromulina* and *Prymnesium*, play an important role in marine ecosystems due to their ability to form harmful and troublesome blooms. *Phaeocystis* is a cosmopolitan and well-known blooming genus of haptophytes, and tends to be more abundant in coastal waters at high latitudes (Schoemann et al. 2005). Their impacts on human economic activity include increased fish mortality, reduced shell fish reproduction and odorous foam on beaches (Lancelot & Mathot 1987). The carbon
fixed by *Phaeocystis* blooms can be exported to the deep sea or transferred to higher trophic levels via zooplankton predation (DiTullio et al. 2000, Schoemann et al. 2005). Moreover, *Phaeocystis* produces substantial amounts of dimethylsulfoniopropionate, the precursor of dimethylsulfide, which influences the global climate (Liss et al. 1994, Schoemann et al. 2005).

The hydrographic conditions of the East China Sea (ECS) are highly diverse and change with the seasons. During the spring of 2009, the Shelf Mixing Water (SMW) and the Yellow Sea Mixing Water (YSMW) were the main currents over the ECS shelf. The SMW, influenced by the Kuroshio flowing northward along the shelf edge of the ECS, was characterized as oligotrophic with high temperature (>19°C) and high salinity (>33.7) (Fig. 1A). The YSMW, which occupied the northern ECS, was colder (<19°C) and more eutrophic than the SMW. The water masses of the ECS in summer consisted mainly of the Changjiang Diluted Water (CDW) and the Taiwan Current Warm Water (TCWW). The CDW occupied the north (salinity ≤31) while the TCWW occupied the south (salinity >31) (Gong et al. 1996, Chiang et al. 2002) (Fig. 1A). Besides natural factors, the influence of human activity on the hydrography of the ECS should also be considered. The Changjiang River (also known as the Yangtze River), the longest river in China and the third longest in the world, injects large amounts of terrestrial nutrients and anthropogenic pollutants into the northern ECS (Dagg et al. 2004), to such an extent that eutrophic water often occupies the coastal waters of China, resulting in algal bloom events involving phytoplankton such as diatoms and dinoflagellates (Tang et al. 2006, Zhou et al. 2008, Gong et al. 2011).

Most studies on phytoplankton distribution in the ECS have involved pigment analysis, but few have focused on their abundances. The surveys on abundances of microphytoplankton (diatom) and prokaryotic picophytoplankton (*Synechococcus*) in the ECS indicate that their distribution patterns vary seasonally according to the dynamics of water masses (Chuang et al. 1999, 2002, Chung et al. 2014). Until now, no study has focused on the diversity and distribution of small eukaryotic phytoplankton in that area by using molecular techniques. Haptophytes are known as important contributors to carbon dioxide fixation in subtropical oceans (Jardillier et al. 2010), and also as important constituents in the phytoplankton community in the ECS according to pigment analysis (Furuya et al. 2003). However, little is known about their abundance, cell size and phylogenetic diversity in the ECS. In the present study, fluorescent in situ hybridization associated with tyramide signal amplification (FISH-TSA) targeting 18S rRNA was used to assess the distribution pattern of haptophytes in the ECS surface waters in the spring and summer of 2009. A haptophyte bloom was observed in the YSMW during spring. To discover the main bloom taxa, we first applied a haptophyte-specific probe to eukaryotic 18S rRNA gene clone libraries to acquire full-length 18S rRNA gene sequences. Haptophyte-specific primers were also applied to amplify a shorter sequence of the 18S rRNA gene. To delineate the distribution patterns in bloom (spring) and non-bloom (summer) conditions, correlation analysis was performed between abundance and haptophyte cell size, and various environmental parameters. Profile data along the blooming transect was examined to aid our understanding of the vertical distribution of haptophytes and their requirements for nutrients and irradiance.

