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Key Points:

- First map of the floe-scale distribution of ice algal chlorophyll *a* in Antarctic pack ice established
- Snow depth, ice thickness, and sea ice freeboard levels significantly affect ice algal chlorophyll *a* distribution
- Described relationships provide a first step toward upscaling, i.e., predictability of ice algal biomass from sea ice physical parameters

Supporting Information:

Supporting Information S1

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Antarctic pack ice algal distribution: Floe-scale spatial variability and predictability from physical parameters

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Abstract Antarctic pack ice serves as habitat for microalgae which contribute to Southern Ocean primary production and serve as important food source for pelagic herbivores. Ice algal biomass is highly patchy and remains severely undersampled by classical methods such as spatially restricted ice coring surveys. Here we provide an unprecedented view of ice algal biomass distribution, mapped (as chlorophyll *a*) in a 100 m by 100 m area of a Weddell Sea pack ice floe, using under-ice irradiance measurements taken with an instrumented remotely operated vehicle. We identified significant correlations (p < 0.001) between algal biomass and concomitant in situ surface measurements of snow depth, ice thickness, and estimated sea ice freeboard levels using a statistical model. The model's explanatory power ($r^2 = 0.30$) indicates that these parameters alone may provide a first basis for spatial prediction of ice algal biomass, but parameterization of additional determinants is needed to inform more robust upscaling efforts.

1. Introduction

Sea ice is a key driver of Southern Ocean physical and biogeochemical processes and plays a pivotal role in Antarctic marine ecosystem function [*Thomas*, 2017]. Antarctic pack ice provides a vast habitat for ice-associated algae which form distinct communities in surface, interior, and bottom layers of sea ice floes [*Meiners et al.*, 2012; *Arrigo*, 2017]. Model estimates suggest that sea ice algal production can contribute significantly to the overall annual primary production of the ice-covered Southern Ocean [*Saenz and Arrigo*, 2014]. However, modeled ice algal production and biomass estimates are subject to considerable uncertainty as field-based observations of ice algal biomass, necessary to evaluate ecosystem models, remain extremely sparse [*Meiners et al.*, 2012]. The high spatial heterogeneity of ice algal distribution averts reliable in situ biomass estimates of biomass [*Mundy et al.*, 2007; *Lange et al.*, 2016].

Physical sea ice properties, such as snow depth, ice thickness, and surface slush layers, are considered key factors in controlling ice algal biomass [*Eicken*, 1992; *Fritsen et al.*, 1998; *Mundy et al.*, 2005; *Vancoppenolle et al.*, 2013], but their role in driving the floe-scale spatial distribution of pack ice algae remains poorly understood. Snow on Antarctic pack ice, for example, reduces light needed for algal photosynthesis but can also suppress pack ice floes below seawater levels, thereby replenishing nutrients and promoting algal growth in surface slush and refrozen slush (= snow ice) layers [*Fritsen et al.*, 1998; *Arrigo et al.*, 2014]. Scavenging of phytoplankton by frazil crystals during ice formation can result in the development of interior communities [*Spindler*, 1994], which can also emerge during ridging and rafting [*Ackley and Sullivan*, 1994]. While generally lower in biomass than bottom and surface communities, interior communities can significantly contribute to integrated biomass values [*Meiners et al.*, 2012]. Thus, thicker sea ice may be associated with increased integrated ice algal biomass [*Meiners et al.*, 2011; *Arrigo et al.*, 2014]. Improving

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measurement capabilities to simultaneously determine sea ice physical properties and in situ ice algal biomass, and establishing statistical relationships between these parameters, is necessary to develop and test the feasibility of upscaling efforts. The ultimate goal of such efforts would be estimation of ice algal biomass from remotely sensed estimates of physical sea ice properties, such as emerging satellite products for snow depth and ice thickness.

