Role of *Calanus sinicus* (Copepoda, Calanoida) on dimethylsulfide production in Jiaozhou Bay

Juan Yu¹,², Jiyuan Tian³, Zhengyu Zhang¹, Guipeng Yang¹,²*, Hongju Chen⁴

¹Key Laboratory of Marine Chemistry Theory and Technology, Ministry of Education; College of Chemistry and Chemical Engineering, Ocean University of China, Qingdao, 266100, People’s Republic of China
²Laboratory for Marine Ecology and Environmental Science, Qingdao National Laboratory for Marine Science and Technology, Qingdao 266071, People’s Republic of China
³College of Food Science and Engineering, Qingdao Agricultural University, Qingdao, 266009, People’s Republic of China
⁴College of Environmental Science and Engineering, Ocean University of China, Qingdao, 266100, People’s Republic of China

Correspondence to: Guipeng Yang (gpyang@ouc.edu.cn)

Abstract. The role of copepod *Calanus sinicus* on the production of dimethylsulfide (DMS)/dimethylsulfiniopropionate (DMSP) in Jiaozhou Bay was evaluated in field measurements and laboratory experiments. Samples at 10 sites in the bay were collected monthly from June 2010 to May 2011 (except for March 2011), and zooplankton species composition was analyzed. Effects of *C. sinicus* grazing on DMS/DMSP production at different conditions (i.e., algal diets, food concentrations, and salinities) were assessed in the laboratory. Data from the field experiment showed that *C. sinicus* was the dominant copepod in Jiaozhou Bay (up to 123 individuals m⁻³ in May 2011) and preferred to graze on diatom. DMS and DMSP concentrations not only depend on phytoplankton abundance, but also phytoplankton species and other factors. In the laboratory experiment, compared with *Gymnodinium* sp. or *Emiliania huxleyi*, *C. sinicus* feeding on *Isochrysis galbana* and *Chaetoceros curvisetus* exhibited increased DMS concentration, whereas high salinity inhibited DMS production. This study indicated that DMSP was transferred from phytoplankton to copepod body, fecal pellet, and seawater through copepod grazing. Our results provided important information to understand the biogeochemical cycle of DMSP in Jiaozhou Bay.

Key words: Copepod; Diet; Dimethylsulfide; Dimethylsulfiniopropionate; Grazing; Salinity

1 Introduction

Dimethylsulfide (DMS) is the most abundant biogenic sulfur gas that may influence planetary climate by forming cloud condensation nuclei that alters global radiation balance (Charlson et al., 1987). The biogeochemical cycling of dimethylsulfiniopropionate (DMSP), which is the main precursor of DMS. The marine environment is the major source of DMSP, and DMSP is synthesized by many marine phytoplankton species as an osmolyte (Kirst, 1996). Therefore, the dynamics of DMSP in the ocean will have an important influence on global DMS production. The conversion of DMSP to DMS is regulated by complex trophic processes in the water column, e.g., algal senescence (Nguyen et al., 1988), phytoplanktonic enzyme catalysis (Niki et al., 2000), bacterial activity (Kiene and Linn, 2000), and zooplankton grazing...
(Dacey and Wakeham, 1986; Wolfe et al., 2000; Yu et al., 2015). Currently, most studies have focused on field research concerning the spatial and temporal distributions and fluxes of DMS/DMSP (Turner et al., 1996; Wong et al., 2005). Many field studies indicated a weak correlation between DMS and parameters directly related to primary producers (Kettle et al., 1999). Numerous biotic factors (i.e., phytoplankton, zooplankton, bacteria, and virus) and abiotic factors (i.e., temperature, salinity, light, and nutrient), e.g., factors other than phytoplankton, might play an important role in DMS and DMSP dynamics. Zooplankton are known to consume for a large fraction (10% to 25%, or higher) of the daily oceanic primary production (Lancelot and Billen, 1985). Copepods, a trophodynamic link between primary and tertiary productions, play a key role in the cycling of materials and energy in marine ecosystems (Kiorboe, 1997). Dacey and Wakeham (1986) were the first to show that copepod grazing stimulated DMS production in a laboratory experiment. Other studies have also documented that zooplankton grazing induced DMS production (Belviso et al., 1990; Christaki et al., 1996; Daly and DiTullio, 1996; Leck et al., 1990) because of sloppy feeding (Dacey and Wakeham, 1986; Tang et al., 2000) or enhanced DMSP lyase activities (Wolfe and Steinke, 1996). The DMSP ingested by zooplankton was accumulated in the body and transferred to the upper food chain, and a portion of the DMSP ingested was transferred to fecal pellets, which were subsequently uncoupled as DMS and DMSP productions.

The calanoid copepod *Calanus sinicus* is distributed in the East China Sea, the Yellow Sea, and the coastal waters of Japan (Brodsky, 1965) and is one of the dominant zooplankton in the East China Sea, the Yellow Sea, and the Jiaozhou Bay. In this study, we performed field measurements and laboratory experiments to investigate the effects of copepod on DMS and DMSP productions in Jiaozhou Bay. The field experiments revealed the relationship between DMS/DMSP and biotic or abiotic parameter, and the laboratory experiments indicated the changes of DMS, dissolved and particulate DMSP (DMSP\(_d\) and DMSP\(_p\)), DMSP in copepod bodies (DMSP\(_c\)), and DMSP in fecal pellets (DMSP\(_f\)) under different conditions of salinities, food concentrations, and diet species. The data obtained in this study will further recognize the role of copepods in the DMSP biogeochemical cycle.

2 Materials and methods

2.1 In situ experiment

2.1.1 Sampling site

Fieldwork was performed at 10 stations of the Jiaozhou Bay from June 2010 to May 2011 once a month, except of March 2011 due to strong wind (Fig. 1). The Jiaozhou Bay (36°7′24.44″ N, 120°14′44.3″ E), a semi-enclosed bay in Qingdao City (China), separates Huangdao District from Qingdao City and borders on Jiaozhou City. The bay is 32 km long and 27 km wide, with a surface area of 362 km².
2.1.2 Abundance and taxonomic composition of zooplankton

Zooplankton samples were collected by vertical tows from the bottom to the surface (the depth varied from 3 m to 28 m according to different stations) using a conical–cylindrical plankton net with a 50 cm mouth opening and a 160 µm mesh size. A flow meter was used to estimate the amount of filtered water. We retrieved the net at 0.3 m s\(^{-1}\) to 0.5 m s\(^{-1}\) at each station. Samples were then rinsed into cod-end buckets, concentrated, and preserved in 5% formalin for qualitative and quantitative analyses of in situ zooplankton. Taxonomy was determined by optical microscopy. In situ abundance was calculated using tow volume estimates, which were determined by net dimensions and flow meter values.

2.1.3 Dilution experiments

Dilution experiments were set up according to the methods of Landry and Hassett (1982) and Wolfe et al. (2000), which are widely used to estimate microzooplankton grazing and phytoplankton growth rates. The technique assumes that increasingly diluted treatments reduce grazer-prey encounter and therefore grazing rates \(g\), without changing specific growth rates \(\mu\) of prey. Net production of a prey biomarker \(B\) is thus given by \(B(t) = B(0)e^{(\mu - d) t}\), where \(d\) is the fraction of unfiltered water; regressing \(1/t \ln(B(t)/B(0))\) vs. \(d\) yields \(\mu\) as the Y-intercept and \(g\) as the negative of the slope (Wolfe et al., 2000).

Depending upon the time allowed for water collection, we conducted dilution experiments aboard at three stations (C3: 120.26°E, 36.13°N; D4: 120.30°E, 36.09°N; E3: 120.30°E, 36.03°N) on each cruise from June 2010 to May 2011 in Jiaozhou Bay (Fig. 1). A dilution series was prepared, consisting of 100, 80, 60, 40, and 20% unfiltered water in 1 L polycarbonate bottles (washed with 10% HCl and distilled water, and rinsed with seawater prior to use). The water was collected using 12 L Niskin bottles on a CTD rosette. Water for dilutions was filtered using gravity through Gelman Suporcap 0.2 µm capsule filters into an acid-washed carboy. These bottles were incubated for 24 h under simulated in situ conditions in a water-bath deck incubator with neutral density screening. Nutrients were added in the bottles to ensure constant phytoplankton growth in the dilution series when the final concentrations of nitrate and phosphate were lower than 0.5 µM and 0.03 µM. We did not pre-screen water to remove mesozooplankton grazers, because copepod numbers were usually low (< 1 L\(^{-1}\)) compared with microzooplankton and our preliminary trials showed that mesograzers had a negligible grazing impact at natural densities compared to microzooplankton. Specific growth and grazing rates of Chl \(a\) were calculated by regressing the time-normalized, log-transformed ratio of final and initial concentrations vs. fraction unfiltered water (Landry and Hassett, 1982).