**MATERIALS AND METHODS**

**Sample collection and determination of hydrographic features**

Samples of surface water (2 to 3 m depth) were collected from onboard RV ‘Ocean Researcher I’ during the spring (29 April to 15 May) and summer (29 June to 13 July) of 2009. To understand the vertical distribution of haptophytes, water samples were collected from the surface down to near-bottom, or a depth of 100 m, in the northern transect (Stns 19 to 24) during the spring cruise (Fig. 1B). Temperature and salinity data were acquired using conductivity, temperature, depth (CTD) instruments, SBE911 plus, (SeaBird), while water transparency and dissolved oxygen concentration were measured using additional sensors (Alphatrace II Transmissometer and SBE 43, respectively) attached to the CTD frame. Seawater was collected in 20 l Niskin bottles mounted on a Rossette sampler equipped with a CTD profiler. Nutrients (NH₄, NO₂, NO₃, SiO₃ and PO₄) were measured in line with previous studies (Morris & Riley 1963, Pai & Yang 1990), but with some modifications (Gong et al. 1995). To determine chlorophyll a (chl a) concentration, 1 or 2 l of seawater was filtered onto a 25 mm diameter GF/F filter (Whatman). After extraction, chl a was retained on the filter by acetone, and the concentration was determined with a fluorometer (Turner Designs) (Parsons et al. 1984).
**Fluorescent in situ hybridization associated with tyramide signal amplification (FISH-TSA)**

Seawater samples (90 or 180 ml) for the FISH-TSA were preserved with formaldehyde (Sigma) at a final concentration of 3.7% for 1 to 24 h. Cells in the fixative samples were subsequently collected on a 0.8 µm pore size polycarbonate membrane (47 mm diameter, Whatman) and stored at ~80°C until the FISH-TSA procedure. The FISH-TSA method was performed according to Not et al. (2002). In brief, the filters were hybridized with a horseradish peroxidase-labeled PRYM02 probe at a final concentration of 5 ng µl⁻¹ in a hybridization buffer (40% deionized formamide, 0.9 M NaCl, 20 mM Tris–HCl, 0.01% sodium dodecyl sulfate [SDS], 2% blocking reagent [Roche Diagnostic Boehringer]) at 35°C for 3 h. The PRYM02 probe (5’-GGA ATA CGA GTG CCC CTG AC-3’) specifically recognizes the 18S rRNA in haptophyte cells (Simon et al. 2000) and can match 95% (22/466) of haptophytes from *in silico* analysis of both SILVA ARB database and our sequences. Non-hybridized probes were subsequently washed off with a washing buffer (56 μM NaCl, 5 mM EDTA, 0.01% SDS, 20 mM Tris-HCl [pH 7.5]) at 37°C for 10 min twice. Signal amplification was performed in the amplification buffer containing Alexa488-tyramide at room temperature for 30 min in the dark. The membrane was observed with an epi-fluorescent microscope (Nikon Optiphot-2) after counterstaining cellular nuclei with 4’,6-diamidino-2-phenylindole (DAPI) and mounting with a glycerol mixing anti-fading reagent AF3 (Citiñuor Ltd) (Porter & Feig 1980).

**Microscopic counts of haptophytes and total nanoflagellates**

Abundances of haptophytes and total nanoflagellates (≥2 µm, divided into heterotrophic and pigmented nanoflagellates) were estimated using different sections of the same filter. Counting of pigmented nanoflagellates excluded cell sizes of <2 µm due to extremely high values (~10⁵ cells ml⁻¹) in some eutrophic water in the ECS. Under excitation at a wavelength of 470 nm (blue), the FISH signals emit a green color. Haptophyte cell size was represented as equivalent spherical diameter, calculated from the length and width of a cell. The biovolume-carbon conversion for a eukaryotic phytoplankton was evaluated using 237 fg carbon µm⁻³ (Worden et al. 2004).

**Phylogenetic analysis of 18S rRNA gene sequences**

The clone libraries for the 18S rRNA gene were constructed at Stns 19, 23 and 12, representing eutrophic, mesotrophic and oligotrophic water, respectively (Fig. 1B). For DNA extraction, between 1.5 and 9.0 l of seawater flowed through a 20 µm nylon mesh, and was then pre-filtered through a 5 µm pore size filter. Cells <5 µm were collected on a 0.8 µm pore size filter. Genomic DNA was extracted by the phenol–chloroform method (Lin et al. 2012). The universal eukaryotic primers EUKA (5’-AAC CTG GTT GAT CCT GCC AGT A-3’) and EUKB (5’-GAT CCT TCT GCA GGT TCA CCT AC-3’) were used in a PCR reaction to amplify the 18S rRNA gene (Medlin et al. 1988). The PCR conditions were as follows: 95°C for 2 min, followed by 30 cycles of denaturing at 95°C for 45 s, annealing at 55°C for 1 min and extension at 68°C for 2 min, with a final extension step at 68°C for 10 min. The subsequent cloning processes are described in Lin et al. (2012). We obtained about 50 successful eukaryotic sequences from each library for the full length of the 18S rRNA gene.