The presence of ice algal photosynthetic pigments affects under-ice irradiance spectra through absorption of specific wavelengths in the photosynthetic active radiation (400–700 nm) range [e.g., *Perovich et al.*, 1993; *Arrigo and Sullivan*, 1994; *Fritsen et al.*, 2011; *Hawes et al.*, 2012; *Lange et al.*, 2016]. Changes in spectra, often characterized by the difference of intensities at two key wavelengths, have been used to develop algorithms to estimate sea ice algal chlorophyll *a* (chl *a*, as proxy for biomass) from under-ice spectra collected with moored and horizontally profiling instruments [*Mundy et al.*, 2007; *Campbell et al.*, 2015; *Lange et al.*, 2016]. Employing a normalized difference index (NDI) approach, *Melbourne-Thomas et al.* [2015, 2016] recently established an NDI algorithm to estimate ice algal chl *a* in Weddell Sea pack ice from irradiance spectra measured beneath sea ice. This algorithm accounted for 79% of the total variation of integrated ice algal chl *a* in Weddell Sea pack ice.

Recent advance in the development of underwater vehicles and under-ice trawls has provided new opportunities to study sea ice physical and biological properties on small to medium scales (up to 1000 m). These include measurements and mapping of sea ice draft [*Wadhams and Doble*, 2008; *Williams et al.*, 2013, 2015] and light transmission [*Nicolaus et al.*, 2012; *Nicolaus and Katlein*, 2013; *Katlein et al.*, 2015a; *Arndt et al.*, 2017]. Biological properties such as the distribution of bottom-ice algae and under-ice algal aggregates [*Gutt*, 1995; *Katlein et al.*, 2015b], and most recently integrated ice algal biomass [*Lange et al.*, 2016], have also been determined.

In this study, we present the first map of the chl *a* concentration of an Antarctic pack ice floe in a 100 m by 100 m area. We analyzed these data in combination with concomitant surface measurements of snow depth, sea ice thickness, and calculated sea ice freeboard estimates in a first attempt to identify physical controls of the distribution of Antarctic pack ice algal biomass at the sea ice floe scale.

2. Material and Methods

2.1. Study Site and Field Procedures

Sampling was carried out as part of the "Winter Ice Study on Key Species" experiment in the Weddell Sea during late austral winter-early austral spring 2013 on the German icebreaker R/V Polarstern (voyage ANTXXIX-7/ PS81). The study site was at an approximate position of 60.78°S and 26.36°W and located in the center of a deformed first year pack ice floe composed of smaller ice floes frozen together [Meyer and Auerswald, 2014]. Remotely operated vehicle (ROV) and concomitant surface surveys were conducted as outlined in detail in a related paper on light transmission through Antarctic pack ice [Arndt et al., 2017] (see also supporting information). Briefly, under-ice irradiance spectra were collected with a ROV equipped with an upward looking hyperspectral radiometer (TriOS, Ramses-ACC) in a designated 100 m by 100 m area during 17 deployments between 18 and 26 September 2013. Atmospheric and ice surface conditions remained stable during this period [Arndt et al., 2017], which allowed merging of all observations for the following analyses. A Long Base Line positioning system, interrogating four ice-tethered transponders, coupled to GPS units (at the ice surface), and spaced around the survey site in a 400 m by 400 m rectangle, was used to determine and capture the ROV position data [e.g., Williams et al., 2015] (for details, see supporting information). The surface GPS data allowed for precise registration of any ice floe drift and rotation in between ROV and surface surveys and were used to merge under-ice survey data with the temporally displaced surface measurements. Snow depth and total ice thickness measurements were conducted on a single day after completion of the last ROV survey to prevent disturbance of the snow surface features. A sled-mounted, GPS-enabled, multifrequency electromagnetic induction instrument (GEM-2, Geophex Ltd.) [Hunkeler et al., 2015] was used to measure total ice thickness (sea ice thickness plus snow depth) in the ROV grid. Snow depths were measured every two steps along the GEM-2 tracks using a GPS-equipped Snow Depth Probe (Snow-Hydro, USA).

In addition to the spatial survey, paired point measurements of under-ice irradiance and ice core chl *a* were carried out at five sites on the ice floe. These were used to evaluate the applicability of an existing NDI

algorithm [*Melbourne-Thomas et al.*, 2015, 2016]. For full details of the sample site selection and NDI evaluation, see the supporting information. Field-based under-ice irradiance measurements, ice core collection, and fluorometric in vitro analysis of ice algal chl *a* exactly followed the methods outlined in *Melbourne-Thomas et al.* [2015].