2.1.4 Abiotic parameters and analyses of chlorophyll \(a\) (Chl \(a\)) and bacteria

The shipboard measurements of temperature and salinity of surface water were obtained with the Sea-Bird 911. The surface water samples were collected by a Niskin sampler and then they were filtered using Whatman GF/F filters (nominal pore size of 0.7 µm) to determine Chl \(a\). Chl \(a\) concentrations were measured using a Hitachi F4500 fluorometer according to the
methods of Parsons et al. (1984). Based on the procedures of Porter and Feig (1980), bacteria were counted by epifluorescence microscopy (Hitachi F4500; total magnification of ×1,000).

2.2 Incubation experiment

2.2.1 Phytoplankton and zooplankton cultures

Phytoplankton species (*Isochrysis galbana*, *Chaetoceros curvisetus*, *Emiliania huxleyi*, and *Gymnodinium* sp.) were provided by the Marine Microalgae Research Center, Ocean University of China in different sizes and DMSP contents. They are widespread species in Jiaozhou Bay and Chinese coastal waters (Li et al., 2005; Zhong et al., 2001). These algae are described in Table 1, and the sizes of algal cells are obtained by measuring at least 200 cells using a calibrated ocular micrometer with a light microscope at a magnification of × 400. Cell volume was estimated by approximating the shape of each species to an elliptic sphere, which subsequently determined the biovolume according to the method of Verity et al. (1992). In this study, we determined the cellular carbon contents in phytoplankton according to the method of Strathmann (1967).

All algae were cultured with f/2 medium (Guillard, 1975) at 60 μmol m⁻² s⁻¹ under a dark/light cycle of 12 h:12 h at a temperature of 15 ± 1 °C. According to the methods of Wolfe and Steinke (1996), the algal culture was detected by epifluorescence microscopy following staining with acridine orange and by plating on 1% peptone agar plates to check for bacterial growth. No bacterial contamination was found in any of the experimental cultures. Algae were fed to copepods during their exponential growth phase. Copepods *C. sinicus* were collected from Jiaozhou Bay, Qingdao, China (120°8′ E, 36°8′ N) in April and May 2011 using a 0.5 m standard ring net equipped with a 160 μm mesh and a solid cod end. The copepods were grown at 15 ± 1 °C in 30 PSU sterilized seawater. In the incubation experiment, adult copepods were utilized as experimental animals.

2.2.2 Ingestion configuration

Four species of algae (*C. curvisetus, I. galbana, Gymnodinium* sp., and *E. huxleyi*) were utilized to assess dietary effects on changes in ingestion rate (IR), clearance rate (CR), and DMS/DMSP production in a salinity of 30 PSU. *I. galbana* was used as a single food source, which was a DMSP-poor and favorite food for *C. sinicus*, and four concentrations of *I. galbana* (10, 15, 20, and 25 × 10⁴ cells mL⁻¹) were set as concentration contrasts to obtain ingestion and DMS/DMSP information from food concentration. Laboratory studies have confirmed that salinity can significantly influence the ingestion rates of copepod and the S-compounds production of phytoplankton (Tang et al., 1999; Yu et al., 2015). Therefore, four levels of salinities (20, 25, 30, and 35 PSU) were set up to investigate the effects of salinity on copepod grazing and the variations of DMSP and DMS concentrations. Single-factor experiments were performed, and all parameters were run individually.
2.2.3 IR and CR

In order to ensure that the copepods were hungry to graze the diets, copepods were starved for 24 h before they were fed with phytoplankton. Copepods were rinsed with sterilized seawater before the beginning of the grazing experiment. Batches of 10 individual copepods in each of 3 replicates were sorted out and transferred to 250 mL polycarbonate bottles. Three bottles containing no grazers were used as controls. All bottles were topped off with suspensions of algae, sealed with parafilm, and fastened onto a spinning plankton wheel (2 rpm) at 15 ± 1 °C in darkness for 24 h. After a 24-h incubation, copepods and fecal pellets were harvested according to the methods of Tang (2001). Water samples were used to analyze cell density, IR, CR, DMS, DMSP\textsubscript{p}, and DMSP\textsubscript{d}. The IRs and CRs were calculated according to the equations of Frost (1972). Because no significant differences were found in algae concentrations between the initial and final control bottles, the growth constant (\(k\)) for algal growth was eliminated from the equations, thus yielding:

\[
CR = \frac{V \times \ln(C_i / C_{2}^\ast)}{Nt}
\]

\[
IR = CR \times C_i
\]

Where \(CR\) is the clearance rate (mL ind\(^{-1}\) h\(^{-1}\)), \(IR\) is the ingestion rate (cells ind\(^{-1}\) h\(^{-1}\)), \(C_i\) is the initial algal concentration in control bottles (cells mL\(^{-1}\)), \(C_{2}^\ast\) is the final algal concentration in the experimental bottles (cells mL\(^{-1}\)), \(t\) is the duration of the experiment (h), \(V\) and \(N\) are the volume (mL) and number of copepods in the experimental bottles (ind), respectively.

Duplicates of 10 mL aliquots of each algal suspension were placed in 40 mL serum bottles that contained 2 mL of 10 mol L\(^{-1}\) KOH solution for DMSP detection. Ten copepods from each bottle were placed individually in serum bottles for DMSP\textsubscript{z} (DMSP in the copepod body) measurement before gut clearance. Fecal pellets were separated from detritus with a mouth micropipette, rinsed with filtered seawater, and concentrated onto a 47 mm Whatman GF/F filter by gravity filtration to determine DMSP\textsubscript{f} (DMSP in the fecal pellets).

We ran preliminary experiment to check the effects of the bacteria on DMS concentration. According to the methods of Agostini et al. (2016), the treatment with antibiotics (0.025 g L\(^{-1}\) penicillin G potassium + 0.08 g L\(^{-1}\) streptomycin sulphate + 0.04 g L\(^{-1}\) neomycin sulphate) were used to inhibit the bacteria in the algal culture. When \textit{C. sinicus} were fed on the four diets (\textit{I. galbana}, \textit{C. curvisetus}, \textit{E. huxleyi}, and \textit{Gymnodinium} sp.), no significant differences were found between DMS concentrations in the control (without antibiotics) and those in the treatment (with antibiotics) (data not shown). Therefore, the copepod cultures were not treated with antibiotics in our laboratory experiment to obtain axenicity. Yost and Mitchelmore (2009) reported that antibiotic treatment negatively affected algal growth, which was the other reason for not using antibiotics.
2.2.4 DMS and DMSP determinations

According to the methods of Andreae and Barnard (1983), DMS and DMSP concentrations were determined using the purge-and-trap technique by a gas chromatograph, which was equipped with a flame photometric detector (Shimadzu GC-14B). For DMS measurement, 10 mL seawater was directly introduced into a glass purge chamber. Gravity filtering of samples for DMSP\textsubscript{d} was obtained according to the method of Kiene and Slezak (2006) with minor modifications. The filtrate and unfiltered seawater for DMSP\textsubscript{d} and total DMSP (DMSP\textsubscript{c}, DMSP\textsubscript{d} + DMSP\textsubscript{p}) measurements were transferred to a 40 mL serum bottle containing 2 mL of 10 mol L\textsuperscript{-1} KOH solution and kept at 4 °C for at least 24 h to complete cleavage. DMSP\textsubscript{d} and DMSP\textsubscript{p} concentrations were determined by the total DMS subtracting the DMS in original seawater. The DMSP\textsubscript{c} and DMSP\textsubscript{e} concentrations were calculated by subtracting DMS and DMSP\textsubscript{d} in filtered water.

2.3 Statistical analysis

Data were expressed as mean ± standard deviation. Student’s \( t \) test and one-way ANOVA were used to determine the differences between control and treatment samples. Pearson correlations were utilized to assess the relationships between DMS/DMSP and IRs.

3 Results

3.1 Field experiment

3.1.1 Abiotic parameter

The temperature and salinity in Jiaozhou Bay are described in Fig. 2A. The salinity changed from 29.7 PSU to 32.3 PSU, and the lowest and highest salinities occurred in August 2010 (summer) and April 2011 (spring), respectively. The changing range of temperature was 1.72 °C to 25.3 °C, and the lowest and highest temperatures were observed in December 2010 (winter) and August 2010 (summer), respectively.