To comprehensively determine the assemblage composition of the haptophyte bloom observed in the spring of 2009, we used 2 methods to clarify the main bloom taxa. A haptophyte-specific probe was applied to 18S rRNA gene clone libraries to acquire full-length sequences. In addition, haptophyte-specific primers were used to amplify a shorter sequence of the 18S rRNA gene. Stn 16 was selected to study assemblage composition due to the abundance of haptophytes that occurred there (10.3×10³ cells ml⁻¹; see Fig. 1E). About 400 colonies were randomly picked from the 18S rRNA gene libraries of both spring and summer samples. After disrupting cells with SDS, the plasmid DNA was transferred to nylon membranes (NEN Life Science Products) and hybridized with a digoxigenin (DIG) labeled PRYM02 probe. These colonies were recognized using alkaline phosphatase-conjugated anti-DIG antibodies (Roche) and reacted with substrate, CDP-Star, to produce a chemiluminescent signal. The signals were then detected by an imaging system (UVP Biospectrum) (Chung et al. 2003). Colonies with a positive signal were chosen for sequencing. For the Stn 16 samples, the universal eukaryotic primers Euk328f (5’-ACC TGG TTG ATC CTG CCA G-3’) and Euk329r (5’-TGA TCC TTC YGC AGG TTC AC-3’) were used in a PCR reaction to acquire full length 18S rRNA gene sequences (Moon-Van der Staay et al. 2000). The PCR conditions were as fol-
Fig. 1. (A) East China Sea (ECS) currents. The spring and summer currents are indicated with white arrows and black arrows, respectively. (B) ECS sampling stations. Surface abundance of haptophytes sampled in both spring and summer are indicated by filled circles. The open circles indicate stations sampled only in summer, while the triangle (Stn 14) indicates spring sampling only. The abundances of pigmented nanoflagellates and haptophytes were measured from surface water (depth of 2 to 3 m; C−F), except in the north transect of Stns 19 to 24 in spring (dashed line) where a vertical profile was determined. The stations marked with a filled star indicate sequences acquired from the cloning of all eukaryotes, while the station marked with an open star indicate acquisition from haptophytes only. Distribution patterns of (C,D) pigmented nanoflagellates (PNF) and (E,F) haptophytes in spring and summer (10^2 cells ml^{-1}). For the sake of clarity, the contour intervals are not equidistant.
lows: 95°C for 15 min, followed by 32 cycles of denaturing at 95°C for 1 min, annealing at 55°C for 2 min and extension at 72°C for 3 min, with a final extension step at 72°C for 10 min. A second approach used the haptophyte-specific primers Prym-429F (5’-GGC GCG AAT TGG CCC CCA GAA-3’) and Prym-887R (5’-GGA ATA CGA GTG CCC CTG AC-3’) to amplify the 18S rRNA gene on PCR (Simon et al. 2000, Coolen et al. 2004) and its amplicon was ~465 bp. The reverse primer Prym-887R is identical to the FISH probe of PRYM02. The touchdown PCR conditions consisted of 10 cycles of 94°C for 15 s, 60 to 51°C for 30 s, 72°C for 2 min, followed by 25 cycles of 94°C for 15 s, 55°C for 30 to 54 s and 72°C for 2 min, with a final extension step of 72°C for 10 min. Sanger’s sequencing reaction was performed on a PRISM 3100 Genetic Analyzer (Applied Biosystems). Full-length 18S rRNA gene sequences were sequenced with 3 primers, 2 vector (T7 and SP6) and one internal (502f, 5’-GGA GGG CAA GTC TGG T-3’) (Worden 2006), while partial sequences were sequenced with only one vector primer, T7.

All sequences were annotated against known sequences deposited in the GenBank database by the BLASTN algorithm (National Center for Biotechnology Information). A check for chimeric sequences was performed if the sequences had an identity <98% against the known 18S rRNA sequences. Chimeric sequences were detected when annotations conflicted (at the order level) for different regions of an individual sequence (Massana et al. 2006), and were then excluded from further analysis. The sequences for phylogenetic analysis were edited and aligned with the ClustalW and Bioedit software. Phylogenetic trees were inferred from the neighbor-joining (NJ) and maximum likelihood (ML) methods, respectively, using 1000 and 100 bootstraps in PHYLIP 3.69 (Felsenstein 2005). The nucleotide substitution model for ML was selected by JModelTest with the Bayesian information criteria (Posada 2008). The haptophyte sequences reported in this paper have been deposited in the GenBank database under accession numbers KF620969 to KF621011 for full-length sequences and KJ608556 to KJ608631 for partial-length sequences.