2.2. Data Preparation

Analysis of under-ice irradiances was restricted to ROV radiometer data with a distance of ≤ 2 m to the sea ice subsurface, resulting in 4482 valid under-ice spectra along 17 dive tracks. All ROV measurements taken within an area of 2 m by 2 m were averaged to account for ROV track overlap [*Arndt et al.*, 2017]. All grid cells overlap for 1 m translating to an effective grid resolution of 1 m (running mean).

Ice thickness was calculated from the total snow + ice GEM-2 data by subtraction of the Snow Depth Probe data from the GEM-2 data. Using nearest neighbor resampling, we interpolated these surface measurements to the positions of the under-ice irradiance measurements.

Sea ice freeboard levels are controlled by the buoyancy of sea ice and the mass of the snow load. We calculated freeboard levels as a function of snow depth and ice thickness using mean density values for seawater, sea ice, and snow [*Yi et al.*, 2011; *Arndt et al.*, 2017] (for details, see supporting information). Negative freeboard values indicate flooding, while negative freeboard magnitudes specify the slush layer thickness.

Collected broadband hyperspectral under-ice irradiance spectra were interpolated to a spectral resolution of 1 nm. From these we calculated the optically derived integrated ice algal biomass values (Chl a_{ROV} , mg m⁻²), using the established NDI to ice algal chl *a* relationship of *Melbourne-Thomas et al.* [2015, 2016], as detailed in the supporting information (Figure S1).

2.3. Statistical Analyses

Empirical variograms were used to explore scales of spatial variability in physical and biological data sets. Variograms describe the variance in spatially binned subsets of the data as a function of varying distances (lag distances). Theoretical exponential variograms were fitted to estimate variogram "range" parameters, i.e., the lag distance at which the model flattens out and reaches 95% of its sill value [*Cressie*, 1991]. The range value provides a measure of the spatial autocorrelation length scale of the respective parameter.

To analyze the relationship between ice algal biomass and sea ice physical parameters (snow depth, ice thickness, and sea ice freeboard as predictor variables), we employed a Generalized Additive Model approach (GAM) [*Wood*, 2006], as relationships were not necessarily expected to be linear, using the *mgcv* R package [*Wood*, 2016]. A Gaussian error distribution was used, with the Chl a_{ROV} and ice thickness values log transformed prior to analysis to normalize the residuals. Outliers at very high and low values of ice thickness and snow depth were removed (1.27% of total data). Spatially autocorrelated data can affect the smoothness estimation of model terms; i.e., the estimation procedure in GAM is likely to undersmooth. We therefore also implemented the same model including a spatial term interaction on *x* and *y* (full tensor product smooth). This model (GAM_{*xy*}) accounted for effects of ROV position, *x* and *y* values, minimizing the effects of spatial autocorrelation on the estimation for the physical parameters (supporting information Figure S2). Significance levels were evaluated using *F* statistics [*Wood*, 2006]. GAM analyses were conducted using the statistical software R (version 3.3.1, R Development Core Team, 2016).

3. Results and Discussion

3.1. Point Measurements

The integrated chl *a* concentrations from five ice cores (point measurements) collected from the sea ice floe under investigation ranged from 1.65 to 11.42 mg m⁻² (mean \pm SD = 4.34 \pm 4.02 mg m⁻²; median = 3.22 mg m⁻²; supporting information Figure S1). Normalized difference indices (NDIs) calculated from concomitant under-ice irradiance measurements showed good agreement with a published NDI to chl *a* relationship for springtime Weddell Sea ice [*Melbourne-Thomas et al.*, 2015, 2016], and thus, this existing algorithm was applied to the ROV data (supporting information Figure S1).

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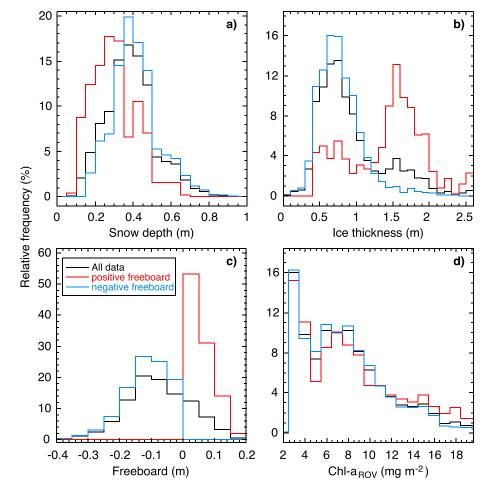


Figure 1. Frequency distributions for (a) snow depth, (b) ice thickness, (c) calculated freeboard levels (for details see text), and (d) integrated ice algal biomass as chlorophyll *a* (Chl a_{ROV}) determined from under-ice irradiance measurements using the algorithm by *Melbourne-Thomas et al.* [2015] for all data points (*n* = 4482, black) and grouped according to freeboard levels (positive freeboard = red (*n* = 1041), negative freeboard levels indicating surface flooding = blue (*n* = 3441)).