3.1.2 Biotic parameters

Chl \( a \) concentrations in surface water ranged from 0.8 \( \mu \text{g L}^{-1} \) to 11.0 \( \mu \text{g L}^{-1} \), with an average value of 3.1 \( \mu \text{g L}^{-1} \). Chl \( a \) concentration peaked on April 2011 (spring) (Fig. 2B). The bacterial abundance in surface water ranged from \( 5.7 \times 10^7 \) cells \( \text{L}^{-1} \) to \( 1.79 \times 10^8 \) cells \( \text{L}^{-1} \) (average of \( 9.68 \times 10^7 \) cells \( \text{L}^{-1} \)), with a peak value in September 2010 (autumn) (Fig. 2B).

A total of 74 species of zooplankton were identified and were categorized into 5 phyla, 6 classes, 17 orders, 44 families, and 55 genera. Among these species, 31 are Arthropoda, 22 are Coelenterata, 17 are planktonic larvae, 2 are Chaetognatha, 1 are Protozoa, and 1 are Tunicata (Table 2). The species composition of zooplankton varied with months, and the number of species ranged from 19 to 35. The lowest and highest numbers of species occurred in February 2011 (winter) and June 2010.
respectively. In total, 22 copepods were identified, and *Acartia pacifica*, *Acartia bifilosa*, *Centropages abdominalis*, *Eurytemora pacifica*, and *C. sinicus* were the dominant copepods (Table 2 and Fig. 3B).

Temporal variations in the mean zooplankton abundance are shown in Fig. 3A. *Noctiluca scintillans* appeared as the dominant zooplankton, contributing 91% of zooplankton abundance on June 2010. Copepods dominated the zooplankton community during winter and spring and accounted for 39% to 83% of the total zooplankton (Fig. 3A). The changing trend of copepod abundance showed that the minimum (7.5 individuals m⁻³) occurred on July 2010, subsequently increased, and peaked (596 individuals m⁻³) on April 2011. The *C. sinicus* abundance ranged from 0.143 individuals m⁻³ to 123 individuals m⁻³, and *C. sinicus* is the dominant copepod from October 2010 to May 2011. The *C. sinicus* abundance showed a similar trend to that of copepod, and *C. sinicus* exhibited the highest and lowest abundances in spring and summer, respectively (Fig. 3B).

### 3.1.3 Dilution experiments

Fig. 4 illustrates the principles of dilution experiments conducted at three stations (C3, D4, and E3) in Jiaozhou Bay during May 2011. At station C3, regression analysis of the dilution series with added nutrients yielded $\mu = 0.45$ d⁻¹, and $g = 0.12$ d⁻¹. At station D4, a nearshore at the east of Jiaozhou Bay, $\mu = 0.49$ d⁻¹, and $g = 0.11$ d⁻¹. In the present example for station E3, the Jiaozhou Bay mouth, $\mu = 0.23$ d⁻¹, and $g = 1.38$ d⁻¹, indicating that mortality in excess of growth ($\mu < g$). Phytoplankton growth rates for the nutrient addition treatments were higher than those for the no-nutrient treatments at the three stations.

The results of dilution experiment at three stations (C3, D4 and E3) are presented in Table 3, and $\mu$ varied from 0.02 d⁻¹ to 1.29 d⁻¹, with the highest value at station E3 (February 2011). $g$ varied from 0.02 d⁻¹ (June 2010) to 1.38 d⁻¹ (May 2011). Phytoplankton growth rates at coastal stations (D4 and E3) were higher than those at offshore station C3 (except July 2010 and May 2011).

### 3.1.4 DMS/DMSP

Changes in DMS/DMSP are presented in Fig. 5. The average concentrations of DMSP⁺, DMSP⁻, and DMS in surface water were 4.0 (1.1–9.9), 10.6 (3.3–21.2), and 1.9 (0.4–3.2) nmol L⁻¹, respectively. Compared with DMSP⁺ and DMS, DMSP⁻ has higher concentrations at the same periods. The highest concentrations of DMSP⁻, DMSP⁺, and DMS in Jiaozhou Bay were observed on September, September, and June 2010, respectively. In comparison, the lowest concentrations of DMSP⁻, DMSP⁺, and DMS were observed on November 2010, April 2011, and April 2011, respectively.

### 3.1.5 Relationships with biotic and abiotic parameters

In Jiaozhou Bay, many factors (biotic and abiotic parameters) could affect DMS and DMSP concentrations, and five factors (salinity, temperature, Chl a concentrations, bacterial abundance, and copepod abundance) were selected to investigate their effects on DMS and DMSP in this study. Salinity was negatively correlated with temperature and DMSP⁻ (p < 0.05). In comparison, Chl a concentrations were positively correlated with the copepod abundance (p < 0.01), and the copepod
abundance was positively correlated with *C. sinicus* abundance (p < 0.05). Furthermore, positive correlations were observed among DMS, DMSP<sub>d</sub>, and DMSP<sub>p</sub> (p < 0.01). In addition, no significant correlation was observed between DMS/DMSP and zooplankton, copepod, *C. sinicus* abundance or Chl *a* concentration (p > 0.05).

### 3.2 Incubation experiment

#### 3.2.1 Dietary effects on IR, CR, and DMS/DMSP

After *C. sinicus* was fed on four different diets, IRs changed from 0.13 × 10<sup>3</sup> cells copepod<sup>−1</sup> h<sup>−1</sup> to 5.55 × 10<sup>3</sup> cells copepod<sup>−1</sup> h<sup>−1</sup> and CRs ranged from 0.26 mL copepod<sup>−1</sup> h<sup>−1</sup> to 0.80 mL copepod<sup>−1</sup> h<sup>−1</sup> (Fig. 6A). Four species of algae affected IRs and CRs of *C. sinicus* differently, with *C. curvisetus* being the optimum diet for the grazing of *C. sinicus* (IR = 5.55 × 10<sup>3</sup> cells copepod<sup>−1</sup> h<sup>−1</sup>; CR = 0.80 mL copepod<sup>−1</sup> h<sup>−1</sup>), *I. galbana* being the second, *E. huxleyi* being the third, and *Gymnodinium* sp. being the fourth.

The ingestion of *C. sinicus* of four species of algae resulted in different DMS productions (Fig. 6B). With regard to *I. galbana* and *C. curvisetus*, *C. sinicus* grazing promoted DMS production in the treatments compared with the controls, e.g., DMS in *C. curvisetus* treatment by *C. sinicus* grazing was 1.7-fold of DMS in the controls. In comparison, DMS production has an opposite changing trend for *E. huxleyi* and *Gymnodinium* sp., in which the treatments showed lower DMS production than that in the controls.

For four algae species, DMSP<sub>p</sub> concentrations in the controls ranged from 48.78 nmol L<sup>−1</sup> to 7,165 nmol L<sup>−1</sup> and those in the treatments ranged from 27.25 nmol L<sup>−1</sup> to 11,055 nmol L<sup>−1</sup>. In the control and treatment groups, DMSP<sub>p</sub> concentrations of *E. huxleyi* and *Gymnodinium* sp. were two to three orders of magnitude higher than those of *I. galbana* and *C. curvisetus* (Fig. 6C). DMSP<sub>p</sub> concentrations in the treatments with copepod exhibited diverse changes for four algae species, DMSP<sub>p</sub> concentrations for *I. galbana* and *C. curvisetus* decreased, and DMSP<sub>p</sub> concentrations for *E. huxleyi* and *Gymnodinium* sp. increased. DMSP<sub>d</sub> in the treatments were higher than those in the controls (p > 0.05). DMSP<sub>d</sub> concentrations in the controls ranged from 10.60 nmol L<sup>−1</sup> to 6,595 nmol L<sup>−1</sup>. In comparison, DMSP<sub>d</sub> concentrations ranged from 11.39 nmol L<sup>−1</sup> to 10,848 nmol L<sup>−1</sup> in the treatments (Fig. 6D). DMSP<sub>d</sub> in the treatments were higher than those in the controls (p > 0.05).

Significant differences for the DMSP<sub>z</sub> and DMSP<sub>f</sub> contents of the four diets were observed (Fig. 6E). DMSP<sub>z</sub> contents decreased according the following order: *E. huxleyi* > *Gymnodinium* sp. > *I. galbana* > *C. curvisetus*. DMSP<sub>z</sub> contents of *C. sinicus* fed on *E. huxleyi* were 23.51-fold of those fed on *C. curvisetus* (Fig. 6E). When compared with DMSP<sub>z</sub> contents, DMSP<sub>f</sub> of four diets had different contents with the following order: *Gymnodinium* sp. > *E. huxleyi* > *I. galbana* > *C. curvisetus*. DMSP<sub>f</sub> contents of *C. sinicus* fed on *Gymnodinium* sp. were 126.3-fold of those fed on *C. curvisetus* (Fig. 6E).