### Statistical analysis

The relationship between biological factors and their ambient environmental parameters were analyzed using Pearson’s correlation (SPSS). Environmental parameters included temperature, salinity, nutrients (NH4, NO2, NO3, SiO3 and PO4), transparency, dissolved oxygen and chl a concentration.

### RESULTS

#### Surface distribution of pigmented nanoflagellates and haptophytes

The cell concentrations of pigmented nanoflagellates (PNF) and haptophytes at each station in the ECS are shown in Fig. 1C–F. Surface abundances of pigmented nanoflagellates varied from 5.6×10² to 4.3×10⁴ cells ml⁻¹ in spring, and from 2.1×10² to 2.8×10⁴ cells ml⁻¹ in summer (Fig. 1C,D). For haptophytes, the range was from 1.5×10² to 2.2×10⁴ cells ml⁻¹ in spring and from 4.8×10⁴ to 3.1×10⁵ cells ml⁻¹ in summer (Fig. 1, Table 1). The abundances showed similar distribution patterns in spring, but not in summer (Fig. 1). In spring, peak haptophyte abundance appeared in the northern transect of the ECS, which was occupied by the YSMW (Figs. 1E & 2). A secondary peak of haptophyte abundance (1.0×10⁴ cells ml⁻¹) appeared at Stn 16 in the middle shelf (Fig. 1E).

The high levels of haptophyte abundance (>10⁴ cells ml⁻¹) indicated haptophyte bloom conditions in the northern region of the ECS in spring. In summer, peak haptophyte abundance occurred in the north-

<table>
<thead>
<tr>
<th>Variable</th>
<th>Spring</th>
<th>Summer</th>
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<tr>
<td>Temperature (°C)</td>
<td>19.4 ± 2.3 (15.7–25.3)</td>
<td>26.6 ± 1.5*** (23.3–28.8)</td>
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<tr>
<td>Salinity</td>
<td>32.6 ± 1.9 (27.9–34.5)</td>
<td>32.3 ± 2.0 (23.8–34.1)</td>
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<td>NH4 (µM)</td>
<td>1.0 ± 0.7 (0.1–3.0)</td>
<td>0.8 ± 0.6 (0.25–2.5)</td>
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<tr>
<td>NO2 (µM)</td>
<td>0.3 ± 0.3 (0–1.4)</td>
<td>0.2 ± 0.3 (0–1.0)</td>
</tr>
<tr>
<td>NO3 (µM)</td>
<td>3.2 ± 5.9 (0–25.3)</td>
<td>2.2 ± 5.7 (0–24.3)</td>
</tr>
<tr>
<td>PO4 (µM)</td>
<td>0.05 ± 0.08 (0–0.36)</td>
<td>0.14 ± 0.19** (0–0.83)</td>
</tr>
<tr>
<td>SiO3 (µM)</td>
<td>6.4 ± 6.3 (0.9–28.7)</td>
<td>6.1 ± 6.2 (1.5–24.5)</td>
</tr>
<tr>
<td>Chlorophyll a (mg m⁻³)</td>
<td>1.96 ± 2.47* (0.24–10.96)</td>
<td>1.14 ± 1.40 (0.14–5.1)</td>
</tr>
<tr>
<td>Haptophyte (10⁴ cells ml⁻¹)</td>
<td>30.9 ± 43.7** (1.6–223.9)</td>
<td>6.1 ± 6.7 (0.5–27.6)</td>
</tr>
</tbody>
</table>

Table 1. Comparison of environmental variables and haptophyte abundance in surface water at the stations for both spring and summer samples (filled circles in Fig. 1B; n = 31). The table shows average values, standard deviations and ranges of these variables. A paired t-test was performed to judge the difference between the means of parameters between spring and summer and the results were considered significantly different at *p < 0.05, **p < 0.01 or ***p < 0.001. The temperature and phosphate concentration were higher in summer while concentrations of chlorophyll a and haptophytes were higher in spring.
ern ECS at the edge of the CDW (Figs. 1F & 2). Generally, there was a low abundance of haptophytes (<1×10^3 cells ml⁻¹) in the southern ECS and offshore area in spring, and also outside the northern plume area in summer (Fig. 1E,F). The average abundance of haptophytes in spring and summer was about 3×10³ cells ml⁻¹ and 6×10² cells ml⁻¹, respectively (Table 1). The abundance of haptophytes was higher in spring than in summer (paired t-test, p < 0.001; Table 1). Overall, haptophytes contributed 52% and 49% to pigmented nanoflagellates, and 32% and 12% to total nanoflagellates abundances (cells ml⁻¹) in spring and summer, respectively.

