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Key Points:

- Global sea-air fluxes of DMS are quantified
- Effect of 25 years of modelled climate change is dominated by reduced wind
- Coccolithophores are estimated to have a small effect on DMS fluxes

Supporting Information:

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Exploiting satellite earth observation to quantify current global oceanic DMS flux and its future climate sensitivity

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Abstract We used coincident Envisat RA2 and AATSR temperature and wind speed data from 2008/2009 to calculate the global net sea-air flux of dimethyl sulfide (DMS), which we estimate to be 19.6 Tg S a⁻¹. Our monthly flux calculations are compared to open ocean eddy correlation measurements of DMS flux from 10 recent cruises, with a root mean square difference of $3.1 \,\mu$ mol m⁻² day⁻¹. In a sensitivity analysis, we varied temperature, salinity, surface wind speed, and aqueous DMS concentration, using fixed global changes as well as CMIP5 model output. The range of DMS flux in future climate scenarios is discussed. The CMIP5 model predicts a reduction in surface wind speed and we estimate that this will decrease the global annual sea-air flux of DMS by 22% over 25 years. Concurrent changes in temperature, salinity, and DMS concentration increase the global flux by much smaller amounts. The net effect of all CMIP5 modelled 25 year predictions was a 19% reduction in global DMS flux. 25 year DMS concentration changes had significant regional effects, some positive (Southern Ocean, North Atlantic, Northwest Pacific) and some negative (isolated regions along the Equator and in the Indian Ocean). Using satellite-detected coverage of coccolithophore blooms, our estimate of their contribution to North Atlantic DMS emissions suggests that the coccolithophore blooms, our estimate of their contribution to North Atlantic annual flux estimate, but may be more important in the summertime and in the northeast Atlantic.

1. Introduction

The global oceans are the largest natural source of atmospheric sulfur through the emission of the gas dimethyl sulfide (DMS) [*Bates et al.*, 1992]. Derived from the growth and decay of phytoplankton, DMS is ubiquitous in the surface ocean, with dissolved concentration grossly supersaturated relative to the overlying air, resulting in consistent emission to the atmosphere. Once in the marine atmosphere, the oxidation of DMS leads to sulfur dioxide, sulfuric acid, and sulfate aerosols, among other products. These acidic and hygroscopic species affect atmospheric chemistry [*Charlson and Rodhe*, 1982], and contribute to the nucleation of new particles as well as the growth of existing particles to cloud condensation nuclei (CCN). Such condensation nuclei affect the Earth's radiation budget (and thus climate) by directly scattering sunlight and indirectly influencing cloud physics and albedo [*Charlson et al.*, 1987].

Despite increasing anthropogenic sulfur emissions, DMS remains the predominant source of sulfur mass to the remote marine atmosphere [Yang et al., 2011b]. There is, however, much debate over the exact role that DMS plays in regulating our climate. Gunson et al. [2006] modeled a 0.8°C cooling for a doubling of DMS emission, and a 1.6°C warming for a halving of DMS emission. Furthermore, their modeled DMS emission increases over time in response to warming and decreases in response to cooling, consistent with a negative climatic feedback. Other studies suggest less climate sensitivity to DMS. For example, Boucher and Lohmann [1995] and Woodhouse et al. [2010] found the radiative properties of clouds to depend more on total sulfate and cloud droplet number concentrations (partly anthropogenic) than on DMS and naturally derived sulfate. Similarly, Quinn and Bates [2011] argued that the climate regulation by DMS is weak mostly because of other sources of CCN, e.g., sea salt and organics, while Clarke et al. [2013] found the main source of CCN in the marine boundary layer (MBL) to be entrainment of preexisting aerosols from the free troposphere rather than nucleation and growth within the MBL. The impact of DMS on clouds is thought to be most significant in regions with minimal competitions from anthropogenic CCN [Twomey, 1991]. Boucher et al. [2003] showed large spatial heterogeneity in their modeled indirect radiative forcing due to DMS-derived aerosols, varying from close to zero to -5 Wm^{-2} . Woodhouse et al. [2013] recently demonstrated highly variable sensitivity of CCN production as a function of DMS emission and argued for the importance of

monitoring of future changes in spatial DMS distributions. *Six et al.* [2013] argued that ocean acidification would reduce DMS emissions, resulting in a radiative forcing of 0.4 Wm^{-2} .

Previous estimates for the present day global sea-to-air flux of DMS range from 15 to 33 Tg S a⁻¹ [*Elliott*, 2009; *Kettle and Andreae*, 2000; *Simó and Dachs*, 2002]. Based on the most recent DMS climatology, *Lana et al.* [2011] estimated a mean annual emission of 28.1 Tg S a⁻¹, with a range from 17.6 to 34.4 Tg S a⁻¹ depending on the choice of transfer velocity parameterization (see subsection 4.1). Predictions of future DMS emission vary significantly, especially on a regional scale [*Cameron-Smith et al.*, 2011; *Halloran et al.*, 2010] due to combined uncertainties in the changes of seawater DMS concentration, winds, temperature, etc. Observations and recent modeling studies indicate that the concentration of DMS in surface seawater is the net result of rapid but variable biological production and consumption processes [*Galí et al.*, 2013]. A small perturbation in DMS source or sink can significantly influence the amount of seawater DMS available for air-sea exchange.

Earth observation (EO) data from satellites have the potential to help predict regional changes in DMS emissions. In the Arctic, where in situ observations are rare, the use of EO data underpinned by field data is currently the most feasible way of reliably monitoring sea-air gas fluxes. EO data have previously been used to derive sea-air fluxes of CO_2 at the global scale [e.g., *Boutin et al.*, 2002], and the improved quasi-global availability of EO data has facilitated recent studies of CO_2 sea-air exchange over a substantial area of the Arctic Ocean [*Arrigo et al.*, 2010; *Else et al.*, 2008]. In comparison, the use of EO data to study sea-air gas exchange of trace gases including DMS is novel. Such methods have the potential to improve our understanding of the global sulfur cycle and its impact on climate.

One specific example of EO data utility is the potential to constrain the contribution to the global DMS flux from coccolithophore blooms. Algal species produce a large range of intracellular dimethylsulfoniopropionate (DMSP), which is the major algal precursor to DMS [*Keller et al.*, 1989]. Diatoms are low DMSP/DMS producers, with intracellular DMSP on the order of 0.86 mmol/mol C [*Stefels et al.*, 2007]. In sharp contrast, the haptophyte phytoplankton group, which contains coccolithophores, has a higher intracellular DMSP content (11 mmol/mol C), which is second only to dinoflagellates (22 mmol/mol C) [*Stefels et al.*, 2007]. In the open ocean, certain coccolithophore blooms are clearly detectable by satellites. Some blooms have been observed to produce substantially elevated DMSP and DMS concentrations and are thought to contribute substantially to the DMS sea-to-air flux [*Malin et al.*, 1993; *Matrai and Keller*, 1993; *Steinke et al.*, 2002a].

Globally and regionally, predicting future DMS emissions requires predictions of (1) surface seawater DMS concentration; (2) changes in forcing parameters (e.g., wind speed, SST, salinity), and (3) the sensitivity of DMS flux to these forcing parameters.