For *C. curvisetus*, *I. galbana*, *Gymnodinium* sp., and *E. huxleyi*, DMSP<sub>z</sub> and DMSP<sub>f</sub> accounted for 0.58%, 3.2%, 0.57%, 0.14% and 0.16%, 1.04%, 0.70%, 0.43% of the total amounts of DMS/DMSP (DMSP<sub>z</sub> + DMSP<sub>f</sub> + DMS + DMSP<sub>d,p</sub>), respectively.
3.2.2 Food concentration experiment

When *I. galbana* concentrations increased, IRs gradually increased, peaked at $15 \times 10^4$ cells mL$^{-1}$, and subsequently declined (ranged from $2.45 \times 10^3$ cells copepod$^{-1}$ h$^{-1}$ to $5.12 \times 10^3$ cells copepod$^{-1}$ h$^{-1}$). In comparison, increased *I. galbana* concentrations induced the decrease in CRs, which ranged from 0.007 mL copepod$^{-1}$ h$^{-1}$ to 0.054 mL copepod$^{-1}$ h$^{-1}$ (Fig. 7A). DMS and DMSP$_d$ increased with the increase in *I. galbana* concentrations (Figs. 7B and 7C). DMS in the treatments were higher than those in the controls ($p < 0.05$). Moreover, DMS in the treatments (20.19–49.84 nmol L$^{-1}$) increased by 1.8% to 11% compared with that in the controls (18.23–48.96 nmol L$^{-1}$). DMSP$_p$ contents in the treatments (292.44–945.76 nmol L$^{-1}$) were lower than those in the controls (385.94–1,319.83 nmol L$^{-1}$) because of grazing activity.

DMSP$_d$ increased initially, peaked at $15 \times 10^4$ cells mL$^{-1}$, and decreased with the increase in *I. galbana* concentrations (Fig. 7D). DMSP$_d$ contents in the treatments (23.54–235.94 nmol L$^{-1}$) were higher than those in the controls (22.65–207.86 nmol L$^{-1}$) ($p > 0.05$), and those in the treatments increased by 97.95% compared with that in the controls ($20 \times 10^4$ cells mL$^{-1}$). A positive correlation between IR and the increment of DMS + DMSP$_d$ was observed ($r = 0.623$, $p = 0.377$, $n = 4$). In addition, a positive correlation between algal concentration and the decrement of DMSP$_p$ was discovered ($r = 0.767$, $p = 0.233$, $n = 4$).

DMSP$_d$ and DMSP$_t$ increased with the increase in *I. galbana* concentrations, peaked at $15 \times 10^4$ cells mL$^{-1}$, and declined (Fig. 7E). A significantly positive relationship was observed between IR and DMSP$_t$ ($r = 0.99$, $p = 0.01$, $n = 4$), accompanied by irrelevance between IR and DMSP$_d$ ($r = 0.07$, $p = 0.93$, $n = 4$). DMSP$_d$ and DMSP$_t$ of four *I. galbana* concentrations ($10 \times 10^4$, $15 \times 10^4$, $20 \times 10^4$, and $25 \times 10^4$ cells mL$^{-1}$) accounted for 1.9%, 4.5%, 3.8%, 2.5% and 3.3%, 2.4%, 1.4%, 1.2% of the sum of DMS/DMSP (DMSP$_d$ + DMSP$_t$ + DMS + DMSP$_{d,p}$) of *I. galbana*, respectively. We observed significant positive correlations between DMS and DMSP$_p$ ($r = 0.893$, $p = 0.003$, $n = 8$), accompanied by significant negative correlations between DMS and DMSP$_d$ ($r = -0.847$, $p = 0.008$, $n = 8$). In addition, significant negative correlations between DMSP$_d$ and DMSP$_p$ were noted in this study ($r = -0.804$, $p = 0.016$, $n = 8$).

3.2.3 Effects of salinity on IRs and DMS/DMSP production

The salinity experiments showed that IRs and CRs increased with the increase in salinities, peaked at 30 PSU (IR = 3.18 × $10^3$ cells copepod$^{-1}$ h$^{-1}$; CR = 0.23 mL copepod$^{-1}$ h$^{-1}$), and declined (Fig. 8A).

In this study, Fig. 8B shows that increased salinities restrained the production of DMS. Moreover, DMS contents in the treatment decreased from 84.54 nmol L$^{-1}$ to 26.54 nmol L$^{-1}$ when salinities increased, which was consistent with the DMS changes in the control (reduction from 41.11 nmol L$^{-1}$ to 25.90 nmol L$^{-1}$). The DMSP$_d$ changes indicated a trend similar to those of DMS contents, that is, increasing salinities decreased DMSP$_d$ contents. Fig. 8D shows that DMSP$_d$ contents decreased from 481.41 nmol L$^{-1}$ to 72.35 nmol L$^{-1}$ in the treatment. In comparison, DMSP$_d$ contents in the control reduced from 423.99 nmol L$^{-1}$ to 66.80 nmol L$^{-1}$. DMS contents of the treatments were higher than those of the controls ($p > 0.05$) and increased by 106%, 51%, 4.7%, and 2.4% at the salinities of 20, 25, 30, and 35 PSU, respectively.
The increasing salinities facilitated the accumulation of DMSP_p in the controls and treatments (Fig. 8C). DMSP_p contents in the treatments increased from 92.20 nmol L⁻¹ to 371.49 nmol L⁻¹, and those in the controls rose from 121.57 nmol L⁻¹ to 532.16 nmol L⁻¹. DMSP_p contents in the controls and treatments at 30 PSU were 2.9- and 1.7-fold of those at 25 PSU, respectively. DMSP_p contents in the treatments were lower than those in the controls because of grazing activity (p > 0.05). Furthermore, data analysis showed a positive correlation between IR and the decrement of DMSP_p (r = 0.945, p = 0.055, n = 4).

DMSP_d contents in the treatments were higher than those in the controls (p > 0.05) (Fig. 8D). The relationship between IR and the increment of DMS + DMSP_d proved to be positive (r = 0.662, p = 0.338, n = 4). In comparison, the correlation between the increments of DMS and DMSP_d was negative (r = −0.955, p = 0.045, n = 4).

DMSP_z ranged from 0.21 nmol L⁻¹ to 5.38 nmol L⁻¹, reached the maximum content at 35 PSU, and reached the minimum content at 20 PSU. When salinities increased, DMSP_z initially increased, subsequently declined, and finally increased (Fig. 8E). Maximum DMSP_z contents at 35 PSU were 25.6-fold, 4.2-fold, and 24.5-fold of those at 20, 25, and 30 PSU, respectively. DMSP_f had a similar trend to DMSP_z in the salinity experiments, and its changing range was 0.78 nmol L⁻¹ to 3.44 nmol L⁻¹. DMSP_f reached the minimum at 20 PSU and the peak at 25 PSU. Compared with 20, 30, and 35 PSU, DMSP_f at 25 PSU increased by 341%, 98%, and 2%, respectively. At 35 PSU, DMSP_z and DMSP_f attained the maximum sum (8.75 nmol L⁻¹), which was 8.84-fold of the minimum at 20 PSU. With regard to DMS/DMSP (DMS + DMSP_d + DMSP_z + DMSP_p) of I. galbana, DMSP_z of the four salinities (20, 25, 30, and 35 PSU) accounted for 0.035%, 0.2%, 0.05%, and 1.1%, and DMSP_f of the four salinities accounted for 0.13%, 0.56%, 0.41%, and 0.70%, respectively. Pearson correlation analysis did not detect a significant correlation between IR and DMSP_z (r = 0.37, p = 0.63, n = 4), which coincided with the relationship between IR and DMSP_f (r = 0.30, p = 0.70, n = 4). In addition, DMSP_p was significantly negatively correlated with DMS (r = −0.721, p = 0.044, n = 8) and DMSP_d (r = −0.918, p = 0.001, n = 8). In comparison, DMSP_p was observed to be positively correlated with DMSP_f (r = 0.491, p = 0.509, n = 4) and DMSP_z (r = 0.705, p = 0.295, n = 4).