Vertical distribution of haptophytes along the cross-shelf transects during the bloom

A haptophyte bloom was observed in the YSMW during the spring period (Fig. 1E). A vertical profile along the transect of Stns 19 to 24 showed that fresh water from the Changjiang was visible at Stn 19, whereas the more saline water of the YSMW occupied Stns 20 to 23 (Fig. 3). The haptophyte abundance peaked at the boundary of the coastal mixing water and oceanic water (Fig. 3). At Stns 19 to 22,
haptophyte abundance peaked at the surface, and then decreased with depth (Fig. 3). Chl a concentrations peaked at the surface in the more coastal water of Stns 19 and 20, where the distribution was similar to that of haptophytes, but peaked in the subsurface (~10 to 20 m) in the more oceanic water of Stns 21 to 23 (Fig. 3).

Haptophyte cell size

Haptophyte cell size in the ECS ranged from 2 to 9 µm, with averages of 3.5 ± 0.5 [SD] µm in spring and 3.1 ± 0.4 µm in summer. The average haptophyte size at each station varied from 2.7 to 5.0 µm in spring and 2.2 to 3.9 µm in summer. In spring, haptophytes with an average size of >4 µm appeared on the coast (Stns 18, 19 and 30) and middle shelf (Stns 15 and 16). However, in the summer, no station had haptophytes with an average size of >4 µm, and those between 3.5 and 4.0 µm appeared mostly near the areas of the CDW (Stns 19, 19A, 21−23 and 40−43) and the coast (Stn 4). Correlation analysis revealed that haptophyte cell size was positively associated with dissolved oxygen, NO3, SiO3 and chl a concentrations, and negatively with temperature, salinity and transparency (details of correlation coefficients and p-values are shown in Table 2).

18S rRNA gene phylogeny of haptophytes

Stns 12, 19 and 23 showed unique hydrographic conditions (Lin et al. 2012), and were selected to construct 18S rRNA gene clone libraries. A higher percentage of haptophyte sequences, 9% (5/53), was retrieved from Stn 23 in spring, reflecting the high relative abundance of haptophytes there. However, no haptophyte sequences were retrieved from the offshore water of Stn 12 in spring or from the coastal water of Stn 19 in summer.

By using a haptophyte-specific probe to screen about 400 clones randomly picked from the eukaryotic 18S rRNA gene libraries of Stn 16, 11 chimeric sequences were excluded, and a total of 24 accurate full-length haptophyte 18S rRNA sequences were retrieved. The haptophyte sequences from Stn 16 during the spring bloom contained 13 sequences of Phaeocystis, 6 sequences of Chrysochromulina, 4 sequences of Haptolina and 1 sequence of Prymnesiaceae. The Phaeocystis sequences were affiliated with P. cordata and P. jahnii, with identities from 98.5 to 99.4% and from 97.8 to 98.1%, respectively. Phylogenetic analysis showed that the sequences related to P. cordata and P. jahnii in the ECS were somewhat distinct from the cultured representatives, particularly those affiliated to P. jahnii (Fig. 4).

The partial sequences of the 18S rRNA gene were amplified with a specific primer pair. In total, 36 and 40 accurate sequences were obtained from spring and summer samples, respectively. All sequences belonged to haptophytes. In spring, Phaeocystis accounted for 42% of the total, with Prymnesium being the second largest contributor, accounting for 25%. In summer, Chrysochromulina was the dominant component (33%) (Table 3). Phylogenetic analysis showed that in spring some Phaeocystis sequences in the ECS affiliated to a clone retrieved from symbiotic Phaeocystis in Acantharia of the West Pacific Ocean (Fig. 5). Some Phaeocystis sequences from the ECS were affiliated to P. cordata, which was consistent with the results from full-length analysis. In this study, these Phaeocystis sequences were defined as P. jahnii-like and P. cordata-like. Most Phaeocystis and Haptolina sequences were acquired from spring samples, whereas most Chrysochromulina sequences were acquired in the summer (Fig. 5, Table 3).