3.2. Spatial Survey

The area surveyed by ROV was part of a large $(>1 \text{ km}^2)$ first year pack ice floe showing highly variable snow depths and ice thicknesses (Figures 1 and 2) and situated in a band of dense pack ice. The study site demonstrated substantial snow accumulation with snow depth ranging from 0.08 m to 0.91 m (mean = 0.39 ± 0.13 m) and a modal snow depth of 0.39 m (Figures 1a and 2a). Sea ice thickness varied between 0.03 m and 2.57 m (mean = 0.93 ± 0.45 m, mode = 0.79 m) with a noteworthy contribution of thicker sea ice as indicated by the tail in the frequency distribution (Figure 1b). Thick sea ice was observed in distinct clusters denoting localized deformation features (Figure 2b). Measured snow depths and ice thicknesses were higher but broadly consistent with climatological mean snow depth and ice thickness values for eastern Weddell Sea pack ice as reported in the ASPeCt observations data set [Worby et al., 2008], the most comprehensive data compilation available. The sampled sea ice area was thus characteristic for a deformed first year Antarctic pack ice floe with heavy snow cover. The heavy snow load resulted in widespread flooding and occurrence of slush layers, as observed during sampling and indicated by the large proportion (76.8%) of negative values in calculated freeboard levels (Figures 1c and 2c). Widespread flooding is a key feature of Weddell Sea ice [Lange et al., 1990] and more generally Antarctic pack ice [Sturm and Massom, 2017]. Snow depth distributions were similar overall for negative and positive freeboard levels, albeit snow depths <0.3 m had a relatively high occurrence in positive freeboard grid cells. Areas of negative freeboard predominantly occurred where ice thicknesses were between 0.4 m and 1.2 m, although negative freeboard was calculated at ice thicknesses up to 2 m. Ice thicknesses associated with positive freeboard levels were

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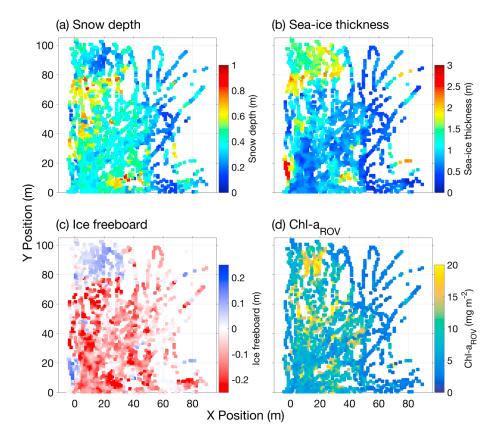


Figure 2. Spatial distribution of sea ice physical properties and ice algal biomass in the ROV survey area. (a) Snow depth as measured with the Snow Depth Probe, (b) ice thickness calculated from the GEM-2 measurements, (c) ice freeboard levels calculated as function of snow depth and ice thickness (for details see text), and (d) integrated ice algal biomass as chlorophyll *a* (Chl *a*_{ROV}) determined from under-ice irradiance measurements using the algorithm by *Melbourne-Thomas et al.* [2015]. Snow depth and sea ice thickness measurements are interpolated to ROV transect lines.

variable and ranged over 0.4 m to 2.5 m. Within the survey area, the 2-D distribution of sea ice freeboard levels showed general agreement with the distribution of sea ice thickness (Figures 2b and 2c), although estimated autocorrelation length scales, i.e., variogram ranges, were distinctly different with values of 28.8 m, 41.7 m, and 19.6 m for snow depth, ice thickness, and freeboard levels, respectively (Figure 3).