Here we use spatially and temporally coincident EO data from the European Space Agency's (ESA) Envisat satellite to estimate the global flux of DMS in 2008 and 2009. We then evaluate the sensitivity of sea-air DMS exchange in the global ocean to changes in temperature, salinity, wind speed, and aqueous DMS concentration, both using fixed scalar changes in conditions and arising from future climate model scenarios.

2. Methods

2.1. Data Inputs

We used EO data from Radar Altimeter 2 (RA2) and Advanced Along Track Scanning Radiometer (AATSR) on ESA's Envisat satellite. RA2 is an altimeter that can provide estimates of wind speed at 10 m height (U_{10}) with a small surface footprint. In the absence of clouds, AATSR provides estimates of sea surface skin temperature SST_{skin} over a much wider swath [*Donlon et al.*, 2007]. The fields of view of the two instruments overlap, providing spatially and temporally coincident measurements. Globally, RA2 has an error standard deviation of 1.25 ms⁻¹ and a bias of -0.28 ms^{-1} [*Queffeulou et al.*, 2010], while AATSR has a standard deviation of 0.16°C and bias of $+0.2^{\circ}$ C [O'Carroll et al., 2008]. We processed the RA2 and AATSR data for 2008–2009 using standard ESA methods and binned them to a 1° \times 1° geographic grid, see [Land et al., 2013] for details.

Climatological values of the bulk concentration of DMS in surface seawater (DMS_w) were obtained on a $1^{\circ} \times 1^{\circ}$ grid from the Lana et al. [2011] climatology, while climatological salinity values were obtained on the same grid from the NOAA National Oceanographic Data Center (http://www.nodc.noaa.gov/OC5/

Table 1. Summary of DMS Concentrations and Fluxes From 10 Cruises, Compared to Fluxes From This Study										
Cruise	TAO	PHASE	BIO	Knorr 06	DOGEE a ^a	DOGEE b ^a	Knorr 07	SO GasEx	VOCALS	Knorr 11
Location	Equatorial Pacific	North Pacific	Sargasso Sea	South Pacific	Northeast Atlantic	Northeast Atlantic	North Atlantic	Southern Ocean	Southeast Pacific	North Atlantic
Latitude ^b	-1	5	30	-16	43	54	48	-51	-20	50
Longitude ^b	-98	-179	-66	-106	-17	-11	-51	-38	-80	-45
Time	Nov 2003	May–Jun 2004	Jul-Aug 2004	Jan 2006	Jun–Jul 2007	July 2007	July 2007	Mar–Apr 2008	Oct-Nov 2008	Jun–Jul 2011
DMS_w^c	2.6 (0.8)	1.7 (0.7)	2.6 (0.4)	3.8 (2.2)	1.2 (0.6)	6.9 (3.1)	2.5 (1.1)	1.6 (0.7)	2.8 (1.1)	4.1 (2.0)
DMS_a^c	0.0064 (0.0020)	0.0036 (0.0020)	0.0056 (0.0022)	0.0140 (0.0152)	0.0053 (0.0034)	0.0203 (0.0073)	0.0125 (0.0087)	0.0052 (0.0024)	0.0024 (0.0021)	0.0167 (0.0127)
DMS_w/DMS_a	406	470	464	271	226	340	200	308	1120	245
In Situ Flux ^d	7.1 (3.7)	5.1 (3.1)	6.2 (2.4)	12.1 (15.0)	2.7 (2.4)	18.1 (9.4)	5.9 (4.9)	2.9 (2.1)	3.4 (1.9)	7.6 (7.6)
EO Flux ^e	3.2 (-1.1)	5.0 (0.0)	5.1 (-0.5)	8.8 (-0.2)	6.6 (1.6)	12.0 (-0.6)	2.5 (-0.7)	2.6 (-0.1)	4.9 (0.8)	6.4 (-0.2)
Publication	Huebert	Marandino	Blomquist	Marandino	Huebert	Huebert	Marandino	Yang	Yang	Bell
	et al. [2004]	et al. [2007]	et al. [2006]	et al. [2009]	et al. [2010]	et al. [2010]	et al. [2008]	et al. [2011a]	et al. [2009]	et al. [2013]

^aThese two are the same cruise but two distinct water masses.

^bCruise Mean, Degrees N or E.

^cCruise Mean, μ mol m⁻³ = nM.

^dCruise mean (standard deviation), μ mol m⁻² day⁻¹.

^eMean of all cruise months in a 5 $^\circ$ imes 5 $^\circ$ window (deviation from cruise mean/cruise standard deviation).

WOA09/pr_woa09.html). Global daily air pressure fields were provided by the European Centre for Mediumrange Weather Forecasting (ECMWF) operational data set (N80 Gaussian gridded analysis on surface levels; ERA-40 format, 6 h fields). The pressure data were reprojected from their original $2.5^{\circ} \times 2.5^{\circ}$ grid to the 1° imes 1° grid using linear interpolation. Daily sea ice coverage data were provided by the Operational Sea Surface Temperature and Sea Ice Analysis (OSTIA) system [Donlon et al., 2011; Stark et al., 2008] based on the EUMETSAT Ocean and Sea Ice Satellite Application Facility (OSI-SAF) daily analyses. These daily global OSTIA sea ice coverage data were averaged from their original $0.05^{\circ} \times 0.05^{\circ}$ grid to the same $1^{\circ} \times 1^{\circ}$ grid. We used a 4096 \times 2048 global land mask ($0.09^{\circ} \times 0.09^{\circ}$) to determine the proportion of land by area in each $1^{\circ} \times 1^{\circ}$ pixel. The land mask is needed to calculate the net fluxes and the use of a higher spatial resolution subgrid for the land mask gives a more accurate representation of the integrated flux from coastal pixels.

2.2. Calculating Sea-Air DMS Fluxes

We estimated the sea-to-air flux of DMS (F) in μ mol S m⁻² day⁻¹ as the product of the gas transfer velocity, K_w (cm h⁻¹), and the difference in DMS concentration (nM) between the base [DMS_{agw}] and the top $[DMS_{aao}]$ of the mass boundary layer at the sea surface, multiplied by 0.24 to convert K_w into m day⁻¹:

$$F = 0.24K_W \left(\left[DMS_{aqw} \right] - \left[DMS_{aq0} \right] \right). \tag{1}$$

 $[DMS_{aqw}]$ is set to the bulk concentration DMS_w from Lana et al. [2011], while $[DMS_{aqv}]$ is set to $H DMS_{ar}$ where H is the dimensionless Henry's Law solubility from liquid to gas, calculated as a function of temperature and salinity using the methods of Johnson [2010], and DMS_a is the bulk concentration of DMS in air, giving

$$F = 0.24K_w (DMS_w - H \times DMS_a).$$
⁽²⁾

 DMS_a (units 10⁻⁹ mol/L of air) is estimated as DMS_w (in units of nM) divided by 400, where both concentrations are expressed in equivalent units. The dimensionless water/air ratio of 400 represents an order of magnitude value based on observations from 10 open ocean cruises (see Table 1). This fixed ratio clearly does not capture variations on small temporal or spatial scales, but is intended to be broadly representative of the ratio on a regional to global level. The relative importance of the $H \times DMS_a$ term varies with temperature (and slightly with salinity) due to changes in H. In 2 years of monthly global data at 1° spatial resolution, $H \times DMS_a$ varies from 2% of DMS_w in the warmest waters to 7% in the coldest.