DISCUSSION

Haptophytes can be ascribed to either pico- or nanosize fractions (Cuvelier et al. 2010, Masquelier et al. 2011). Preservation with formaldehyde can shrink cells, and therefore the size of haptophytes in our study may be underestimated (Choi & Stoecker 1989). In this study, we regarded haptophytes as nanoplanckton due to their observed size range of 2 to

### Table 2. Pearson’s correlations (r) were performed between haptophyte cell size and environmental variables in surface water in spring (n = 32) and summer (n = 47), respectively. Values marked with asterisks were considered significant, **p < 0.01 or ***p < 0.001

<table>
<thead>
<tr>
<th>Variables</th>
<th>Cell size</th>
<th>Spring</th>
<th>Summer</th>
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<tbody>
<tr>
<td>Temperature (°C)</td>
<td>−0.457**</td>
<td>−0.463**</td>
<td></td>
</tr>
<tr>
<td>Salinity</td>
<td>−0.690***</td>
<td>−0.463***</td>
<td></td>
</tr>
<tr>
<td>Transparency (%)</td>
<td>−0.579**</td>
<td>−0.455**</td>
<td></td>
</tr>
<tr>
<td>Dissolved oxygen (µM)</td>
<td>0.669***</td>
<td>0.575***</td>
<td></td>
</tr>
<tr>
<td>NH4 (µM)</td>
<td>0.248</td>
<td>0.086</td>
<td></td>
</tr>
<tr>
<td>NO2 (µM)</td>
<td>0.324</td>
<td>0.272</td>
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</tr>
<tr>
<td>NO3 (µM)</td>
<td>0.584***</td>
<td>0.156</td>
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<tr>
<td>PO4 (µM)</td>
<td>0.298</td>
<td>0.283</td>
<td></td>
</tr>
<tr>
<td>SiO3 (µM)</td>
<td>0.609***</td>
<td>0.458**</td>
<td></td>
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<tr>
<td>Chlorophyll a (mg m−3)</td>
<td>0.540**</td>
<td>0.583***</td>
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Previous studies have reported that haptophytes account for nearly half of the picophytoplankton carbon biomass at high latitudes in the Atlantic and Pacific oceans (Cuvelier et al. 2010), and contribute 38% of the eukaryotic abundance in the Atlantic water mass of the English Channel (Masquelier et al. 2011). In our study, haptophytes were detected at every station in the ECS (58 to 22,392 cells ml⁻¹), and overall contributed 32% and 12% to the abundances of total nanoflagellates (cells ml⁻¹) in spring and summer, respectively. These results indicate that haptophytes are ubiquitous and an important component of eukaryotic phytoplankton in various marine ecosystems.

Even though the hydrographic conditions in the ECS are highly dynamic, the major water masses that occupy the ECS in the spring (SMW and YSMW) and summer (TCWW and CDW) were simply characterized using the salinity and temperature criteria defined by Gong et al. (1996) and Chiang et al. (2002) (Fig. 2). In spring, high numbers of haptophyte cells were observed in the YSMW (Fig. 2). Moreover, haptophyte abundance was negatively correlated with temperature in spring (r = −0.53, p < 0.01), but did not correlate with nutrients. This suggests that the high abundance of haptophytes is driven by cold water in spring in the ECS. During summer, the haptophyte abundance peaked near

Fig. 4. Haptophyte phylogeny constructed using neighbor-joining with near-full length 18S rRNA gene sequences. A total of 1399 bp alignment positions were analyzed using maximum likelihood (ML) and neighbor-joining (NJ) methods with 1000 and 100 bootstraps, respectively. Only nodes with >70% support are shown; filled circles indicate significant support by both ML and NJ; gray circles and open circles indicate significant support by only ML or only NJ, respectively. The TrN+I+G substitution model (I = 0.316 and α = 0.487) was used and Micromonas pusilla (AY954997), Rhodomonas salina (EU926158), Thalasiosira pseudonana (DQ093367) serve as an outgroup. The scale bar represents 0.02 nucleotide changes per position. The names of the East China Sea clones contain the cruise numbers (897, spring; 905, summer) and station numbers (Stn 12, 16, 19 or 23). The blue and red sequences represent spring and summer clones, respectively. Cultured or identified sequences are in gray and the environmental sequences are in black.
Phaeocystis is the dominant genus in spring in the ECS according to 18S rRNA gene data (Figs. 4 & 5, Table 3). The large haptophytes (>4 µm) found in coastal water (Stns 18, 19 and 30) and the middle shelf (Stns 15 and 16) may therefore consist of Phaeocystis. Overall, haptophyte cell sizes were smaller in summer than in spring (t-test, p < 0.001) and the average sizes in summer were all <4 µm. The results of correlation analysis suggest that the level of eutrophication influences haptophyte cell size in both spring and summer (Table 2). This lends support to the idea that small phytoplankton with a high surface-volume ratio have an advantage over other phytoplankton in oligotrophic conditions in the competition for nutrients (Raven 1998). In addition, a positive relationship between cell size and haptophyte abundance implies that the types of haptophyte tend to be those with a larger cell size in bloom conditions (r = 0.36, p < 0.01).