Estimated integrated ice algal chl *a* concentrations (Chl a_{ROV} , calculated according to *Melbourne-Thomas et al.* [2015, 2016]) ranged from 1.96 to 19.95 mg m⁻² (mean = 7.22 ± 4.00 mg m⁻²) with a median value of 6.62 mg m⁻². This relatively large range contrasts the smaller range encountered in the data set for five ice cores sampled from this ice floe (supporting information Figure S1). Classical ice coring surveys typically collect only a small number of cores [*Miller et al.*, 2015]. Our results highlight that small ice core sample sizes might result in large biases of floe-scale ice algal biomass estimates due to insufficient spatial representation of the collected ice cores. This has recently been discussed in detail by *Lange et al.* [2016] for Arctic pack ice. The Chl a_{ROV} values showed a highly skewed frequency distribution with a high relative contribution of low values (Figure 1d), consistent with the ASPeCt-Bio observational data set in which about 50% of all ice cores (n = 1300) show an integrated ice algal chl *a* value of <3 mg m⁻² [*Meiners et al.*, 2012].

The map of Chl a_{ROV} shows a highly variable distribution with no simple association to the spatial distribution of snow depth, ice thickness, or freeboard levels (Figure 2d). The variogram-based range value for Chl a_{ROV} is 17.3 m (Figure 3), which may provide some indication of the "patch" scale of integrated ice algal biomass in the studied sea ice floe. The difference between this range and the calculated range values for snow depth, ice thickness, and freeboard levels highlights the complexity in relationships between sea ice physical parameters and ice algal biomass as reported in previous studies and reviews [e.g., *Arrigo*, 2014; *Meiners*

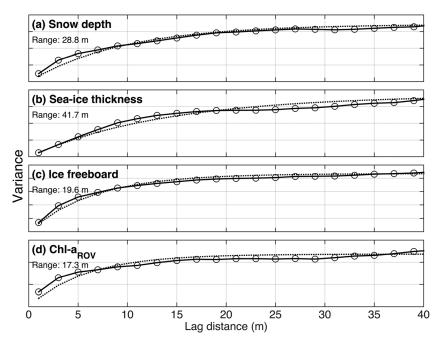


Figure 3. Variograms for (a) snow depth, (b) ice thickness, (c) calculated freeboard level, and (d) integrated ice algal biomass (as chlorophyll *a* determined from under-ice irradiance measurement (Chl a_{ROV})). Open circles with black lines indicate empirical variograms, and dotted lines indicate theoretical exponential variograms from which the range parameters were estimated. Variances are shown in arbitrary units starting at zero.

and Michel, 2017; Thomas, 2017]. Snow on sea ice, for example, influences ice algal biomass in multiple ways. Due to its high albedo and light attenuation properties, snow is a key factor controlling the light available to ice algal communities [Leu et al., 2015; Campbell et al., 2015]. Early in the growth season, snow cover can result in light limitation of ice algal growth, whereas later in the season snow cover can prevent photoinhibition of ice algal communities which are generally shade adapted [Arrigo, 2014, 2017]. In addition, snow cover can insulate sea ice, thus preventing internal ice melting and bottom ablation [Sturm and Massom, 2017], thereby stabilizing ice algal habitats. In the present study, integrated ice algal biomass (Chl a_{ROV}) was negatively correlated with snow depth, particularly at snow depths above 0.4 m (GAM: $s(Z_{snow})$, p < 0.0001, Figure 4a; GAM_{xv}: $s(Z_{snow})$, p = 0.007; supporting information Figure S2a), which is consistent with the lightlimiting role of snow during the winter-early spring transition. Our late-winter/early-spring data (September) contrast with summer (December-January) results of Arrigo et al. [2014], which showed a positive relationship between snow depth and integrated ice algal biomass for Amundsen Sea ice. In combination, these studies support the notion of a seasonally changing influence of snow on ice algal biomass in the Antarctic pack ice zone, as recently shown for Arctic landfast sea ice [Leu et al., 2015; Campbell et al., 2015]. We hypothesize that early in the season snow on Antarctic pack ice limits ice algal accumulation, while later in the season snow cover supports ice algal accumulation.