4 Discussion

4.1 Effects of C. sinicus on DMS/DMSP in the field study

In Jiaozhou Bay, C. sinicus was the dominant copepod in 74 species of zooplankton identified from years 2010 to 2011. In this study, no significant correlations were observed between zooplankton, copepod, and/or C. sinicus abundance and DMS/DMSP concentrations in the field study, illustrating that Jiaozhou Bay was a complex ecosystem with different abundances and types of phytoplankton, zooplankton, and/or copepod in the natural environment. Many kinds of copepods inhabited Jiaozhou Bay, and their IRs depended on copepod species, e.g., IRs of Harpacticus sp. (Yu et al., 2015) were 10-fold of those of C. sinicus in this study. The gut contents of C. sinicus were checked, and the results showed that the C. sinicus preferred to graze on diatom Chaetoceros curvisetus and Thalassiosira nordenskoldi (data not shown), suggesting that diatom (DMSP-poor algae) were the preferable diet for C. sinicus. Zheng et al (2014) and Luo et al (2016) have
investigated the species composition and abundance of phytoplankton in 2010 and 2011 Jiaozhou Bay, respectively. According to their data, the changing trends of species composition, abundance of phytoplankton and dinoflagellate/diatom ratio from June 2010 to May 2011 were presented in Table 4 and Fig. 9. The predominancy of dinoflagellate Ceratium fusus was 0.10 in September 2010 (Table 4). The dinoflagellate/diatom ratios in the three months (July, August and September) were high among the whole year, and the abundances of dinoflagellate and diatom in September were the highest among the three months (Fig. 9), what is more, the bacterial abundance in September was the highest among the year (Fig. 2). Therefore, the occurrence of high abundances of dinoflagellate and bacteria might were the reason of high DMS and DMSP in September 2010. In February, April and May 2011, the dominant phytoplankton were diatom Rhizosolenia delicatula, Skeletonema costatum, and Skeletonema costatum, and the predominancies were 0.7, 0.99 and 0.68, respectively (Table 4).

Although the phytoplankton abundances and Chl a contents were high during January 2011 to May 2011, the DMSPₚ and DMSPₚ concentrations were lower than those in September 2010, suggesting that DMSP concentration not only depend on phytoplankton abundance, but also phytoplankton species and other factors. We evaluated the effects of several agents (i.e., food, diet concentration, and salinity) on DMS and DMSP productions in the laboratory study. Our incubation data showed that copepod grazing increased DMS production, which was consistent with previous investigations on the effects of copepod grazing on DMS production (Dacey and Wakeham, 1986; Yu et al., 2015). Consistent with our field study results, no significant correlations between mesozooplankton abundance and the distribution of DMS or DMSPₚ were also observed in the Gulf of Maine and St. Lawrence in previous studies (Cantin et al., 1996; Matrai and Keller, 1993). Cantin et al. (1996) concluded that mesozooplankton grazing played a minor role in DMS and DMSPₚ productions in the Gulf of St. Lawrence. The variable conditions in the natural environment can explain the reason for the minor role of zooplankton on DMS/DMSP production and the inconsistent results of field and incubation studies.

4.2 Dietary effects on copepod grazing and DMS/DMSP production

Different diets contain various DMSP contents, which affected DMS production induced by copepod grazing. DMSP-rich algae (E. huxleyi and Gymnodinium sp.) were not the preferable food for copepod C. sinicus in the incubation study because the released acrylic acid from DMSP in algae prevents them from copepod grazing. Based on the species of the diets, algae produced the corresponding DMS and DMSP (DMSPₚ and DMSPₚ) contents, which might be different. For example, when comparing Gymnodinium sp. and E. huxleyi with C. curvisetus and I. galbana, we determined that cellular DMSPₚ in Gymnodinium sp. and E. huxleyi were one to two orders of magnitude higher than those in C. curvisetus and I. galbana (see Table 1). In terms of I. galbana, our detection results on cellular DMSPₚ production were consistent with those reported by Niki et al. (2000), indicating that cellular DMSPₚ production in given algae was approximately invariable.

When Gymnodinium sp. and E. huxleyi (DMSP-rich phytoplankton) were grazed by C. sinicus, DMSP released from algae protected them from being grazed and stimulated the increase in DMSPₚ. In the natural environment, acrylic acid or DMSP depressed copepod appetites and forced them to ingest other DMSP-poor phytoplankton. Thus, DMSP-poor C. curvisetus became the favorite diet of copepods among four algae species, elucidating that the C. sinicus fed on C. curvisetus, which
evidently promoted DMS production in this study. Further studies have revealed that DMSP$_d$ was a feeding inhibitor, and DMSP$_p$, DMSP$_a$, DMS, and acrylic acid constituted a cellular antioxidant system involved in the scavenging of hydroxyl radicals (Strom et al., 2003; Sunda et al., 2002). Non-DMS-producing phytoplankton species in a mixture of prey were preferentially selected by grazers, and single DMS/DMSP-rich diet decreased the food intake of copepods, e.g., copepod *Harpacticus* sp. had inferior IRs and PPRs when fed on DMS/DMSP-rich alga *Prymnesium parvum* (Wolfe et al., 1997; Wolfe and Steinke, 1996; Yu et al., 2015).

Our results showed that DMSP in copepod bodies and fecal pellets accounted for 0.035% to 4.5% and 0.13% to 3.3% of DMS and DMSP in this study, illustrating that the ingestion of *C. sinicus* transferred DMSP from phytoplankton to the copepod bodies and fecal pellets. When compared with *Harpacticus* sp., lower DMSP$_f$ and DMSP$_e$ contributions in *C. sinicus* attributed to the lower IRs of *C. sinicus*.

Based on the *C. sinicus* abundance on May 2011, the maximum DMSP$_z$ (0.02–7 nmol L$^{-1}$) and DMSP$_f$ (0.1–12 nmol L$^{-1}$) of *C. sinicus* in the seawater of Jiaozhou Bay were achieved. The results of our study and that of Tang (2001) showed that DMSP$_p$ in copepod bodies and fecal pellets was an essential part of DMSP flux in the ocean. In addition, another sink of DMSP in the ocean was achieved by microbial processes. The results of Tang et al. (2001) and Dong et al. (2013) indicated that copepods and their pellets harbored a dense population of DCB (DMSP-consuming bacteria), which played an important role in DMSP degradation.

DMSP$_z$ and DMSP$_f$ contents were strongly associated with *C. sinicus* grazing. Their changing trends were consistent with that of IR, which depended on the critical concentration of phytoplankton. IRs of *C. sinicus* increased steadily with the increase in algal concentration below the critical concentration and decreased above the critical concentration. Critical concentration differed depending on the copepod and algal species, which were confirmed by this study and other investigations (Yu et al., 2015).

### 4.3 Effects of salinity on copepod grazing and DMS/DMSP

In this study, low salinity induced high DMS production in seawater, whereas high salinity increased DMSP$_p$ (intracellular DMSP) and decreased DMSP$_d$, which were consistent with the conclusion of Variamuthy et al. (1985). Our results were in accordance with the observation of a benthic diatom documented by van Bergeijk et al. (2003) and *Skeletonema costatum* documented by Yang et al. (2011). Intracellular DMSP, as an osmotically active compound, was accumulated or released to help algal cells adjust their osmotic potential when salinity increased or decreased (Kirst, 1996). *C. sinicus* grazing promoted DMS production in this study, which was consistent with the results obtained by many investigators (Belviso et al., 1990; Christaki et al., 1996; Daly and DiTullio, 1996; Leck et al., 1990). The decrement of DMSP$_p$ changed into DMS and DMSP$_d$ by copepod grazing, which was consistent with the positive relationships between IR and the increment of DMS + DMSP$_d$ and/or the decrement of DMSP$_p$. The increment of DMS was significantly negatively correlated with the increment of DMSP$_d$, indicating that the cleavage of DMSP$_d$ was the source of DMS by DMSP lyase. Many reports on the location of DMSP lyase in the cell have been published. Wolfe and Steinke (1996) indicated that the DMSP lyase location of *E. huxleyi*
CCMP 370 was in the membrane bound inside cells. Stefels and Dijkhuizen (1996) reported that DMSP lyase of *Phaeocystis* was membrane-bound and located extracellularly. Cellular locations and functions of DMSP lyase might differ depending on algal species.

This study confirmed that DMS production was affected by primary producers, e.g., algae. DMS/DMSP in algae transferred to the food web by predation, and several researchers investigated the effects of zooplankton grazing on DMS production with copepods and krill, indicating that the breakage of algal cells through sloppy feeding may increase DMSP$_d$ production (Dacey and Wakeham, 1986; Kaamatsu et al., 2004). When DMSP$_p$ concentrations changed, DMSP$_f$ and DMSP$_z$ concentrations were altered correspondingly, which was verified by other results in which the DMSP defecation rate of copepod *Acartia tonsa* feeding on *Tetraselmis impellucida* (prasinophyte) was closely related to food concentration and DMSP$_z$ content (Tang, 2001). Salinity changes altered the osmotic pressure surrounding copepod and algae cells, which in turn adjusted DMSP (intracellular DMSP$_p$, DMSP in tissues, and DMSP in gut content). Copepods contained more DMSP at high salinity, indicating the osmoregulatory function of DMSP (Tang et al., 1999).