In addition to the large number of Phaeocystis sequences discovered in spring, Chrysochromulina was also an important component in the ECS (Figs. 4 & 5, Table 3). This is consistent with previous research based on studies of 28S rRNA and 18S rRNA genes that showed that Chrysochromulina contributes a large proportion of haptophytes in the open ocean (Liu et al. 2009, Cuvelier et al. 2010). With respect to the phylogeny of the 18S rRNA gene, the high diversity detected in Chrysochromulina implies that some of the genetic clades do not have a cultured representative (Fig. 4). Moreover, some novel haptophyte sequences located at the base of the phylogenetic tree, including 2 sequences reported here.
Table 3. Clone numbers for different taxonomic groups of haptophytes at Stn 16 found by using specific primers to amplify the 18S rRNA gene. A total of 36 and 40 sequences were obtained in the spring and summer, respectively. The number in parentheses represents the percentage of total haptophyte sequences.

<table>
<thead>
<tr>
<th>Seasons</th>
<th>Clone numbers</th>
<th>Haptolina</th>
<th>Prymnesium</th>
<th>Chrysochromulina</th>
<th>Phaeocystis</th>
<th>Others</th>
</tr>
</thead>
<tbody>
<tr>
<td>Spring</td>
<td>36</td>
<td>7 (19%)</td>
<td>9 (25%)</td>
<td>5 (14%)</td>
<td>15 (42%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Summer</td>
<td>40</td>
<td>4 (10%)</td>
<td>4 (10%)</td>
<td>13 (33%)</td>
<td>8 (20%)</td>
<td>11 (28%)</td>
</tr>
</tbody>
</table>

Fig. 5. Haptophyte phylogeny constructed using neighbor-joining with partial length 18S rRNA gene sequences. A total of 449 bp alignment positions were analyzed using maximum likelihood (ML) and neighbor-joining (NJ) methods with 1000 and 100 bootstraps, respectively. Only nodes with >70% support are shown; filled circles indicate significant support by both ML and NJ; gray circles and open circles indicate significant support by only ML and only NJ, respectively. The TrNef+G substitution model used ($\alpha = 0.473$) and outgroup was as for Fig. 4. The scale bar represents 0.03 nucleotide changes per position. The filled squares represent the full-length sequences also shown in Fig. 4.
and 2 environmental sequences (Biosope_T58.080 and Biosope_T65.100) from the South East Pacific Ocean (Fig. 4), confirm the extensive phylogenetic diversity and novelty of haptophytes in the world’s oceans (Shi et al. 2009).