 K_w is the reciprocal of the total resistance to gas transfer on both sides of the air/water interface [Liss and Slater, 1974], given by

$$1/K_w = 1/k_w + H/k_a \tag{3}$$

Daily maps of the in-water gas transfer velocity k_w were derived from the SST_{skin} and U₁₀ data using the wind-based parameterization of Goddijn-Murphy et al. [2012], who compared all available DMS eddy correlation measurements of gas-transfer velocity with altimeter-derived U_{10} :

$$k_{\rm w} = (2.2U_{10} - 3.4) \left(\frac{S_c}{660}\right)^{-0.5},\tag{4}$$

where S_c is the Schmidt number of DMS, calculated as a function of temperature and salinity using the methods of *Johnson* [2010]. We calculated k_a for DMS as a function of wind speed and temperature using the methods of *Johnson* [2010].

2.2.1. Net DMS Fluxes

Using equations (2–4), we calculated the flux *F* per unit area of open water. Land and sea ice coverage data were used with *F* to calculate the net DMS flux from a 1° × 1° grid cell. Open water area in a given cell was calculated by subtracting land surface area and sea ice surface area, if any, from the cell. Following the *Takahashi et al.* [2009] approach for CO₂, we considered <10% sea ice coverage to have a negligible effect on the DMS flux from a cell, and the calculated flux was unmodified. Sea ice coverage >90% still allows some flux of DMS due to leads, polynyas etc. (where sea-air DMS fluxes may be very strong, *Else et al.* [2011]), so in this case coverage was set to 90%. Multiplying *F* (per unit area) by the remaining ice free ocean area gives the net DMS flux from the cell. This differs from the methodology of *Lana et al.* [2011], who set DMS flux to zero if sea ice coverage exceeded 75%, neglecting ice effects otherwise. In 2 years of monthly global 1° gridded data, the RMS difference in the annual hemisphere-integrated flux between the two methods of dealing with partial ice cover is negligible on a global scale (0.003 Tg S a⁻¹ in the northern hemisphere and 0.02 Tg S a⁻¹ in the southern hemisphere – compare with figures in Table 3).

2.2.2. Missing Data

In a given cell on a given day, it may be that not all the required data are present. RA2 has a small footprint, so daily coverage of RA2 data is generally smaller than that of AATSR. AATSR is vulnerable to cloud contamination, so it is quite rare for both data sets to give coincident valid data. The salinity and DMS_w data sets are also incomplete, with a small amount of missing data around some coastlines. To address this problem, we adopted a hierarchical strategy:

- 1. If all daily composite data sets are valid at a given position, we use these values;
- 2. If RA2 or AATSR daily data are missing, we use the monthly composite of that data set.
- 3. If monthly composite data are missing, we use a composite of that month over the whole time series (currently 2 years).
- 4. If time series composite data are missing, in the case of U_{10} , we substitute U_{10} from the daily ECMWF data. If SST_{skin} is missing, the cell flux is not directly calculable. The same is true if DMS_w or S is missing, but this only happens for a very small number of pixels around coastlines, where land masks differ slightly. In these cases, we estimate the cell flux as the monthly average flux in the corresponding 10° latitude band and apply it to the cell. This averaging occurs for 2.2% of the global open water area and 17% of the area in the Arctic over the 2 years.

2.3. Sensitivity to Temperature, Salinity, Wind Speed, and DMS Concentration

We initially varied SST_{skin} , salinity, U_{10} and DMS_w independently and uniformly over all months and positions. Though these are not realistic scenarios, they give us a simple insight into the sensitivity of the global flux to changes in individual climate variables. In order to give a more realistic simulation of the effects of predicted climate change, we then used output from the fully coupled oceanatmosphere Met. Office Hadley Centre Earth System model HadGEM2-ES [*Collins et al.*, 2008] downloaded from version 5 of the Coupled Model Intercomparison Project (CMIP5), using the RCP8.5 scenario [*Taylor et al.*, 2012]. This makes monthly global predictions from 2006 to 2100 assuming a scenario of high anthropogenic CO₂ emissions. We considered variations in *SST*, salinity, U_{10} , and DMS_w . The model *SST* is the temperature a few metres below the sea surface, not SST_{skin} . However, the difference is a few tenths of a degree, averaging 0.14° C for wind speeds greater than 6 ms⁻¹ [*Donlon et al.*, 2002], so the temporal changes described in subsection 3.2 are largely unaffected. Each parameter was reprojected to the $1^{\circ} \times 1^{\circ}$ grid and averaged over 6 years in each month of the year. The data for 2006–2011 were then compared with those for 2031–2036 to estimate the effect of 25 years of change. Averaging over 6 years reduces the effect of transient features in a given year, such as the El Niño-Southern Oscillation. We used pixel by pixel changes in this case rather than a uniform Table 2. Monthly Coverage of *E. Huxleyi* Blooms in the North Atlantic (75°W-11°E, 35–68°N) and Northeast Atlantic (40°W-11°E, 60–68°N) During April–September^a

		Coverage (%)					
	April	May	June	July	August	September	Annual
North Atlantic	0.003	0.08	0.29	0.95	0.44	0.14	0.16
Northeast Atlantic	0.003	0.005	0.36	3.7	2.8	0.87	0.65

^aCoverage in other months is assumed to be negligible.

global change because there is strong spatial variability, especially in salinity, U_{10} and DMS_{w} , which exhibit significant regions of both positive and negative trends.

The sensitivity of DMS flux to plausible changes in each of the four parameters was investigated by applying 25 years of change to one parameter while keeping the others fixed at their original values. *SST* variations were applied to k_w , k_a , H and hence F through the temperature dependence of S_c (k_a is not expected to have a large temperature dependence because the airside Schmidt number is largely temperature independent). For consistency, salinity variations were applied to k_w , H and hence F through this was not expected to have much effect. The salinities at the surface and at a depth of a few meters were assumed to covary. U_{10} and DMS_w changes were applied as percentages, since in both cases variations can be negative or positive and values can be close to zero. U_{10} variations were applied to k_w and k_a , and DMS_w variations were applied directly in equation (2). Finally, we applied 25 years of change to all four parameters (*SST*, salinity, U_{10} , and DMS_w) simultaneously.

This analysis enabled us to gain insight into the impact of plausible changes in temperature, salinity, wind speed, and aqueous DMS concentration from 2006–2011 to 2031–2036.

2.4. Contribution of Coccolithophores to the North Atlantic DMS Flux

The *Lana et al.* [2011] climatology was constructed from all available data and no overarching sampling strategy was employed to coordinate the data collection. Some past research activities have focussed on reduced sulfur cycling and certain fieldwork thus specifically targeted coccolithophore blooms [e.g., *Steinke et al.*, 2002b]. Other DMS data have been collected along transect lines, objectively analysing whatever seawater masses are encountered. Bloom features thus have the potential to be underrepresented or overrepresented within the climatology, which is not scaled to account for the spatial and temporal frequency of phytoplankton blooms.