Chl a_{ROV} showed a positive correlation with sea ice thickness (GAM: $s(log[Z_{ice}])$, p < 0.0001, Figure 4b; GAM_{xy}: $s(log[Z_{ice}])$, p < 0.0001; supporting information Figure S2b), particularly at an ice thickness above 1 m. Internal ice algal communities are a key feature of Antarctic pack ice [*Arrigo*, 2017] and have previously been shown to have an increased probability of occurrence in thicker sea ice, e.g., due to sea ice deformation processes such as rafting [*Ackley and Sullivan*, 1994; *Meiners et al.*, 2012]. Moreover, microalgae are incorporated into sea ice during ice formation through various processes including scavenging [*Spindler*, 1994] and retention of particulate material during thermodynamic ice growth [*Janssens et al.*, 2016]. Thus, most Antarctic pack ice types contain at least low levels of algal chl *a*, and an increase in sea ice thickness results in an increase in integrated chl *a* values.

We note that our GAM analyses suggest a likely interaction between ice thickness and snow depth, whereby thicker ice can support higher snow loadings. This may moderate the negative snow effects on integrated algal biomass early in the season and allows for the final outcome that thicker ice harbors more integrated

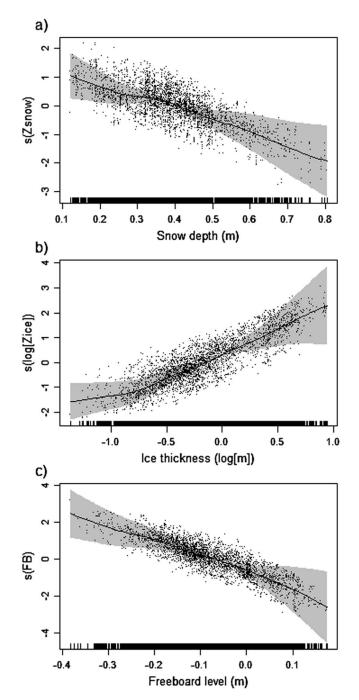


Figure 4. Smooths of generalized additive modeling (GAM) terms showing the effect of physical parameters on integrated ice algal biomass (Chl a_{ROV} , determined from under-ice irradiance measurements). (a) Snow depth, (b) ice thickness, and (c) calculated freeboard levels. Locations of observations (n = 4425) are shown as vertical lines on the *x* axes. Solid lines are the estimates of the smooths, shaded areas are standard errors of the estimated smooths, and points are the observation partial residuals. Note the natural log scale on *x* axis in Figure 4b.

algal biomass even with a higher snow load. Our data set contains a low amount of data (<2.4%) with high (>2.0 m) and low (<0.2 m) ice thicknesses. Future surveys of a larger range of ice thicknesses would be valuable to explore this observation and determine potential thresholds.

Surface flooding and subsequent snow ice formation are estimated to affect large parts of the Antarctic ice pack [Wadhams et al., 1987; Sturm and Massom, 2017], and snow ice is considered to contribute a significant proportion to the total Antarctic sea ice volume [Jeffries et al., 1998; Maksym and Markus, 2008]. The flooding water may originate from sea ice brines, if the ice is permeable [Golden et al., 1998; Jutras et al., 2016], or can be seawater moving laterally from ice floe edges and cracks [Massom et al., 2001]. Flooding of snow and surface ice may seed these layers with microalgae and increases nutrient availability. Both field observations [Fritsen et al., 1994] and modeling studies [Saenz and Arrigo, 2012, 2014] suggest that surface flooding and freeze cycles are critically important for ice algal biomass accumulation in Antarctic pack ice surface layers. Consistent with these studies, the Chl a_{ROV} data in our study were significantly negatively correlated with calculated freeboard levels (GAM: s(FB), p < 0.0001, Figure 4c; GAM_{xv}: s(FB), p < 0.0001; supporting information Figure S2c), i.e., potentially flooded areas with slush layers showed increased integrated ice algal biomass.

3.3. Conclusion

In conclusion, our study provides an unprecedented view of the spatial variability of ice algal biomass (as chl

a) in Antarctic pack ice and constitutes an important step toward estimating ice algal biomass from sea ice physical parameters. We observed a large difference between the mean and range of ice algal biomass estimates based on ice core sampling versus optically derived measurements, further highlighting the

uncertainty in ice algal biomass estimates from limited ice coring surveys [*Lange et al.*, 2016]. We suggest that newly available methods for floe-scale ice algal biomass estimation, e.g., using instrumented underwater vehicles as demonstrated in this study, will be better suited for comprehensive biomass estimates on relevant spatial scales.