5 Conclusions

In the present study, field and incubation experiments were performed to investigate the effects of *C. sinicus* grazing on DMS production in Jiaozhou Bay. Copepods (*C. sinicus*) was the dominant copepod in Jiaozhou Bay and preferred to graze on diatom. Appropriate diets and salinities facilitated DMS/DMSP production, e.g., *C. sinicus* feeding on *I. galbana* and *C. curvisetus* exhibited increased DMS production at 30 PSU in the laboratory experiment. *C. sinicus* grazing promoted the productions of DMS and DMSP$_d$, and DMS was released mainly from DMSP$_d$ and low salinity increased DMS production. Copepods and fecal pellets supplied substantial DMSP$_p$ into the water column and were important to the biogeochemical cycling of DMS.

*Competing interests.* The authors declare that they have no conflict of interest.

*Acknowledgements.* This work was financially supported by the National Key Research and Development Program of China (Grant No. 2016YFA0601302), the National Natural Science Foundation for Creative Research Groups (Grant No. 41521064), the Major International Joint Research Project of NSFC (Grant No. 41320104008), the AoShan Talents Program supported by the Qingdao National Laboratory for Marine Science and Technology (Grant No. 2015ASTP), and the National Natural Science Foundation of China (Grant No. 41776085). This study is MCTL Contribution No. 156.

*References*

Agostini, V.O., Macedo, A.J., and Muxagata E.: Evaluation of antibiotics as a methodological procedure to inhibit free-living and biofilm bacteria in marine zooplankton culture, An. Acad. Bras. Ciênc., 88, 733–746.
http://dx.doi.org/10.1590/0001-3765201620150454, 2016.

Andreae, M.O. and Barnard, W.R.: Determination of trace quantities of dimethyl sulfide in aqueous solutions, Anal. Chem., 55, 608–612, https://doi.org/10.1021/ac00255a006, 1983.

Belviso, S., Kim, S.K., Rassoulzadegan, F., Kraja, B., Nguyen, B.C., Mihalopoulos, N., and Buat-Menard, P.: Production of dimethylsulmonium propionate (DMSP) and dimethylsulfide (DMS) by a microbial food web, Limnol. Oceanogr., 35, 1810–1821, https://doi.org/10.4319/lo.1990.35.8.1810, 1990.

Brodsky, K.A.: Variability and systematic of the species of the genus Calanus Copepoda. 1. *Calanus pacificus* Brodsky and *Calanus sinicus* Brodsky sp. n, Exp. Fauna Seas., 3, 22–71, 1965.

Cantin, G., Levasseur, M., Gosselin, M., and Michaud, S.: Role of zooplankton in the mesoscale distribution of surface dimethylsulfide concentrations in the Gulf of St. Lawrence, Canada, Mar. Ecol. Prog. Ser., 141, 103–117, https://doi.org/10.3354/meps141103, 1996.

Charlson, R.J., Lovelock, J.E., Andreae, M.O., and Warren, S.G.: Oceanic phytoplankton, atmospheric sulphur, cloud albedo, and climate, Nature, 326, 655–661, https://doi.org/10.1038/326655a0, 1987.

Christaki, U., Belviso, S., Dolan, J.R., and Corn, M.: Assessment of the role of copepods and ciliates in the release to solution of particulate DMSP, Mar. Ecol. Prog. Ser., 141, 119–127, https://doi.org/10.3354/meps141119, 1996.

Dacey, J.W.H. and Wakeham, S.G.: Oceanic dimethylfide: production during zooplankton grazing on phytoplankton, Science, 233, 1314–1316, https://doi.org/10.1126/science.233.4770.1314, 1986.

Daly, K.L. and DiTullio, G.R.: Particulate dimethylsulfinopropionate removal and dimethylsulfide production by zooplankton in the Southern Ocean, in: Biological and environmental chemistry of DMSP and related sulfonium compounds, edited by: Kiene, R.P., Visscher, P.T., Keller, M.D., Kirst, G.O., Springer, Boston, MA, 223–238, 1996.

Dong, Y., Yang, G.-P., and Tang, K.M.: Dietary effects on abundance and carbon utilization ability of DMSP-consuming bacteria associated with the copepod *Acartia tonsa* Dana, Mar. Biol. Res., 9, 809–814, https://doi.org/10.1080/17451000.2013.765587, 2013.

Frost, B.W.: Effects of size and concentration of food particles on the feeding behavior of the marine planktonic copepod *Calanus Pacificus*, Limnol. Oceanogr., 17, 805–815, https://doi.org/10.4319/lo.1972.17.6.0805, 1972.

Guillard, R.R.L.: Culture of phytoplankton for feeding marine invertebrates, in: Culture of Marine Invertebrate Animals, edited by: Smith, W.L., Chanley, M.H., Springer, Boston, MA, 29–60, 1975.

Kasamatsu, N., Kawaguchi, S., Watanabe, S., Odate, T., and Fukuchi, M.: Possible impacts of zooplankton grazing on dimethylsulfide production in the Antarctic Ocean, Can. J. Fish. Aquat. Sci., 61, 736–743, https://doi.org/10.1139/f04-072, 2004.

Kettle, A.J., Andreae, M.O., Amouroux, D., Andreae, T.W., Bates, T.S., Berresheim, H., Bingemer, H., Boniforti, R., Curran, M.A.J., DiTullio, G.R., Helas, G., Jones, G.B., Keller, M.D., Kiene, R.P., Leck, C., Levasseur, M., Malin, G., Maspero, M., Matrai, P., McTaggart, A.R., Mihalopoulos, N., Nguyen, B.C., Novo, A., Putaud, J.P., Rapsomanikis, S., Roberts, G., Schebeske, G., Sharma, S., Simó, R., Staubes, R., Turner, S., and Uher, G.: A global database of sea surface
dimethylsulfide (DMS) measurements and a procedure to predict sea surface DMS as a function of latitude, longitude, and month, Global Biogeochem. Cy., 13, 399–444, https://doi.org/10.1029/1999GB900004, 1999.

Kiene, R.P. and Linn, L.J.: Distribution and turnover of dissolved DMSP and its relationship with bacterial production and dimethylsulfide in the Gulf of Mexico, Limnol. Oceanogr., 45, 849–861, https://doi.org/10.4319/lo.2000.45.4.0849, 2000.

Kiene, R.P. and Slezak, D.: Low dissolved DMSP concentrations in seawater revealed by small-volume gravity filtration and dialysis sampling, Limnol. Oceanogr.: Methods, 4, 80–95, https://doi.org/10.4319/mom.2006.4.80, 2006.

Kiørboe, T.: Population regulation and role of mesozooplankton in shaping marine pelagic food webs, Hydrobiologia, 363, 13–27, https://doi.org/10.1023/A:1003173721751, 1997.

Kirst, G.O.: Osmotic adjustment in phytoplankton and macroalgae, in: Biological and environmental chemistry of DMSP and related sulfonium compounds, edited by: Kiene, R.P., Visscher, P.T., Keller, M.D., Kirst, G.O., Springer, Boston, MA, 121–129, 1996.

Lancelot, C. and Billen, G.: Carbon-nitrogen relationships in nutrient metabolism of coastal marine ecosystem, Adv. Aquat. Microbiol., 3, 263–321, 1985.

Landry, M.R. and Hassett, R.P.: Estimating the grazing impact of marine microzooplankton, Mar. Biol., 67, 283–288, https://doi.org/10.1007/BF00397668, 1982.

Leck, C., Larsson, U., Bågander, L.E., Johansson, S., and Hadju, S.: Dimethyl sulfide in the Baltic Sea: Annual variability in relation to biological activity, J. Geophy. Res., 95, 3353–3363, https://doi.org/10.1029/JC095iC03p03353, 1990.

Li, Y., Li, R., Wang, Z., Zhu, M., Sun, P., and Xia, B.: A preliminary study on phytoplankton community structure and its changes in the Jiaozhou Bay, Adv. Mar. Sci., 23, 328–334, 2005. (in Chinese with English abstract)

Luo, X., Sun, X.-X., Zheng, S., and Zhao, Y.-F.: On phytoplankton community structure and causative factors in the Jiaozhou Bay in 2011, Oceanologia Limnologia Sinica, 47, 915–923, https://doi.org/, 2016. (in Chinese with English abstract)

Matrai, P.A. and Keller, M.D.: Dimethylsulfide in a large-scale coccolithophore bloom in the Gulf of Maine, Contin. Shelf. Res., 13, 831–843, https://doi.org/10.1016/0278-4343(93)90012-M, 1993.