Table 3. Annual Mean DMS Flux (Tg S a^{-1}) per 10 Degree LatitudinalBand in 2008–2009, Compared With the Climatology of Lana et al.[2011]

[2011]		
Latitude	2008–2009 Mean Flux	Climatology
90°-80°N	0.0	0.0
$80^{\circ}-70^{\circ}N$	0.1	0.1
70°-60°N	0.2	0.2
60°-50°N	0.6	0.9
50°—40°N	0.9	1.5
$40^{\circ}-30^{\circ}N$	1.0	1.5
$30^{\circ}-20^{\circ}N$	1.1	1.4
$20^{\circ}-10^{\circ}N$	1.8	2.6
10°-0°N	2.1	2.6
$0^{\circ}-10^{\circ}S$	2.1	2.2
$10^{\circ}-20^{\circ}S$	2.5	3.5
$20^{\circ}-30^{\circ}S$	2.1	3.0
$30^{\circ}-40^{\circ}S$	1.9	2.7
$40^{\circ}-50^{\circ}S$	1.6	2.8
$50^{\circ}-60^{\circ}S$	1.1	2.1
$60^{\circ}-70^{\circ}S$	0.5	0.9
$70^{\circ}-80^{\circ}S$	0.1	0.1
80°-90°S	0.0	0.0
Northern hemisphere	7.8	10.8
Arctic (north of 66°N)	0.2	-
Global	19.6	28.1

The coccolithophore Emiliania huxleyi is an ideal phytoplankton species for Earth Observation. When the plankton grow and divide at a high rate, they shed their calcium carbonate plates (liths) into the surrounding seawater [Paasche, 2001]. In bloom conditions, the liths create a strong backscattering signal that can be observed by satellite remote sensing [Balch et al., 1991], and can have a significant local effect on radiative forcing [Gondwe et al., 2001]. Recent work has facilitated the routine identification of coccolithophore blooms in the North Atlantic [Shutler et al., 2010]. Shutler et al. [2013] used this approach to estimate the proportion of time and space that coccolithophore blooms are present within the North Atlantic (75°W–11°E, 35–68°N), see Table 2.

Observations of surface water DMS values within phytoplankton blooms containing high coccolithophore cell density range between 4.8 and 25.0 nM (Table 1). How long this



increase in DMS is sustained over the course of a coccolithophore bloom is not well known. We combined satellitederived spatial coverage of North Atlantic coccolithophore blooms (calculated using the methods of *Shutler et al.* [2013] and summarized in Table 2) with a range of estimates of the increase in DMS levels due to the bloom and the duration of the increased DMS, to estimate coccolithophore bloomdriven sea-to-air DMS flux over plausible ranges of bloom DMS

Figure 1. Annual mean sea-air DMS flux (μ mol S m⁻² day⁻¹) over 2008–2009.

enhancement and duration, in the hope that future work will constrain these ranges. We calculated the increase in annual DMS flux assuming that blooms increase DMS by 5, 15, and 25 nM while a bloom is identified from satellite in a given pixel, and this effect is assumed to continue for 1–20 days, day 1 being the day of satellite detection.

3. Results and Discussion

3.1. Integrated DMS Flux

The mean daily open water DMS flux *F* over 2008–2009 is shown in Figure 1, and its seasonal variations are shown in Figure 2, which also shows eddy correlation measurements of DMS flux from 10 cruises (cruise mean, not seasonal mean). Cruises that overlap two seasons are shown in both. The annual integrated DMS fluxes over 10° latitude bands are listed in Table 3, along with the annual global, northern hemisphere, and Arctic (north of 66°N) DMS fluxes. These figures are compared with the equivalent climatological values from *Lana et al.* [2011]. The cruise data shown in Figure 2 are summarized in Table 1, which also shows the mean EO modeled DMS flux *F* ("EO Flux") over the cruise months in 2008–2009 in a 5°×5° window around the mean cruise position.

Eight out of 10 of the EO fluxes are less than the in situ fluxes, implying a possible bias. However, all but two of the differences are less than the standard deviation of in situ flux, and a paired t-test reveals no significant difference between the two (p = 0.16). As a further check, we estimated the bias between the two data sets by calculating the mean of each, weighted by the reciprocal of the standard deviation of in situ flux. The in situ mean was 5.2 μ mol m⁻² day⁻¹ and the EO mean was 4.9 μ mol m⁻² day⁻¹, both with a standard error of 1.1 μ mol m⁻² day⁻¹, implying an EO underestimate of 7 ± 28%.

The root mean square difference between the mean in situ fluxes and the mean EO flux is 3.1 μ mol m⁻² day⁻¹, similar to the global mean flux of 3.5 μ mol m⁻² day⁻¹ calculated from our data. Much of this variability can be accounted for by differences in DMS between ship and climatology (RMS difference in DMS is 32%, equating to a 32% difference in flux, see Section 3.2) and differences in wind speed between ship and satellite (RMS difference in wind speed is 16%, equating to a 21% difference in flux), while SST has much less effect (RMS difference of 0.9°C, suggesting a flux difference of 2.5%).

Lana et al. [2011] found a global flux of 28.1 Tg S a⁻¹, while our analysis suggests a global flux of 19.6 Tg S a⁻¹. In the northern hemisphere, *Lana et al.* [2011] estimate a flux of 10.8 Tg S a⁻¹ while we find 7.8 Tg S a⁻¹. A significant part of these differences can be ascribed to the choice of flux parameterization. This study uses a linear relationship between k_w and wind that has been derived from direct measurements of DMS flux [*Goddijn-Murphy et al.*, 2012], while *Lana et al.* [2011] use the quadratic relationship of *Nightingale et al.* [2000], resulting in increased fluxes at high wind speeds. It has been proposed that the difference in the functional form of these gas exchange parameterizations results from bubble-mediated gas transfer at high winds, which is enhanced for insoluble gases [*Woolf et al.*, 2007]. Highly insoluble gases were used to estimate the nonlinear relationship between k and wind speed in *Nightingale et al.* [2000]. In contrast, DMS is moderately soluble, and



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recent eddy correlation measurements at high wind speeds do not suggest gas-transfer velocity enhancements at high-wind speeds [*Bell et al.*, 2013; *Goddijn-Murphy et al.*, 2012; *Marandino et al.*, 2009; *Yang et al.*, 2011a]. When we recalculate our fluxes using *Nightingale et al.* [2000], we obtain a global flux of 23.1 Tg S a⁻¹ and a northern hemisphere flux of 8.8 Tg S a⁻¹.

3.2. Sensitivity to Predicted Changes in Temperature, Salinity and Wind Speed

In the initial sensitivity study, we changed a single parameter by a fixed amount, independent of location or month. In all cases, we found the effect to be linear ($r^2 > 0.997$), so only the effect per unit change in the parameter is quoted. The effect of increasing SST_{skin} uniformly was an increase in mean global DMS flux of 0.55 Tg S °C⁻¹ (+2.8% °C⁻¹). The effect of increasing salinity uniformly was a decrease in global flux of 0.028 Tg S PSU⁻¹ (-0.14% PSU⁻¹). The effect of increasing $\log(U_{10})$ uniformly (i.e., increasing U_{10} by a uniform proportion) was an increase in global flux of 0.25 Tg S per unit of (natural) $\log(U_{10})$, hence a 1% increase in U₁₀ results in an increase in global flux of 0.25 Tg S (+1.3%), reflecting the approximately linear relationship between flux and wind speed. The effect of increasing $\log(DMS_w)$ uniformly was an increase in global flux of 20 Tg S per unit of (natural) $\log(DMS_w)$, hence a 1% increase in DMS_w results in an increase in global flux of 0.19 Tg S (+1.0%), reflecting the approximately linear relationship between flux and DMS_w .