Our GAM analyses showed highly significant floe-scale relationships between measured physical sea ice properties and integrated ice algal biomass for this late-winter/early-spring Antarctic pack ice study. The model had moderate explanatory power (GAM: $R_{adj}^2 = 0.30$, 30.8% of the deviance explained; GAM_{xy}: $R_{adi}^2 = 0.55, 55.7\%$ of the deviance explained), considering the relatively few predictor variables. This suggests feasibility of spatial upscaling, such as the estimation of local-to-regional Antarctic ice algal biomass from emerging remote sensing snow depth and ice thickness products [Kurtz and Markus, 2012]. Such advances promise a new independent way to estimate ice algal biomass and will help to inform and evaluate ice algal primary production models. However, broad-scale prediction of Antarctic pack ice algal biomass from physical sea ice properties alone needs further assessment. To achieve this, an improved knowledge of the spatial variability of ice algal biomass and an improved understanding of the temporal scales of coupling/decoupling between sea ice physical processes and ice algal biomass development are both required. Furthermore, additional physical sea ice properties, such as under-ice rugosity and sea ice temperature (as proxy for porosity), which are expected to influence ice algal distribution need to be measured and accounted for. The ready availability and swift emergence of underwater vehicle technologies should support a rapid proliferation of independent data sets at local-to-regional scales, and enable sampling of a wider range of icescapes as well as targeting seasonal differences and progressions.

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Supporting Information for

Antarctic pack-ice algal distribution: floe-scale spatial variability and predictability from physical parameters

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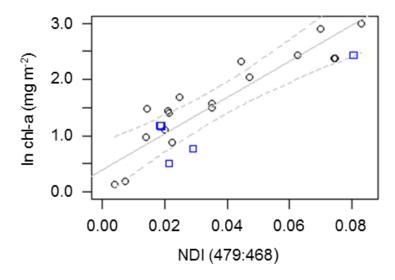
Description of methods used for ice-core sampling, optical point-measurements and chlorophyll-a analysis:

Paired under-ice irradiance measurements and ice core samples were collected from 5 sites on the ice floe under investigation. These bio-optical sampling sites were chosen such that their immediate surroundings (5 m) showed no snow-cover disturbances, were level, and free of major surface deformations. On each sampling site we deployed a TriOS Ramses ACC VIS radiometer (Advanced Cosine Collector, 180° field of view) 0.15 m beneath the subsurface of the ice floe through an access hole (0.11 m diameter) using a retractable L-shaped stainless steel arm. The effect of the hole on the light measurements was assumed to be negligible due the position of the radiometer 1.2 m north (i.e., directed towards the sun) of the access hole. Under-ice high-resolution (400 -700 nm, 3.3 nm band width) irradiance spectra were recorded with a laptop computer using TriOS MSDA XE software version 7.5.1. Results used for further analyses represent average spectra calculated from of a minimum of 10 replicate radiometer measurements. At each site one ice core was collected directly above the radiometer location using a manual Kovacs Mark II ice core system (0.09 m core diameter). The sampled ice core was cut into 4 sections which were placed in clean polyethylene containers and transported back to the ship's laboratories. On the ship, sampled ice core sections were melted at 4° C in the dark within 24 - 36 hours of collection. After the ice cores had melted, the samples were gently mixed and samples taken for the determination of ice algal pigment concentration. For pigment analysis, 0.2 to 4.0 litres of melted ice core subsamples were filtered onto Whatman GF/F glass fibre filters, extracted with 90% (v/v) acetone and analysed for chl-a with a Turner Designs 10AU fluorometer according to standard protocols (Melbourne-Thomas et al., 2015). Ice core section data (chl-a in µg 1^{-1}) were integrated over the entire ice thickness and are expressed as integrated chl-a values (mg m^{-2}).

Calculation of Normalized Difference Indices:

The calculation of the Normalized Difference Index (NDI) to integrated ice core chl-a relationship exactly followed Melbourne-Thomas et al. (2015) and considered their corrigendum to the equations (*Melbourne-Thomas et al.*, 2016). Measured under-ice transmitted irradiance spectra were interpolated to consecutive integer (1 nm) wavelengths in the range of 400 to 700 nm, and normalised to the integrated under-ice irradiance over the 400 to 700 nm wavelength range in order to minimise the amplitude component of spectral variability and to focus on differences in spectral shape. The Normalized Difference Index for the wavelength pair 479 nm and 468 nm was then calculated as:

NDI (479:468) = $[E_d(479nm) - E_d(468nm)] / [E_d(479nm) + E_d(468nm)]$ (1) In which E_d is the normalized intensity of under-ice irradiance in the specific wavelength bands (*Mundy et al.*, 2007; *Melbourne-Thomas et al.*, 2015, 2016; *Lange et al.*, 2016). NDI (479:468) values were related to log-normalized integrated chl-a values (SI, Fig. S1).