The haptophyte bloom was observed in the YSMW during spring, and approximately half of the haptophyte sequences belonged to Phaeocystis at Stn 16 (Table 3). Phaeocystis sequences were also retrieved from the coastal water and continental shelf water (Stns 16, 19 and 23; Figs. 4 & 5), confirming it as the dominant genus in the ECS during spring. Large numbers of Phaeocystis sequences were retrieved from the spring samples, but only a few were obtained from the summer samples (Table 3). Previous studies have indicated that Phaeocystis is a spring-bloom genus, particularly at high latitudes (DiTullio et al. 2000, Monchy et al. 2012, Vogt et al. 2012). P. pouchetii and P. antarctica are well-known bloom species occurring in polar areas, and P.oglobosa is prevalent from tropical to temperate areas (Schoemann et al. 2005, Monchy et al. 2012, Vogt et al. 2012, Rousseau et al. 2013). In our study, the Phaeocystis sequences from the ECS were related to P. jahnii and P. cordata, both originally found in Mediterranean water. Scanning electron microscopy inspection of the FISH filter at Stn 16 revealed no colonies of Phaeocystis, but pentagonal stars of Phaeocystis and heart-shaped P. cordata-like cells were found (data not shown). Cells on a filter preserved with a high concentration of formaldehyde may dissolve colonies, making their detection difficult. Until now, all of the sequences affiliated to P. jahnii and P. cordata have been retrieved from latitudes between 10 and 45°. A recent study on the phylogeny of the symbiotic relationship between Phaeocystis and Acantharia demonstrated that genetic distances of symbiont Phaeocystis were significantly correlated to its biogeography, but had no relationship with the host taxonomy (Decelle et al. 2012). Overall, our results are consistent with previous studies showing that species of Phaeocystis occurring in polar and non-polar areas are distinct (Schoemann et al. 2005, Decelle et al. 2012). Few studies have reported blooms of P. jahnii and P. cordata, which may be due to their small size, or may indicate that they cannot form, or seldom form, colonies in the natural environment (Zingone et al. 1999, Medlin & Zingone 2007). Molecular approaches appear to be useful tools for studying the distributions of P. jahnii and P. cordata.
Temperature may be an important factor in controlling the distribution of *P. jahnii* and *P. cordata* during spring in the ECS. Laboratory studies indicate that the optimal temperature for the maximum growth rate of *Phaeocystis* is around 16°C (Schoemann et al. 2005). A large number of 16S rRNA gene sequences related to *P. jahnii* and *P. cordata* were simultaneously retrieved from spring samples in the Gulf of Naples, implying either an unknown relationship or that they are controlled by the same resources. The temperature in the Gulf of Naples at that time was around 15°C, similar to that of our spring sampling (McDonald et al. 2007). Our study indicated that peak haptophyte abundance occurred at 15.7°C in spring, and 18S rRNA gene data showed that *Phaeocystis* was the dominant genus (Figs. 2, 4 & 5). Zingone et al. (1999) stated that the presence of *Phaeocystis* in the Gulf of Naples cannot be attributed to temperature because they also appear in autumn when the temperature exceeds 23°C. Moreover, the environmental clones related to *P. jahnii* retrieved from the Indian and Atlantic oceans (IND58.21 and AMT15.15_10m_412) came from warm waters (>25°C) (Not et al. 2008, Kirkham et al. 2011). However, no study has ever reported that *Phaeocystis* thrives in high-temperature water.

Previous studies have demonstrated that a *Phaeocystis* bloom is induced by nitrate excess (Breton et al. 2006), whereas a diatom bloom is controlled by silicate (Breton et al. 2006). However, no study has ever reported that *Phaeocystis* blooms are induced by nitrate excess (Breton et al. 2006), whereas a diatom bloom is controlled by silicate. This study reported a haptophyte bloom event during the spring in a subtropical system of the northwestern Pacific Ocean. Overall, the biomass of haptophytes in the ECS ranged from 1 to 194 µg l⁻¹ in spring and 0.09 to 21 µg l⁻¹ in summer. A high level of haptophyte abundance appeared in the YSMW in spring, of which 18S rRNA genes were affiliated with *Phaeocystis jahnii* and *P. cordata*. The peak of haptophyte abundance occurred at the boundary between coastal mixing water and oceanic water, and also at the station with the lowest temperature in the spring. We do not know whether the high level of haptophytes was due to their being brought in by Yellow Sea currents, or whether they simply thrived in the favorable environment in the ECS during spring. However, future studies in the ECS should look at the fate of carbon and sulfur produced by haptophytes in spring as they make up a large part of total nanoplanckton.

**CONCLUSIONS**

This study reported a haptophyte bloom event during the spring in a subtropical system of the northwestern Pacific Ocean. Overall, the biomass of haptophytes in the ECS ranged from 1 to 194 µg l⁻¹ in spring and 0.09 to 21 µg l⁻¹ in summer. A high level of haptophyte abundance appeared in the YSMW in spring, of which 18S rRNA genes were affiliated with *Phaeocystis jahnii* and *P. cordata*. The peak of haptophyte abundance occurred at the boundary between coastal mixing water and oceanic water, and also at the station with the lowest temperature in the spring. We do not know whether the high level of haptophytes was due to their being brought in by Yellow Sea currents, or whether they simply thrived in the favorable environment in the ECS during spring. However, future studies in the ECS should look at the fate of carbon and sulfur produced by haptophytes in spring as they make up a large part of total nanoplanckton.

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