The modeled 25 year changes in *SST*, salinity, U_{10} and DMS_w in four seasons, each with 3 months, are shown in Figures 3–6. Some of the spatial and temporal variability of *SST*, salinity and DMS_w changes can be explained simply as a function of latitude, independent of longitude or month (22, 23, and 21% of variance explained for *SST*, salinity and DMS_w , respectively). Figure S1 (supporting information) shows the mean annual changes in *SST*, salinity, wind speed, and DMS_w over all longitudes as a function of latitude and month.

The effects on global annual integrated flux of applying 25 years of predicted changes in SST, salinity, U_{10} and DMS_w are shown in Table 4, along with the effects of changing all four parameters simultaneously. The



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Figure 3. Change in SST (°C) using CMIP5 data over the 25 years from 2006–2011 to 2031–2036.



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Figure 4. Change in salinity (PSU) using CMIP5 data over the 25 years from 2006–2011 to 2031–2036.



-0.10

-0.12

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-0.06

-0.04



Figure 5. Change in log₁₀(10 m wind speed) using CMIP5 data over the 25 years from 2006–2011 to 2031–2036.

-0.08



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Figure 6. Change in log₁₀(DMS concentration) using CMIP5 data over the 25 years from 2006–2011 to 2031–2036.

Table 4. Changes in Global Annual					
Mean DMS Flux (Tg S a^{-1}) Due to 25					
Years of Changes in SST, Salinity, 10 m					
Wind Speed and DMS Concentration					
Parameter	25 Year Change (Tg S a^{-1}				
CCT	1046(1220)				

+0.46 (+2.3%)
+0.10 (+0.5%)
-4.3 (-22%)
+0.33 (+1.7%)
-3.7 (-19%)

effects on annual mean *F* of applying 25 years of predicted changes are shown in Figure S2 (supporting information). It can immediately be seen from supporting information Figure S2 that the effect of salinity changes is by far the smallest except very close to the coast. The effects of *SST* changes are typically small and positive, increasing in areas where the flux is already large (Figure 1). The effects on flux of U_{10} changes are large and negative in most regions (due to the model predicting decreased wind speed), although less so in polar regions. The effect of *DMS_w* changes is highly variable, with localized "hot spots" of positive or negative change. The effect of changes in

all four parameters is dominated by the effect of wind, so the net effect is generally negative with some positive "hot spots," notably in the eastern Atlantic around 40°N, which has increased DMS_w (Figure 6), and in the Sea of Okhotsk north of Japan, which has increased wind speed and DMS_w (Figures 5 and 6). Applying 25 years of predicted changes in *SST*, U_{10} , and DMS_w affects seasonal mean *F* as shown in Figures 7–9. The effect of applying all four changes simultaneously is shown in Figure 10. The effect of salinity was generally as small in each season as the annual effect shown in supporting information Figure S2b, so this is not shown individually.

Our results suggest a decrease in global DMS flux in the 25 year scenario examined, mostly due to physical changes, i.e., reduced wind speed. It should be noted that if the surface DMS concentrations were to increase more than predicted by the CMIP5 model due to biological processes/feedbacks unaccounted for in the current model, the DMS flux would increase correspondingly.

3.3. Uncertainties and Error Propagation

Uncertainties and bias in the remote sensing data used to calculate *F* could have a significant impact on the resultant DMS fluxes. Previous work on CO₂ fluxes [*Land et al.*, 2013] has shown that the effect of random errors in *SST* and U_{10} is of the order of 1% of the total uncertainty estimate in the Arctic, and a similarly small proportion is likely for global DMS fluxes, hence this error source is neglected. To investigate the effect of possible bias in the input data sets, we follow *Land et al.* [2013] and take the precautionary step of using



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Figure 7. Changes in seasonal mean sea-air DMS flux (µmol S m⁻² day⁻¹) due to 25 years of predicted changes in SST.



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Figure 8. Changes in seasonal mean sea-air DMS flux (μ mol S m⁻² day⁻¹) due to 25 years of predicted changes in 10 m wind speed, on the same scale as Figure 7.



Figure 9. Changes in seasonal mean sea-air DMS flux (μ mol S m⁻² day⁻¹) due to 25 years of predicted changes in DMS concentration, on the same scale as Figures 7 and 8.



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Figure 10. Changes in seasonal mean sea-air DMS flux (μ mol S m⁻² day⁻¹) due to 25 years of predicted changes in SST, salinity, wind speed, and DMS concentration on the same scale as Figures 7–9.

the square root of the sum of squares of the published global bias and standard deviation for each data set. This allows for the possibility of regional biases that cancel in the global bias but contribute to the global standard deviation. The standard deviations of *SST* and U_{10} are 0.16° C and 1.25 ms^{-1} , respectively, and the global biases are 0.2° C and 0.28 ms^{-1} , combining to $\sqrt{0.16^2 + 0.2^2} = 0.26^{\circ}$ C and $\sqrt{1.25^2 + 0.28^2} = 1.28 \text{ ms}^{-1}$. These should be considered to be upper estimates of the error due to bias. The mean and standard deviation of *DMS*_w are given regionally by *Lana et al.* [2011]. We converted these to a regional error in $\ln(DMS_w)$ and found the area weighted global average of $\delta \ln(DMS_w)$ to be 0.12, or a 12% error in DMS_w . We can use the sensitivities found in subsection 3.2 to translate these errors in *SST* and DMS_w into errors in global DMS flux. These are 0.14 Tg S (0.7%) and 2.4 Tg S (12%), respectively. The fixed error in wind speed is incompatible with the proportional change used in subsection 3.2, so instead we offset all wind speeds by 1.28 ms^{-1} and recalculated the global flux, resulting in a change in global flux of 4.8 Tg S (25%). Assuming them to be uncorrelated, these errors sum to 5.4 Tg S (27%). Correlation between errors may result in higher total errors.

Goddijn-Murphy et al. [2012] give an RMS error of $\delta k_w = 5.5 \left(\frac{S_c}{660}\right)^{-0.5}$ for their *k* parameterization. Again assuming the worst case of bias equal to the RMS error, we recalculated fluxes using k values δk_w greater than predicted, resulting in an increase in global DMS flux of 9.4°Tg S (48%). Clearly this dominates the uncertainty in global flux, which then becomes 10.9 Tg S (55%).

3.4. Contribution of Coccolithophores to the North Atlantic DMS Flux

Table S1 (supporting information) shows the increase in annual mean flux over the whole North Atlantic due to DMS increases within detected coccolithophore blooms of 5, 15, and 25 nM, and with durations of 1–20 days. It is also shown as a percentage of the annual mean flux calculated from the *Lana et al.* [2011] climatology. Figure 11a shows the percentage data as a contour plot, while Figure 12 shows the spatial distribution of the increase in flux for the most conservative case (5 nM increase for 1 day). It should be noted that the most extreme case at the upper right of Figure 11a, an increase of 25 nM for 20 days, is highly unlikely to be a good representation of North Atlantic blooms as a whole; we include these extremes to ensure that the unknown actual values are represented in supporting information Table S1 and Figures 11

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Figure 11. (a) Change in North Atlantic annual mean flux; (b) change in monthly mean flux in the northeast Atlantic ($40^{\circ}W-11^{\circ}E$, $60-68^{\circ}N$) in July. Both are shown as a function of increase in DMS concentration within a coccolithophore bloom and duration from the last satellite bloom detection, expressed as a percentage of the mean flux calculated from the *Lana et al.* [2011] climatology. For reference, the annual mean flux in the North Atlantic is $4.44 \ \mu$ mol m⁻² day⁻¹ and the mean flux in the northeast Atlantic in July is 10.20 \ \mu mol m⁻² day⁻¹. Note that these include the effects of coccolithophores.

and supporting information S4. For a given increase and duration, this approach is likely to be an underestimate of the true DMS flux as satellite backscatter cannot identify all coccolithophores (some do not shed their liths) or other DMS-producing, noncoccolithophore species. Also, it should be noted that the method of *Shutler et al.* [2013] specifically targets anomalously high reflectances, and so does not detect any background concentration of coccolithophores, which may also contribute a DMS flux.