Nguyen, B.C., Belviso, S., Mihalopoulos, N., Gostan, J., and Nival, P.: Dimethyl sulfide production during natural phytoplanktonic blooms, Mar. Chem., 24, 133–141, https://doi.org/10.1016/0304-4203(88)90044-8, 1988.

Niki, T., Kunugi, M., and Otsuki, A.: DMSP-lyase activity in five marine phytoplankton species: its potential importance in DMS production, Mar. Biol., 136, 759–764, https://doi.org/10.1007/s002279900235, 2000.

Parsons, T.R., Maita, Y., and Lalli, C.M. (Eds.): A manual of chemical and biological methods for seawater analysis, Pergamon Press, Oxford, 184 pp., 1984.

Porter, K.G. and Feig, Y.S.: The use of DAPI for identifying and counting aquatic microflora. Limnol. Oceanogr., 25, 943–948, https://doi.org/10.4319/lo.1980.25.5.0943, 1980.

Stefels, J. and Dijkhuizen, L.: Characteristics of DMSP-lyase in Phaeocystis sp. (Prymnesiophyceae), Mar. Ecol. Prog. Ser., 131, 307–313, https://doi.org/10.3354/meps131307, 1996.
Strathmann, R.R.: Estimating the organic carbon content of phytoplankton from cell volume or plasma volume, Limnol. Oceanogr., 12, 411–418, https://doi.org/10.4319/lo.1967.12.3.0411, 1967.

Strom, S., Wolfe, G., Slajer, A., Lambert, S., and Clough, J.: Chemical defense in the microplankton II: Inhibition of protist feeding by β-dimethylsulfoniopropionate (DMSP), Limnol. Oceanogr., 48, 230–237, https://doi.org/10.4319/lo.2003.48.1.0230, 2003.

Sunda, W., Kieber, D.J., Kiene, R.P., and Huntsman, S.: An antioxidant function for DMSP and DMS in marine algae, Nature, 418, 317–320, https://doi.org/10.1038/nature00851, 2002.

Tang, K.W., Dam, H.G., Visscher, P.T., and Fenn, T.D.: Dimethylsulfoniopropionate (DMSP) in marine copepods and its relation with diets and salinity, Mar. Ecol. Prog. Ser., 179, 71–79, https://doi.org/10.3354/meps179071, 1999.

Tang, K.W., Fenn, T.D., Visscher, P.T., and Dam, H.G.: Regulation of body dimethylsulfoniopropionate (DMSP) content by the copepod Temora longicornis: a test of four mechanisms, Mar. Biol., 136, 749–757, https://doi.org/10.1007/s002270000309, 2000.

Tang, K.W., Visscher, P.T., and Dam, H.G.: DMSP-consuming bacteria associated with the calanoid copepod Acartia tonsa (Dana), J. Exp. Mar. Biol. Ecol., 256, 185–198, https://doi.org/10.1016/S0022-0981(00)00314-2, 2001.

Tang, K.W.: Defecation of dimethylsulfoniopropionate (DMSP) by the copepod Acartia tonsa as functions of ambient food concentration and body DMSP content, J. Plankton Res., 23, 549–553, https://doi.org/10.1093/plankt/23.5.549, 2001.

Turner, S.M., Malin, G., Nightingale, P.D., and Liss, P.S.: Seasonal variation of dimethyl sulphide in the North Sea and an assessment of fluxes to the atmosphere, Mar. Chem., 54, 245–262, https://doi.org/10.1016/0304-4203(96)00028-X, 1996.

Vairavamurthy, A., Andreae, M.O., and Iverson, R.L.: Biosynthesis of dimethylsulfide and dimethylpropiothetin by Hymenomonas carterae in relation to sulfur source and salinity variations, Limnol. Oceanogr., 30, 59–70, https://doi.org/10.4319/lo.1985.30.1.0059, 1985.

van Bergeijk, S.A., Van der Zee C., and Stal, L.J.: Uptake and excretion of dimethylsulphoniopropionate is driven by salinity changes in the marine benthic diatom Cylindrotheca closterium, Eur. J. Phycol., 38, 341–349, http://dx.doi.org/10.1080/09670260310001612600, 2003.

Verity, P.G., Robertson, C.Y., Tronzo, C.R., Andrews, M.G., Nelson, J.R., and Sieracki, M.E.: Relationships between cell volume and the carbon and nitrogen content of marine photosynthetic nanoplankton, Limnol. Oceanogr., 37, 1434–1446, https://doi.org/10.4319/lo.1992.37.7.1434, 1992.

Wolfe, G.V. and Steinke, M.: Grazing-activated production of dimethyl sulfide (DMS) by two clones of Emiliania huxleyi. Limnol. Oceanogr. 41, 1151–1160, https://doi.org/10.4319/lo.1996.41.6.1151, 1996.

Wolfe, G.V., Levasseur, M., Cantin, G., and Michaud, S.: DMSP and DMS dynamics and microzooplankton grazing in the Labrador Sea: application of the dilution technique, Deep-Sea Res. I, 47, 2243–2264, https://doi.org/10.1016/S0967-0637(00)00028-5, 2000.
Wolfe, G.V., Levasseur, M., Cantin, G., and Michaud, S.: DMSP and DMS dynamics and microzooplankton grazing in the Labrador Sea: application of the dilution technique, Deep Sea Res. I, 47, 2243–2264, https://doi.org/10.1016/S0967-0637(00)00028-5, 2000.

Wolfe, G.V., Steinke, M., and Kirst, G.O.: Grazing-activated chemical defence in a unicellular marine alga, Nature, 387, 894–897, https://doi.org/10.1038/43168, 1997.

Wong, C.S., Wong, S.E., Richardson, W.A., Smith, G.E., Arychuk, M.D., and Page, J.S.: Temporal and spatial distribution of dimethylsulfide in the subarctic northeast Pacific Ocean: a high-nutrient-low-chlorophyll region, Tellus B, 57, 317–331, https://doi.org/10.1111/j.1600-0889.2005.00156.x, 2005.

Yang, G., Li, C., and Sun, J.: Influence of salinity and nitrogen content on production of dimethylsulfoniopropionate (DMSP) and dimethylsulfide (DMS) by Skeletonema costatum, Chin. J. Oceanol. Limn., 29, 378–386, https://doi.org/10.1007/s00343-011-0087-6, 2011.

Yost, D.M. and Mitchelmore, C.L.: Dimethylsulfoniopropionate (DMSP) lyase activity in different strains of the symbiotic alga Symbiodinium microadriaticum, Mar. Ecol. Prog. Ser., 386, 61–70, https://doi.org/10.3354/meps08031, 2009.

Yu, J., Tian, J.-Y., and Yang, G.-P.: Effects of Harpacticus sp. (Harpacticoida, copepod) grazing on dimethylsulfoniopropionate and dimethylsulfide concentrations in seawater, J. Sea Res., 99, 17–25, https://doi.org/10.1016/j.seares.2015.01.004, 2015.

Zheng, S., Sun, X.-X., Zhao, Y.-F., and Sun, S.: Annual variation of species composition and abundance distribution of phytoplankton in 2010 in the Jiaozhou Bay, Mar. Sci., 38, 1–6, 2014. (in Chinese with English abstract)

Zhong, S.-L., Wang, Y.-P., Gao, S., Cao, Q.-Y., and Zhu, Y.-H.: Distribution patterns of nannofossil gephyrocapsa oceanica in surficial sediments of Jiaozhou Bay, southern Shandong Peninsula, China, Acta Palaeontol. Sin., 40, 505–513, 2001.
Table 1. Sizes and DMSP of phytoplankton cells.

| Algae            | Dimension (µm) | Biovolume (µm³) | Carbon (pg cell⁻¹) | DMSP (fmol cell⁻¹) |
|------------------|----------------|-----------------|--------------------|--------------------|
| C. curvisetus    | 7.8 × 4.6      | 9.04 × 1⁰       | 1.15 × 1⁰          | 1.0 × 1⁻¹           |
| I. galbana       | 6.0 × 5.8      | 1.11 × 1²       | 2.05 × 1¹          | 4.0 × 1⁰            |
| Gymnodinium sp.  | 14.0 × 12.0    | 1.13 × 1³       | 1.53 × 1²          | 3.6 × 1¹            |
| E. huxleyi       | 2.7 × 3.2      | 1.25 × 1⁰       | 2.94 × 1⁰          | 3.0 × 1¹            |
Table 2. Species of zooplankton in Jiaozhou Bay.