<u>Figure S1:</u> Normalized Difference Index (NDI) to integrated chlorophyll *a* concentration relationships. Black circles, regression line and 99% confidence intervals show the established relationship of Melbourne-Thomas et al. 2015 [In (Chla_{ROV}) = $0.39 + 31.7 \times NDI_{(479:468)}$, R² = 0.79, p < 0.0001). Blue squares indicate measurements taken during this study (n = 5). For details see text.

Calculation of sea-ice freeboard levels:

Sea-ice freeboard levels (F) were calculated according to Arndt et al. (2017), as:

 $F = (I x (\rho_w - \rho_i) - S x \rho_s) / \rho_w$ for positive freeboard.
(2)

Negative freeboard levels (-F), e.g., the flooded snow layer thickness were calculated as: -F = S - ((I+S)/(1+($\rho_s / (\rho_w - \rho_i)$))). (3)

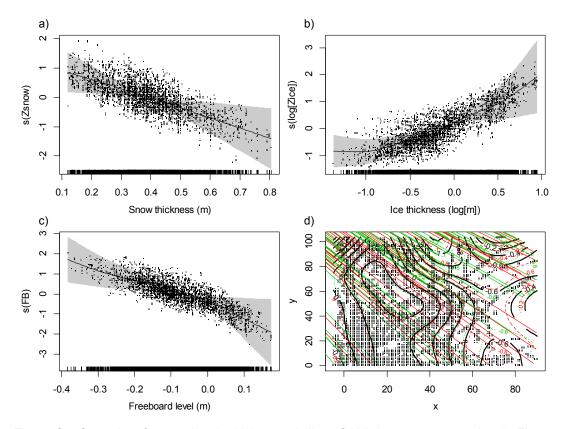
The latter equation accounts for a change in density of snow once flooded. In these equations, S, I, ρ_s , ρ_w and ρ_i denote snow depth (m), ice thickness (m), density of snow (= 300 kg m⁻³), density of water (= 1023.9 kg m⁻³) and density of ice (= 915.1 kg m⁻³), respectively.

Details on the instrumentation of the Remotely Operated Vehicle (ROV) used for under-ice surveys

Under-ice surveys were carried out using a SAAB SeaEye Falcon ROV instrumented with an upward-looking TriOS Ramses-VIS hyperspectral radiometer (320-950 nm) and an upward-looking Valeport500 sonar to measure under-ice irradiance and the distance between the sensor and the bottom of the ice, respectively. ROV position was determined using a long baseline (LBL) acoustic positioning system consisting of a Woods Hole Oceanographic Institution (WHOI) FSK micro-modem equipped with an ITC 3013 transducer (mounted to the ROV) and interrogating four Benthos XT-6001 transponders (15 m depth).

Statistical model results including spatial interaction

Figure S2 shows output from the GAM_{xy} model that includes a spatial smooth term (full tensor product interaction) on the x and y coordinates, in addition to the smooth functions for snow depth, ice thickness and ice freeboard. Note that the smooth functions for the biophysical terms remain almost identical to those shown in Fig. 4 (main text), with slightly widened confidence intervals but no substantive changes to the significance of terms. This indicates that even accounting for the strong spatial autocorrelation in the



dataset (GAM_{xy}: te(x,y), p < 0.0001, Fig. S2d) the algal relationships with biophysical drivers are strong.

Figure S2: Smooths of generalized additive modelling (GAM_{xy}) terms presented as in Figure 4 but including also d) the smooth interaction on the spatial x-, and y- co-ordinates. In panel d) the black lines indicate the tensor smooth estimate te(x,y), the red and green dashed lines indicate ± 1 S.E, and the points indicate the locations of ROV observations (n = 4425). Note natural log scale on x-axis in panel b.

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