Our calculations suggest that the contribution to the annual DMS flux by coccolithophore blooms identified using satellite backscattering may be a relatively small proportion of the *Lana et al.* [2011] climatological flux, which includes some contributions from coccolithophores. This suggests that noncoccolithophore species and coccolithophore species not identifiable from space are collectively an important driver of North Atlantic seawater DMS concentrations. An alternative explanation for our analysis is that the ad hoc sampling approach for data in the DMS climatology has led to an inherent bias toward DMS blooms and a subsequent overestimate of the DMS flux. Objective sampling tracks (i.e., non bloom-focussed) for future DMS data collection would address this issue.

In summer, DMS flux from coccolithophores has the potential to be substantial (Table 2), particularly in the northeast Atlantic where blooms are more prevalent (Figure 12). To illustrate this, the increase in DMS flux due to coccolithophore blooms in the northeast quadrant of the North Atlantic (32°W to 11°E, 52 to 68°N, shown as a black box in Figure 12) in July is shown in Figure 11b as a percentage of the equivalent flux calculated from the *Lana et al.* [2011] climatology.

The progression of coccolithophore bloom activity is illustrated in supporting information Figure S3, showing the distribution of bloom effect on monthly DMS flux from April to August. Blooms start in the Celtic



Figure 12. Change in North Atlantic annual average DMS flux (μ mol S m⁻² day⁻¹) with a 5 nM increase in DMS concentration during a coccolithophore bloom and a duration of 1 day. White areas have no coccolithophore blooms detected, brown is land.

Sea and English Channel in April, moving to the shelf break west of Ireland in May, the shelf break north of Ireland, the North Sea and the Greenland Sea between Greenland and Iceland in June, concentrating in a small region west of Iceland and the southern North Sea in July and August. To investigate the local impact of coccolithophore blooms, we calculated the mean percentage effect on DMS flux for all pixels with nonzero effect, i.e., pixels identified as being bloom affected (pixels not colored white or brown in Figure 12 and supporting information Figure S3). Results are shown annually in supporting information Figure S4a and for July (the month of greatest effect on North Atlantic fluxes) in supporting information Figure S4b. The annual effect ranges from 1 to 38%, while the effect in July ranges from 5 to 147%.

To more accurately quantify the contribution of coccolithophore blooms to regional DMS emissions, detailed investigations are required into the relationship between bloom intensity (spectral backscattering) and seawater DMS concentration.

4. Conclusions

In this paper, we describe a new method for calculating net DMS fluxes from EO data using a *k* parameterization that is calibrated for DMS, showing the applicability of EO data to the monitoring of DMS net fluxes. Specifically, we look at the effect of predicted changes in climate on the flux of DMS from ocean to atmosphere. It shows that the effect of 25 years of changes predicted by a CMIP5 climate model is globally dominated by the prediction of decreased surface wind speed, which results in a 22% decrease in global DMS flux, while changes in temperature, salinity, and DMS concentration have smaller positive effects, with a combined effect of a 19% decrease in global flux. We also evaluate DMS emission from satellite-observed coccolithophore blooms in the North Atlantic, which is small compared to the annual climatology, but may be more important in the summertime and in the northeast Atlantic.

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References

- Arrigo, K. R., S. Pabi, G. L. van Dijken, and W. Maslowski (2010), Air-sea flux of CO2 in the Arctic Ocean, 1998–2003, J. Geophys. Res., 115, G04024, doi:10.1029/2009JG001224.
- Balch, W. M., P. M. Holligan, S. G. Ackleson, and K. J. Voss (1991), Biological and optical properties of mesoscale coccolithophore blooms in the Gulf of Maine, *Limnol. Oceanogr.*, 36, 629–643.
- Bates, T. S., B. K. Lamb, A. Guenther, J. Dignon, and R. E. Stoiber (1992), Sulfur emissions to the atmosphere from natural sources, J. Atmos. Chem., 14, 315–337, doi:10.1007/BF00115242.
- Bell, T. G., W. De Bruyn, S. D. Miller, B. Ward, K. Christensen, and E. S. Saltzman (2013), Air/sea DMS gas transfer in the North Atlantic: Evidence for limited interfacial gas exchange at high wind speed, Atmos. Chem. Phys. Discuss., 13(5), 13,285–13,322, doi:10.5194/acpd-13-13285-2013.
- Blomquist, B. W., C. W. Fairall, B. J. Huebert, D. J. Kieber, and G. R. Westby (2006), DMS sea-air transfer velocity: Direct measurements by eddy covariance and parameterization based on the NOAA/COARE gas transfer model, *Geophys. Res. Lett.*, 33, L07601, doi:10.1029/ 2006GL025735.
- Boucher, O., and U. Lohmann (1995), The sulfate-CCN-cloud albedo effect, *Tellus Ser. B*, 47(3), 281–300, doi:10.1034/j.1600-0889.47.issue3.1.x.
- Boucher, O., C. Moulin, S. Belviso, O. Aumont, L. Bopp, E. Cosme, R. Von Kuhlmann, M. G. Lawrence, M. Pham, and M. S. Reddy (2003), DMS atmospheric concentrations and sulphate aerosol indirect radiative forcing: A sensitivity study to the DMS source representation and oxidation, Atmos. Chem. Phys, 3, 49–65.
- Boutin, J., J. Etcheto, L. Merlivat, and Y. Rangama (2002), Influence of gas exchange coefficient parameterisation on seasonal and regional variability of CO2 air-sea fluxes, *Geophys. Res. Lett*, 29(8), 1182, doi:10.1029/2001GL013872.
- Cameron-Smith, P., S. Elliott, M. Maltrud, D. Erickson, and O. Wingenter (2011), Changes in dimethyl sulfide oceanic distribution due to climate change, *Geophys. Res. Lett.*, 38, L07704, doi:10.1029/2011GL047069.
- Charlson, R. J., and H. Rodhe (1982), Factors controlling the acidity of natural rainwater, *Nature*, *295*(5851), 683–685, doi:10.1038/295683a0. Charlson, R. J., J. E. Lovelock, M. O. Andreae, and S. G. Warren (1987), Oceanic phytoplankton, atmospheric sulphur, cloud, *Nature*, *3526*, 655–661. doi:10.1038/326655a0.
- Clarke, A. D., S. Freitag, R. M. C. Simpson, J. G. Hudson, S. G. Howell, V. L. Brekhovskikh, T. Campos, V. N. Kapustin, and J. Zhou (2013), Free troposphere as a major source of CCN for the equatorial pacific boundary layer: Long-range transport and teleconnections, *Atmos. Chem. Phys.*, 13, 7511–7529, doi:10.5194/acp-13-7511-2013.
- Collins, W. J., N. Bellouin, M. Doutriaux-Boucher, N. Gedney, T. Hinton, C. D. Jones, S. Liddicoat, G. Martin, F. O'Connor, and J. Rae (2008), Evaluation of the HadGEM2 model, *Hadley Cent. Tech. Note, 74*, Met. Office Hadley Centre, Exeter, U. K.
- Donlon, C. J., P. J. Minnett, C. Gentemann, T. J. Nightingale, I. J. Barton, B. Ward, and M. J. Murray (2002), Toward improved validation of satellite sea surface skin temperature measurements for climate research, *J. Clim.*, *15*(4), 353–369, doi:10.1175/1520-0442(2002)015<0353: TIVOSS>2.0.CO;2.
- Donlon, C. J., I. Robinson, K. S. Casey, J. Vazquez-Cuervo, E. Armstrong, O. Arino, C. Gentemann, D. May, P. LeBorgne, and J. Piollé (2007), The global ocean data assimilation experiment high-resolution sea surface temperature pilot project, *Bull. Am. Meteorol. Soc, 88*(8), 1197–1213, doi:10.1175/BAMS-88-8-1197.
- Donlon, C. J., M. Martin, J. Stark, J. Roberts-Jones, E. Fiedler, and W. Wimmer (2011), The operational sea surface temperature and sea ice analysis (OSTIA) system, *Remote Sens. Environ*, 116, 140–158, doi:10.1016/j.rse.2010.10.017.
- Elliott, S. (2009), Dependence of DMS global sea-air flux distribution on transfer velocity and concentration field type, J. Geophys. Res., 114, G02001, doi:10.1029/2008JG000710.
- Else, B. G. T., T. N. Papakyriakou, M. A. Granskog, and J. J. Yackel (2008), Observations of sea surface fCO2 distributions and estimated airsea CO2 fluxes in the Hudson Bay region (Canada) during the open water season, J. Geophys. Res, 113, C08026, doi:10.1029/ 2005JG000031.