| Phylum          | Species                          | 2010.06 | 2010.07 | 2010.08 | 2010.09 | 2010.10 | 2010.11 | 2010.12 | 2011.01 | 2011.02 | 2011.04 | 2011.05 |
|-----------------|----------------------------------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|---------|
| Planktonic larvae | Trehospona larvae                | 1       |         |         |         |         |         |         |         |         |         |         |
|                 | Polychea larva                   | 1       | 1       | 1       | 1       | 1       | 1       | 1       | 1       | 1       |         |         |
|                 | Gastropod post larva             | 1       | 1       | 1       |          |         |         |         |         |         |         |         |
|                 | Bivalva larva                    | 1       | 1       | 1       | 1       |         |         |         |         |         |         |         |
|                 | Naupliar larva (Copepoda)        | 1       |         |         |         |         |         |         |         |         |         |         |
|                 | Naupliar larva (Clintipedia)     | 1       |         |         |         |         |         |         |         |         |         |         |
|                 | Cypria larva                     | 1       |         |         |         |         |         |         |         |         |         |         |
|                 | Macra larva                      | 1       | 1       | 1       | 1       |         |         |         |         |         |         |         |
|                 | Branchyrous zona larva           | 1       | 1       | 1       | 1       | 1       |         |         |         |         |         |         |
|                 | Megalopa larva                   | 1       | 1       | 1       |         |         |         |         |         |         |         |         |
|                 | Porcellana zona larva            | 1       | 1       | 1       |         |         |         |         |         |         |         |         |
|                 | Atlan larva                      | 1       | 1       | 1       |         |         |         |         |         |         |         |         |
|                 | Ophiopluteus larva               | 1       |         |         |         |         |         |         |         |         |         |         |
|                 | Echinopluteus larva              | 1       |         |         |         |         |         |         |         |         |         |         |
|                 | Echinothurius larva              | 1       | 1       | 1       | 1       |         |         |         |         |         |         |         |
|                 | Fish eggs                        | 1       |         |         |         |         |         |         |         |         |         |         |
|                 | Fish larva                       | 1       | 1       | 1       | 1       |         |         |         |         |         |         |         |

“1” means appearance.
Table 3. Growth and grazing rates of chlorophyll $a$ estimated from the dilution technique.

| Sampling time | Station | $\nu$ (d$^{-1}$) | $g$ (d$^{-1}$) | $r^2$ |
|---------------|---------|------------------|----------------|-------|
| 2010.06       | C3      | 0.31             | 0.26           | 0.52  |
|               | D4      | 0.44             | 0.16           | 0.20  |
|               | E3      | 0.68             | 0.65           | 0.57  |
| 2010.07       | C3      | 0.36             | 0.02           | 0.15  |
|               | D4      | 0.33             | 0.55           | 0.46  |
|               | E3      | 0.66             | 0.33           | 0.65  |
| 2010.08       | C3      | 0.38             | 0.27           | 0.59  |
|               | D4      | 0.61             | 0.60           | 0.76  |
|               | E3      | 0.71             | 0.45           | 0.48  |
| 2010.09       | C3      | 0.27             | 0.40           | 0.10  |
|               | D4      | 0.27             | 0.10           | 0.20  |
|               | E3      | 0.28             | 0.59           | 0.40  |
| 2010.10       | C3      | 0.85             | 0.22           | 0.47  |
|               | D4      | 0.88             | 0.44           | 0.39  |
|               | E3      | 0.93             | 0.96           | 0.28  |
| 2010.11       | C3      | 0.12             | 0.31           | 0.54  |
|               | D4      | 0.12             | 0.43           | 0.42  |
|               | E3      | 0.49             | 0.29           | 0.43  |
| 2010.12       | C3      | 0.13             | 0.37           | 0.61  |
|               | D4      | 0.96             | 0.15           | 0.25  |
|               | E3      | 0.36             | 0.30           | 0.71  |
| 2011.01       | C3      | 0.06             | 0.28           | 0.60  |
|               | D4      | 0.13             | 0.33           | 0.23  |
|               | E3      | 0.45             | 0.13           | 0.20  |
| 2011.02       | C3      | 0.02             | 0.49           | 0.25  |
|               | D4      | 1.12             | 1.15           | 0.65  |
|               | E3      | 1.29             | 0.67           | 0.40  |
| 2011.04       | C3      | 0.25             | 0.48           | 0.41  |
|               | D4      | 0.52             | 0.94           | 0.52  |
|               | E3      | 0.48             | 0.66           | 0.70  |
| 2011.05       | C3      | 0.45             | 0.12           | 0.59  |
|               | D4      | 0.49             | 0.11           | 0.52  |
|               | E3      | 0.23             | 1.38           | 0.66  |
Table 4. Dominant species and the predominancy of phytoplankton in the Jiaozhou Bay from June 2010 to May 2011 (cited from Zheng et al. (2014) and Luo et al. (2016)).

| Dominant species                  | Predominancy |
|-----------------------------------|--------------|
| Cossinodiscus asteromphalus       | 0.04 0.23 0.13 0.1 0.03 |
| C. wollei                       | 0.04 0.03 |
| Cossinodiscus spp.               | 0.03 0.06 0.03 0.07 0.13 |
| Actinoecys ehrenbergii           | 0.03 0.12 0.14 |
| Lauderia annulata                | 0.16 |
| Skeletonema costatum             | 0.17 0.33 0.03 |
| Leptocylindrus danicus           | 0.21 0.04 0.04 |
| Rhizosolenia delicatula          | 0.07 0.7 |
| R. imbricata                     | 0.13 0.02 |
| R. stolterfoltii                 | 0.04 0.04 0.11 |
| R. alata f. indica               | 0.02 |
| Gomardula flaccida               | 0.1 |
| Schroederella delicatula         | 0.05 |
| Thalassiosira nordenskottdii      | 0.07 |
| T. pacifica                      | 0.02 |
| Chaetoceros debilis              | 0.02 |
| C. didymus                       | 0.02 |
| C. densus                        | 0.45 0.06 0.14 |
| C. curvisetus                    | 0.05 |
| C. lorentzianus                  | 0.05 0.02 |
| C. pseudocurvisetus              | 0.05 |
| C. teres                         | 0.04 0.02 |
| Chaetoceros spp                  | 0.06 0.02 |
| Biddulphia sinensis              | 0.07 0.05 0.03 0.02 |
| Eucampia zoodiacus               | 0.99 0.68 |
|Skeletonema costatum              | 0.02 0.02 |
| Asteroecia, Kariama               | 0.05 0.02 0.03 |
| Dictyos sp.                      | 0.18 0.05 0.05 |
| Navicula membranacea             | 0.02 0.06 |
| Pseudonitzchia pungens           | 0.02 |
| Nitzchia paradoxa                 | 0.02 0.06 0.17 0.05 |
| Protoperidinium depressum        | 0.02 0.02 |
| Ceratium fusus                   | 0.12 0.04 0.1 0.03 0.02 |
| C. macrocera                      | 0.03 |
| C. tripos                        | 0.05 0.02 |
| Noctiluca scintillans            | 0.03 0.21 |

(Note: blank area means the predominancy < 0.02)
Figure 1. Location of fieldwork sampling stations in Jiaozhou Bay.
Figure 2. Monthly changes in temperature and salinity (A) and Chl a content and bacterial abundance (B) in Jiaozhou Bay.
Figure 3. Monthly changes in the mean abundances of zooplankton (A) and dominant copepod (B) in Jiaozhou Bay.
Figure 4. Examples of experimental analyses for dilution experiments conducted at three stations (C3, D4, E3) in Jiaozhou Bay during May 2011. Regression lines are fit to the data (filled circles) for incubations with added nutrients. Net growth rates for undiluted samples incubated without added nutrients are also shown (open circles).
Figure 5. Mean DMS and DMSP concentrations in surface seawater of 10 stations in Jiaozhou Bay.
Figure 6. Effects of *Calanus sinicus* grazing on IR and CR (A), DMS (B), DMSP$_p$ (C), DMSP$_d$ (D), and DMSP$_{z,f}$ (E) when they preyed on different diets. Error bars represent the standard deviation ($n = 3$).
Figure 7. Effects of *C. sinicus* grazing on IR and CR (A), DMS (B), DMSP$_p$ (C), DMSP$_d$ (D), and DMSP$_{z,f}$ (E) when they preyed on different concentrations of *I. galbana*. Error bars represent the standard deviation ($n = 3$).
Figure 8. Effects of *C. sinicus* grazing on IR and CR (A), DMS (B), DMSP$_p$ (C), DMSP$_d$ (D), and DMSP$_{z,f}$ (E) when they preyed at different salinities. Error bars represent the standard deviation ($n = 3$).
Figure 9. Total abundance of phytoplankton (diatom and dinoflagellate) (A) and the dinoflagellate/diatom ratio (B) in Jiaozhou Bay from June 2010 to May 2011 (cited from Zheng et al. (2014) and Luo et al. (2016)).