 Else, B. G. T., T. N. Papakyriakou, R. J. Galley, W. M. Drennan, L. A. Miller, and H. Thomas (2011), Wintertime CO2 fluxes in an Arctic polynya using eddy covariance: Evidence for enhanced air-sea gas transfer during ice formation, *J. Geophys. Res., 116*, C00G03, doi:10.1029/2010JC006760.
 Galí, M., R. Simó, M. Vila-Costa, C. Ruiz-González, J. M. Gasol, and P. Matrai (2013), Diel patterns of oceanic dimethylsulfide (DMS) cycling: Microbial and physical drivers, *Global Biogeochem. Cycles, 27*, 620–636, doi:10.1002/gbc.20047.

Goddijn-Murphy, L. M., D. K. Woolf, and C. Marandino (2012), Space-based retrievals of air-sea gas transfer velocities using altimeters; calibration for dimethyl sulfide, J. Geophys. Res, 117, C08028, doi:10.1029/2011JC007535.

Gondwe, M., W. Klaassen, W. Gieskes, and H. de Baar (2001), Negligible direct radiative forcing of basin-scale climate by coccolithophore blooms, *Geophys. Res. Lett.*, 28(20), 3911–3914, doi:10.1029/2001gl012989.

Gunson, J. R., S. A. Spall, T. R. Anderson, A. Jones, I. J. Totterdell, and M. J. Woodage (2006), Climate sensitivity to ocean dimethylsulphide emissions, *Geophys. Res. Lett.*, 33, L07701, doi:10.1029/2005GL024982.

Halloran, P. R., T. G. Bell, and I. J. Totterdell (2010), Can we trust empirical marine DMS parameterisations within projections of future climate?, *Biogeosciences*, 7(5), 1645–1656, doi:10.5194/bg-7-1645-2010.

Huebert, B. J., B. W. Blomquist, J. E. Hare, C. W. Fairall, J. E. Johnson, and T. S. Bates (2004), Measurement of the sea-air DMS flux and transfer velocity using eddy correlation, *Geophys. Res. Lett.*, 31, L23113, doi:10.1029/2004GL021567.

Huebert, B. J., B. W. Blomquist, M. X. Yang, S. D. Archer, P. D. Nightingale, M. J. Yelland, J. Stephens, R. W. Pascal, and B. I. Moat (2010), Linearity of DMS transfer coefficient with both friction velocity and wind speed in the moderate wind speed range, *Geophys. Res. Lett.*, 37, L01605, doi:10.1029/2009GL041203.

Johnson, M. T. (2010), A numerical scheme to calculate temperature and salinity dependent air-water transfer velocities for any gas, Ocean Sci., 6(4), 913–932, doi:10.5194/os-6-913-2010.

Keller, M. D., W. K. Bellows, and R. R. L. Guillard (1989), Dimethyl sulfide production in marine phytoplankton, in *Biogenic Sulfur in the Environment*, vol. 393, edited by E. S. Saltzman and W. J. Cooper, pp. 167–182, AGU, Washington, D. C.

Kettle, A. J., and M. O. Andreae (2000), Flux of dimethylsulfide from the oceans: A comparison of updated data sets and flux models, J. Geophys. Res., 105(D22), 26,793–26,808, doi:10.1029/2000JD900252.

Lana, A., T. G. Bell, R. Simó, S. M. Vallina, J. Ballabrera-Poy, A. J. Kettle, J. Dachs, L. Bopp, E. S. Saltzman, and J. Stefels (2011), An updated climatology of surface dimethlysulfide concentrations and emission fluxes in the global ocean, *Global Biogeochem. Cycles*, 25, GB1004, doi:10.1029/2010GB003850.

Land, P. E., J. D. Shutler, R. D. Cowling, D. K. Woolf, P. Walker, H. S. Findlay, R. C. Upstill-Goddard, and C. J. Donlon (2013), Climate change impacts on sea-air fluxes of CO2 in three Arctic seas: A sensitivity study using Earth observation, *Biogeosciences*, 10(12), 8109–8128, doi: 10.5194/bg-10-8109-2013.

Liss, P. S., and P. G. Slater (1974), Flux of gases across the air-sea interface, Nature, 247, 181–184, doi:10.1038/247181a0.

Malin, G., S. Turner, P. Liss, P. Holligan, and D. Harbour (1993), Dimethylsulphide and dimethylsulphoniopropionate in the Northeast Atlantic during the summer coccolithophore bloom, Deep Sea Res., Part I, 40(7), 1487–1508, doi:10.1016/0967-0637(93)90125-M.

Marandino, C. A., W. J. De Bruyn, S. D. Miller, and E. S. Saltzman (2007), Eddy correlation measurements of the air/sea flux of dimethylsulfide over the North Pacific Ocean, J. Geophys. Res., 112, D03301, doi:10.1029/2006JD007293.

Marandino, C. A., W. J. De Bruyn, S. D. Miller, and E. S. Saltzman (2008), DMS air/sea flux and gas transfer coefficients from the North Atlantic summertime coccolithophore bloom, *Geophys. Res. Lett.*, 35, L23812, doi:10.1029/2008GL036370.

Marandino, C. A., W. J. De Bruyn, S. D. Miller, and E. S. Saltzman (2009), Open ocean DMS air/sea fluxes over the eastern South Pacific Ocean, Atmos. Chem. Phys., 9(2), 345–356, doi:10.5194/acp-9-345-2009.

Matrai, P. A., and M. D. Keller (1993), Dimethylsulfide in a large-scale coccolithophore bloom in the Gulf of Maine, Cont. Shelf Res., 13(8), 831–843, doi:10.1016/0278-4343(93)90012-M.

Nightingale, P. D., G. Malin, C. S. Law, A. J. Watson, P. S. Liss, M. I. Liddicoat, J. Boutin, and R. C. Upstill-Goddard (2000), In situ evaluation of air-sea gas exchange parameterizations using novel conservative and volatile tracers, *Global Biogeochem. Cycles*, 14(1), 373–387, doi: 10.1029/1999GB900091.

O'Carroll, A. G., J. R. Eyre, and R. W. Saunders (2008), Three-way error analysis between AATSR, AMSR-E, and in situ sea surface temperature observations, J. Atmos. Oceanic Technol, 25(7), 1197–1207, doi:10.1175/2007JTECHO542.1.

Paasche, E. (2001), A review of the coccolithophorid Emiliania huxleyi (Prymnesiophyceae), with particular reference to growth, coccolith formation, and calcification-photosynthesis interactions, *Phycologia*, 40(6), 503–529, doi:10.2216/i0031-8884-40-6-503.1.

Queffeulou, P., A. Benramy, and D. Croize-Fillon (2010), Analysis of seasonal wave height anomalies from satellite data over the global oceans, in *ESA Living Planet Symposium*, edited by H. Lacoste-Francis, ESA Publ. Div., Bergen, Norway.

Quinn, P. K., and T. S. Bates (2011), The case against climate regulation via oceanic phytoplankton sulphur emissions, *Nature*, 480(7375), 51–56, doi:10.1038/nature10580.

Shutler, J. D., M. G. Grant, P. I. Miller, E. Rushton, and K. Anderson (2010), Coccolithophore bloom detection in the north east Atlantic using SeaWiFS: Algorithm description, application and sensitivity analysis, *Remote Sens. Environ.*, 114(5), 1008–1016, doi:10.1016/ j.rse.2009.12.024.

Shutler, J. D., P. E. Land, C. W. Brown, H. S. Findlay, C. J. Donlon, M. Medland, R. Snooke, and J. C. Blackford (2013), Coccolithophore surface distributions in the North Atlantic and their modulation of the air-sea flux of CO2 from 10 years of satellite Earth observation data, *Bio-geosciences*, 10(4), 2699–2709, doi:10.5194/bg-10-1-2013.

Simó, R., and J. Dachs (2002), Global ocean emission of dimethylsulfide predicted from biogeophysical data, *Global Biogeochem. Cycles*, 16(4), 1018, doi:10.1029/2001GB001829.

Six, K. D., S. Kloster, T. Ilyina, S. D. Archer, K. Zhang, and E. Maier-Reimer (2013), Global warming amplified by reduced sulphur fluxes as a result of ocean acidification, *Nat. Clim. Change*, 3(11), 975–978, doi:10.1038/nclimate1981.

Stark, J. D., C. Donlon, A. O'Carroll, and G. Corlett (2008), Determination of AATSR biases using the OSTIA SST analysis system and a matchup database, *J. Atmos. Oceanic Technol*, 25(7), 1208–1217, doi:10.1175/2008JTECHO560.1.

Stefels, J., M. Steinke, S. Turner, G. Malin, and S. Belviso (2007), Environmental constraints on the production and removal of the climatically active gas dimethylsulphide (DMS) and implications for ecosystem modelling, *Biogeochemistry*, 83(1-3), 245–275, doi:10.1007/s10533-007-9091-5.

Steinke, M., G. Malin, S. W. Gibb, and P. H. Burkill (2002a), Vertical and temporal variability of DMSP lyase activity in a coccolithophorid bloom in the northern North Sea, *Deep Sea Res., Part II*, 49(15), 3001–3016, doi:10.1016/S0967-0645(02)00068-1.

Steinke, M., G. Malin, S. D. Archer, P. H. Burkill, and P. S. Liss (2002b), DMS production in a coccolithophorid bloom: Evidence for the importance of dinoflagellate DMSP lyases, Aquat. Microb. Ecol., 26, 259–270, doi:10.3354/ame026259.

Takahashi, T., et al. (2009), Climatological mean and decadal change in surface ocean pCO2, and net sea-air CO2 flux over the global oceans, *Deep-Sea Res., Part II*, *56*(8-10), 554–577, doi:10.1016/j.dsr2.2008.12.009.

Taylor, K. E., R. J. Stouffer, and G. A. Meehl (2012), An overview of CMIP5 and the experiment design, *Bull. Am. Meteorol. Soc.*, 93(4), 485, doi: 10.1175/BAMS-D-11-00094.1.

Twomey, S. (1991), Aerosols, clouds and radiation, *Atmos. Environ.. Part A*, *25*(11), 2435–2442, doi:10.1016/0960-1686(91)90159-5. Woodhouse, M. T., K. S. Carslaw, G. W. Mann, S. M. Vallina, M. Vogt, P. R. Halloran, and O. Boucher (2010), Low sensitivity of cloud condensa-

tion nuclei to changes in the sea-air flux of dimethyl-sulphide, *Atmos. Chem. Phys., 10*(16), 7545–7559, doi:10.5194/acp-10-7545-2010. Woodhouse, M. T., G. W. Mann, K. S. Carslaw, and O. Boucher (2013), Sensitivity of cloud condensation nuclei to regional changes in dimethyl-sulphide emissions, *Atmos. Chem. Phys, 13,* 2723–2733, doi:10.5194/acp-13-2723-2013.

Woolf, D. K., I. S. Leifer, P. D. Nightingale, T. S. Rhee, P. Bowyer, G. Caulliez, G. De Leeuw, S. E. Larsen, M. Liddicoat, and J. Baker (2007), Modelling of bubble-mediated gas transfer: Fundamental principles and a laboratory test, J. Mar. Syst, 66(1-4), 71–91, doi:10.1016/ j.jmarsys.2006.02.011.

Yang, M., B. W. Blomquist, and B. J. Huebert (2009), Constraining the concentration of the hydroxyl radical in a stratocumulus-topped marine boundary layer from sea-to-air eddy covariance flux measurements of dimethylsulfide, *Atmos. Chem. Phys.*, *9*(23), 9225–9236, doi:10.5194/acp-9-9225-2009.

Yang, M., B. W. Blomquist, C. W. Fairall, S. D. Archer, and B. J. Huebert (2011a), Air-sea exchange of dimethylsulfide in the Southern Ocean: Measurements from SO GasEx compared to temperate and tropical regions, *J. Geophys. Res*, *116*, C00F05, doi:10.1029/2010JC006526.

Yang, M., B. J. Huebert, B. W. Blomquist, S. G. Howell, L. M. Shank, C. S. McNaughton, A. D. Clarke, L. N. Hawkins, L. M. Russell, and D. S. Covert (2011b), Atmospheric sulfur cycling in the southeastern Pacific-longitudinal distribution, vertical profile, and diel variability observed during VOCALS-REX, Atmos. Chem. Phys, 11(10), 5079–5097, doi:10.5194/acp-11-5079-2011.